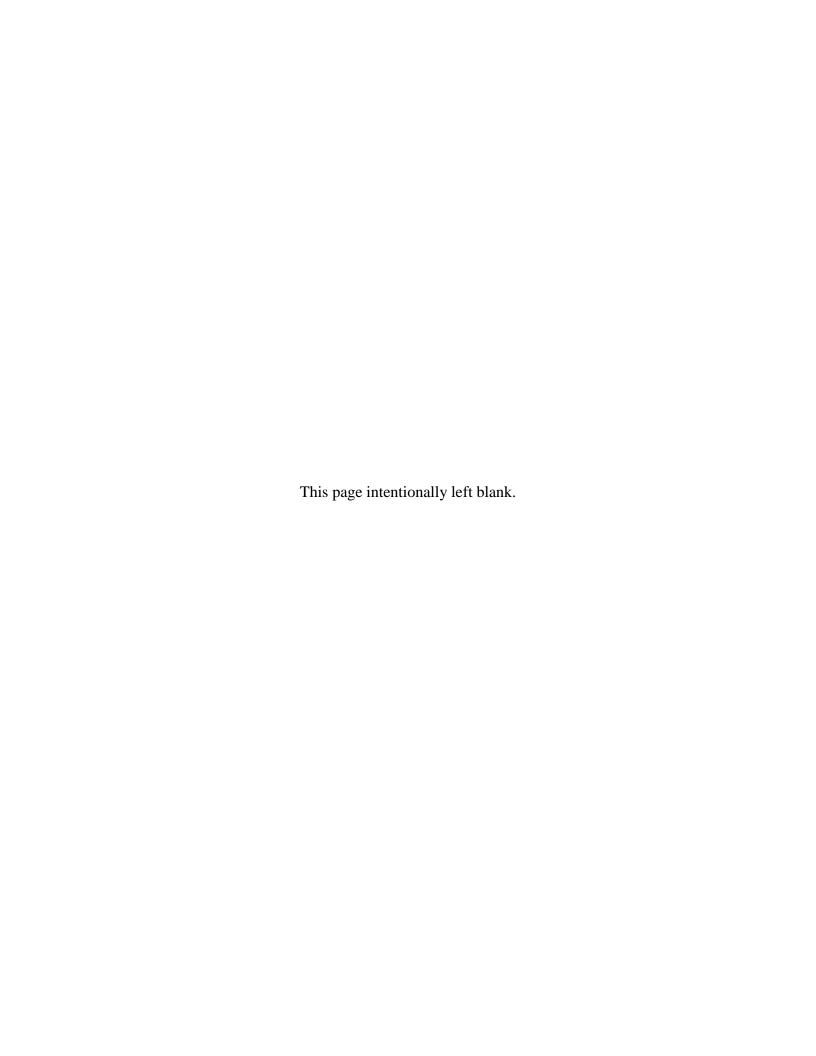


# Chapter 6 Potential Environmental Impact

**Final** 

**Supplemental Generic Environmental Impact Statement** 



# **Chapter 6 – Potential Environmental Impacts**

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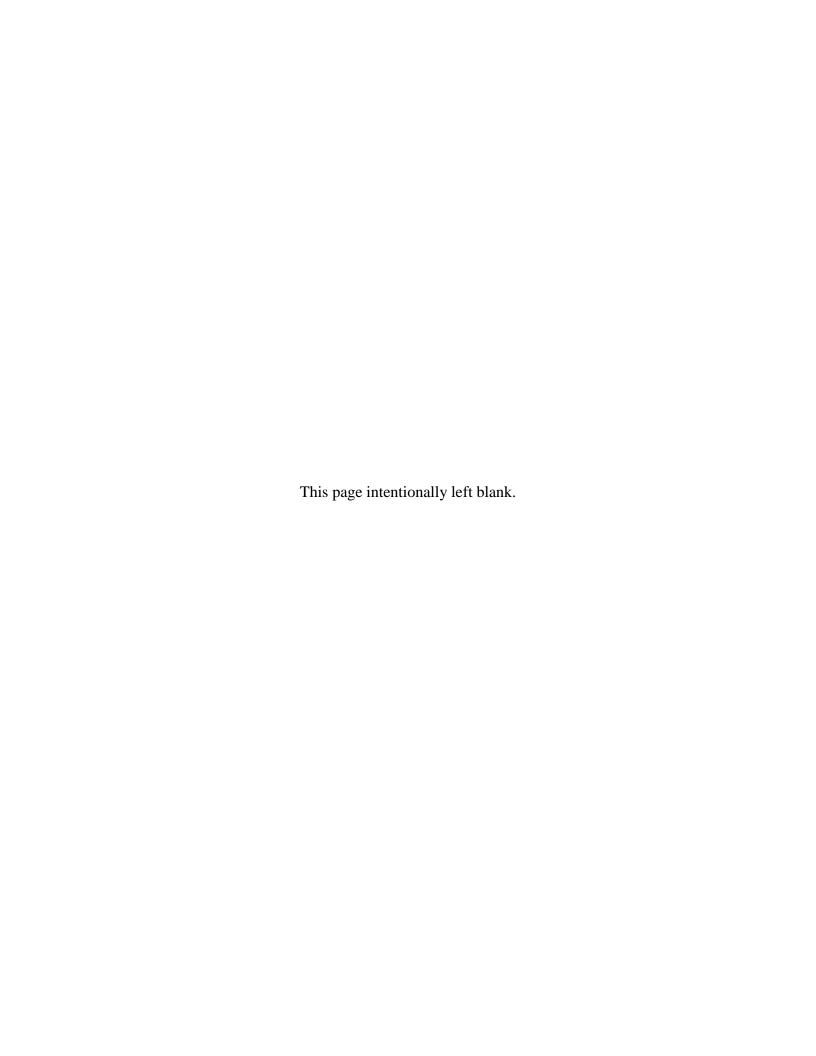
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# **Chapter 6 POTENTIAL ENVIRONMENTAL IMPACTS**

This revised Draft SGEIS incorporates by reference the 1992 Generic Environmental Impact Statement on the Oil, Gas and Solution Mining Regulatory Program - including the draft volumes released in 1988, the final volume released in 1992 - and the 1992 Findings Statement. Therefore, the text in this Supplement is not exhaustive with respect to potential environmental impacts, but instead focuses on new, different or additional information relating to potential impacts of horizontal drilling and high-volume hydraulic fracturing.

#### **6.1** Water Resources

Protection of water resources is a primary emphasis of the Department. Water resource matters that may be impacted by activities associated with high-volume hydraulic fracturing are identified and discussed in Chapter 2.

Adverse impacts to water resources might reasonably be anticipated in the context of unmitigated high-volume hydraulic fracturing due to: 1) water withdrawals affecting surface or groundwater, including wetlands; 2) polluted stormwater runoff; 3) surface chemical or petroleum spills; 4) pit or surface impoundment failures or leaks; 5) groundwater contamination associated with improper well drilling and construction; and 6) improper waste disposal. NYC's subsurface water supply infrastructure that is located in areas outside the boundary of the NYC Watershed could also be impacted by unmitigated high-volume hydraulic fracturing. Potential surface water impacts discussed herein are applicable to all areas that might be developed for natural gas resources through high-volume hydraulic fracturing.

Three water resources issues were the subject of extensive comment during the public scoping process:

- 1) Potential degradation of NYC's surface drinking water supply;
- 2) Potential groundwater contamination from the hydraulic fracturing procedure itself; and
- 3) Adverse impacts to the Upper Delaware Scenic and Recreational River.

Geological factors as well as standard permit requirements that the Department proposes to impose that would limit or avoid the potential for groundwater contamination from high-volume hydraulic fracturing are discussed in Chapters 5, 7 and 8.

#### 6.1.1 Water Withdrawals

Water for hydraulic fracturing may be obtained by withdrawing it from surface water bodies or new or existing water-supply wells drilled into aquifers. Without proper controls on the rate, timing and location of such withdrawals, modifications to groundwater levels, surface water levels, and stream flow could result in adverse impacts to aquatic ecosystems, downstream flow levels, drinking water assured yields, wetlands, and aquifer recharge. While surface-water bodies are still the primary source of water supplies for the drilling of Marcellus wells in Pennsylvania, municipal and public water-supply wells have been used there as well.

#### 6.1.1.1 Reduced Stream Flow

Potential effects of reduced stream flow caused by withdrawals could include:

- insufficient supplies for downstream uses such as public water supply;
- adverse impacts to quantity and quality of aquatic, wetland, and terrestrial habitats and the biota that they support; and
- exacerbation of drought effects.

Unmitigated withdrawals could adversely impact fish and wildlife health due to exposure to unsuitable water temperature and dissolved oxygen concentrations, particularly in low-flow or drought conditions. It could also affect downstream dischargers whose effluent limits are linked to the stream's flow rate. Water quality could be degraded and adverse impacts on natural aquatic habitat increased if existing pollutants from point sources (e.g., discharge pipes) and/or non-point sources (e.g., runoff from farms and paved surfaces) become concentrated.

# 6.1.1.2 Degradation of a Stream's Best Use

New York State water use classifications are provided in Section 2.3.1. All of the uses are dependent upon sufficient water in the stream to support the specified use. As noted, uncontrolled withdrawals of water from streams in connection with high-volume hydraulic

fracturing has the potential to adversely impact stream water supply and thus stream water use classifications.

# 6.1.1.3 Impacts to Aquatic Habitat

Habitat for stream organisms is provided by the shape of the stream channel and the water that flows through it. It is important to recognize that the physical habitat (e.g., pools, riffles, instream cover, runs, glides, bank cover, etc.) essential for maintaining the aquatic ecosystem is formed by periodic disturbances that exist in the natural hydrograph; the seasonal variability in stream flow resulting from annual precipitation and associated runoff. Maintaining this habitat diversity within a stream channel is essential in providing suitable conditions for all the life stage of the aquatic organisms. Stream fish distribution, community structure, and population dynamics are related to channel morphology. Streamflow alterations that modify channel morphology and habitat would result in changes in aquatic populations and community shifts that alter natural ecosystems. Creating and maintaining high quality habitat is a function of seasonally high flows because scour of fines from pools and deposition of bedload in riffles is most predominant at high flow associated with spring snowmelt or high rain runoff. Periodic resetting of the aquatic system is an essential process for maintaining stream habitat that would continuously provide suitable habitat for all aquatic biota. Clearly, alteration of flow regimes, sediment loads and riparian vegetation would cause changes in the morphology of stream channels. Any streamflow management decision would not impair flows necessary to maintain the dynamic nature of a river channel that is in a constant state of change as substrates are scoured, moved downstream and re-deposited.

# 6.1.1.4 Impacts to Aquatic Ecosystems

Aquatic ecosystems could be adversely impacted by:

- changes to water quality or quantity;
- insufficient stream flow for aquatic biota stream habitat; or
- the actual water withdrawal infrastructure.

Native aquatic species possess life history traits that enable individuals to survive and reproduce within a certain range of environmental variation. Flow depth and velocity, water temperature,

substrate size distribution and oxygen content are among the myriad of environmental attributes known to shape the habitat that control aquatic and riparian species distributions. Streamflow alterations can impact aquatic ecosystems due to community shifts made in response to the corresponding shifts in these environmental attributes. The perpetuation of native aquatic biodiversity and ecosystem integrity depends on maintaining some semblance of natural flow patterns that minimize aquatic community shifts. The natural flow paradigm states that the full range of natural intra- and inter-annual variation of hydrologic regimes, and associated characteristics of timing, duration, frequency and rate of change, are critical in sustaining the full native biodiversity and integrity of aquatic ecosystems.

Improperly installed water withdrawal structures can result in the entrainment of aquatic organisms, which can remove any/all life stages of fish and macroinvertebrates from their natural habitats as they are withdrawn with water. While most of the water bodies supplying water for high-volume hydraulic fracturing contain species of fish whose early life stages are not likely to be entrained because of their life history and behavioral characteristics, fish in their older life stages could be entrained without measures to avoid or reduce adverse impacts. To avoid adverse impacts to aquatic biota from entrainment, intake pipes can be screened to prevent entry into the pipe. Additionally, the loss of biota that becomes trapped on intake screens, referred to as impingement, can be minimized by properly sizing the intake to reduce the flow velocity through the screens. Depending on the water body from which water is being withdrawn, the location of the withdrawal structure on the water body and the site-specific aquatic organisms requiring protection, project-specific technologies may be required to minimize the entrainment and impingement of aquatic organisms. Technologies and operational measures that are proven effective in reducing these impacts include but are not limited to narrow-slot width wedge-wire screens (0.5 mm-2.0 mm), fine mesh screening, low intake velocities (0.5 feet per second (fps) or less), and seasonal restrictions on intake operation. Transporting water from the water withdrawal location for use off-site, as discussed in Section 6.4.2.2, can transfer invasive species from one water body to another via trucks, hoses, pipelines, and other equipment. Screening of the intakes can minimize this transfer; however, additional site-specific mitigation considerations may be necessary.

#### 6.1.1.5 Impacts to Wetlands

The existence and sustainability of wetland habitats directly depend on the presence of water at or near the surface of the soil. The functioning of a wetland is driven by the inflow and outflow of surface water and/or groundwater. As a result, withdrawal of surface water or groundwater for high-volume hydraulic fracturing could impact wetland resources. These potential impacts depend on the amount of water within the wetland, the amount of water withdrawn from the catchment area of the wetland, and the dynamics of water flowing into and out of the wetland. Even small changes in the hydrology of the wetland can have significant impacts on the wetland plant community and on the animals that depend on the wetland. It is important to preserve the hydrologic conditions and to understand the surface water and groundwater interaction to protect wetland areas.

# 6.1.1.6 Aquifer Depletion

The primary concern regarding groundwater withdrawal is aquifer depletion that could affect other uses, including nearby public and private water supply wells. This includes cumulative impacts from numerous groundwater withdrawals and potential aquifer depletion from the incremental increase in withdrawals if groundwater supplies are used for hydraulic fracturing. Aquifer depletion may also result in aquifer compaction which can result in localized ground subsidence. Aquifer depletion can occur in both confined and unconfined aquifers.

The depletion of an aquifer and a corresponding decline in the groundwater level can occur when a well, or wells in an aquifer are pumped at a rate in excess of the recharge rate to the aquifer. Essentially, surface water and groundwater are one continuous resource; therefore, it also is possible that aquifer depletion can occur if an excessive volume of water is removed from a surface water body that recharges an aquifer. Such an action would result in a reduction of recharge which could potentially deplete an aquifer. This "influent" condition of surface water recharging groundwater occurs mainly in arid and semi-arid climates, and is not common in New York, except under conditions such as induced infiltration of surface water by aquifer withdrawal (e.g., pumping of water wells). 268

<sup>&</sup>lt;sup>268</sup> Alpha, 2009, p. 3-19, with updates from DEC.

Aquifer depletion can lead to reduced discharge of groundwater to streams and lakes, reduced water availability in wetland areas, and corresponding impacts to aquatic organisms that depend on these habitats. Flowing rivers and streams are merely a surface manifestation of what is flowing through the shallow soils and rocks. Groundwater wells impact surface water flows by intercepting groundwater that otherwise would enter a stream. In fact, many New York headwater streams rely entirely on groundwater to provide flows in the hot summer months. It is therefore important to understand the hydrologic relationship between surface water, groundwater, and wetlands within a watershed to appropriately manage rates and quantities of water withdrawal.<sup>269</sup>

Depletion of both groundwater and surface water can occur when significant water withdrawals are transported out of the basin from which they originated. These transfers break the natural hydrologic cycle, since the transported water never makes it downstream nor returns to the original watershed to help recharge the aquifer. Without the natural flow regime, including seasonal high flows, stream channel and riparian habitats critical for maintaining the aquatic biota of the stream may be adversely impacted.

# 6.1.1.7 Cumulative Water Withdrawal Impacts<sup>270</sup>

As noted in later in this chapter, it is estimated that within 30 years there could be up to 40,000 wells developed with the high-volume hydraulic fracturing technology. This could result in substantial water usage in the study area. There are several potential types of impacts, when considered cumulatively, that could result from these estimated new withdrawals associated with natural gas development. Those are:

- Stream flow, surface water and groundwater depletion;
- Loss of aquifer storage capacity due to compaction;
- Water quality degradation;
- Wetland hydrology and habitat;

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<sup>&</sup>lt;sup>269</sup> Alpha, 2009, p. 81.

<sup>&</sup>lt;sup>270</sup> Alpha, 2009 pp. 3-28.

- Fish and aquatic organism impacts;
- Significant habitats, endangered, rare or threatened species impacts; and
- Existing water users and reliability of their supplies.

Evaluation of the overall impact of multiple water withdrawals based on the projection of maximum activity consider the existing water usage, the non-continuous nature of withdrawals for natural gas development, and the natural replenishment of water resources. Natural replenishment is described in Section 2.3.8.

The DRBC and SRBC have developed regulations, policies, and procedures to characterize existing water use and track approved withdrawals. Changes to these systems also require Commission review. Review of the requirements of the DRBC and SRBC indicates that the operators and the reviewing authority would perform evaluations to assess the potential impacts of water withdrawal for well drilling, and consider the following issues and information.

- Comprehensive project description that includes a description of the proposed water withdrawal (location, volume, and rate) and its intended use;
- Existing water use in the withdrawal area;
- Potential impacts, both ecological and to existing users, from the new withdrawal;
- Availability of water resources (surface water and/or groundwater) to support the proposed withdrawals;
- Availability of other water sources (e.g., treated waste water) and conservation plans to meet some or all of the water demand:
- Contingencies for low flow conditions that include passby flow criteria;
- Public notification requirements;
- Monitoring and reporting;
- Inspections;
- Mitigation measures;
- Supplemental investigations, including but not limited to, aquatic surveys;

- Potential impact to significant habitat and endangered rare or threatened species; and
- Protection of subsurface infrastructure.

#### Existing Regulatory Scheme for Water Usage and Withdrawals

The DRBC and SRBC use a permit system and approval process to regulate existing water usage in their respective basins. The DRBC and SRBC require applications in which operators provide a comprehensive project description that includes the description of the proposed withdrawals. The project information required includes site location, water source(s), withdrawal location(s), proposed timing and rate of water withdrawal and the anticipated project duration. The operators identify the amount of consumptive use (water not returned to the basin) and any import or export of water to or from the basin. The method of conveyance from the point(s) of withdrawal to the point(s) of use is also defined.

There are monitoring and reporting requirements once the withdrawal and consumptive use for a project has been approved. These requirements include metering withdrawals and consumptive use, and submitting quarterly reports to the Commission. Monitoring requirements can include stream flow and stage measurements for surface water withdrawals and monitoring groundwater levels for groundwater withdrawals.

The recently enacted Water Resources Law extends the Department's authority to regulate all water withdrawals over 100,000 gpd throughout all of New York State. This law applies to all such withdrawals where water would be used for high-volume hydraulic fracturing. Withdrawal permits issued in the future by the Department, pursuant to the regulations implementing this law, would include conditions to allow the Department to monitor and enforce water quality and quantity standards, and requirements. The Department is beginning the process for enacting regulations on this new law. These standards and requirements may include: passby flow; fish impingement and entrainment protections; protections for aquatic life; reasonable use; water conservation practices; and evaluation of cumulative impacts on other water withdrawals. The Department intends to seek consistency in water resource management within New York between the DRBC, SRBC and the Department.

Surface water and groundwater are withdrawn daily for a wide range of uses. New York ranks as one of the top states with respect to the total amount of water withdrawals. Figure 6.1 presents a graph indicating the total water withdrawal for New York is approximately 9 to 10 billion gpd, based on data from 2000. Figure 6.2 presents fresh water use in New York, including the projected peak water use for high-volume hydraulic fracturing.

The DRBC reports on the withdrawal of water for various purposes. The daily water withdrawals, exports, and consumptive uses in the Delaware River Basin are shown in Figure 6.3. The total water withdrawal from the Delaware River Basin was 8,736 MGD, based on 2003 water use records. The highest water use was for thermoelectric power generation at 5,682 million gpd (65%), followed by 875 million gpd (10%) for public water supply, 650 million gpd (7.4%) for the NYC public water supply, 617 million gpd (7%) for hydroelectric, and 501 million gpd (5.7%) for industrial purposes. The amount of water used for mining is 70 million gpd (0.8%). The "mining" category typically includes withdrawals for oil and gas drilling; however, DRBC has not yet approved water withdrawal for Marcellus Shale drilling operations. The information in Figure 6.3 shows that 4.3% (14 million gpd) of the water withdrawn for consumptive use is for mining and 88% (650 million gpd) of water exported from the Delaware River Basin is diverted to NYC.

Whereas certain withdrawals, like many public water supplies are returned to the basin's hydrologic cycle, out-of-basin transfers, like the NYC water-supply diversion, some evaporative losses, and withdrawals for hydraulic fracturing, are considered as 100% consumptive losses because this water is essentially lost to the basin's hydrologic cycle.

#### Withdrawals for High-Volume Hydraulic Fracturing

Current water withdrawal volumes when compared to withdrawal volumes associated with current natural gas drilling indicates that the historical percentage of withdrawn water that goes to natural gas drilling is very low. The amount of water withdrawn specifically for high-volume hydraulic fracturing also is projected to be relatively low when compared to existing overall levels of water use. The total volume of water withdrawn for high-volume hydraulic fracturing in New York would not be known with precision until applications are received, reviewed, and potentially approved or rejected by the appropriate regulatory agency or agencies, but can be

estimated based on activity in Pennsylvania and projections of potential levels of well drilling activity in New York.

Between July 2008 and February 2011, average water usage for high-volume hydraulic fracturing within the Susquehanna River Basin in Pennsylvania was 4.2 million gallons per well, based on data for 553 wells. 271 Current practice is to use 80% - 90% fresh water and 10% - 20% recycled flowback water for high-volume hydraulic fracturing. 272 Average fresh water use as 85% of the total used per well is consistent with statistics reported by the SRBC.<sup>273</sup> This would equate to average fresh water use of 3.6 million gallons per well (85% of 4.2 million gallons). Industry projects a potential peak annual drilling rate in New York of 2,462 wells, a level of drilling that is projected to be at the very high end of activity. Although some of these wells may be vertical wells which require less water than horizontal wells where high-volume hydraulic fracturing is planned, all of the wells reflected in the peak drilling rate will be conservatively considered to be horizontal wells for the purpose of this analysis. Multiplying the peak projected annual wells by current average use per well results in calculated peak annual fresh water usage for high-volume hydraulic fracturing of 9 billion gallons. Total daily fresh water withdrawal in New York has been estimated at approximately 10.3 billion gallons. <sup>274</sup> This equates to an annual total of about 3.8 trillion gallons. Based on this calculation, at peak activity high-volume hydraulic fracturing would result in increased demand for fresh water in New York of 0.24%. The potential relationship between water use for high-volume hydraulic fracturing and other purposes is shown in Figure 6.2.

While projected water withdrawals and consumptive use of water are modest relative to overall water withdrawals in New York, there remains the potential for adverse impacts particularly when withdrawals take place during low-flow or drought conditions. Adverse impacts previously discussed may also occur when high or unsustainable withdrawals take place in localized ground or surface water that lack adequate hydrologic capacity.

<sup>271</sup> SRBC 2011.

<sup>&</sup>lt;sup>272</sup> ALL Consulting, 2010, p. 74.

<sup>&</sup>lt;sup>273</sup> Richenderfer, 2010, p. 30.

<sup>&</sup>lt;sup>274</sup> Kenny et al, 2009, p.7.

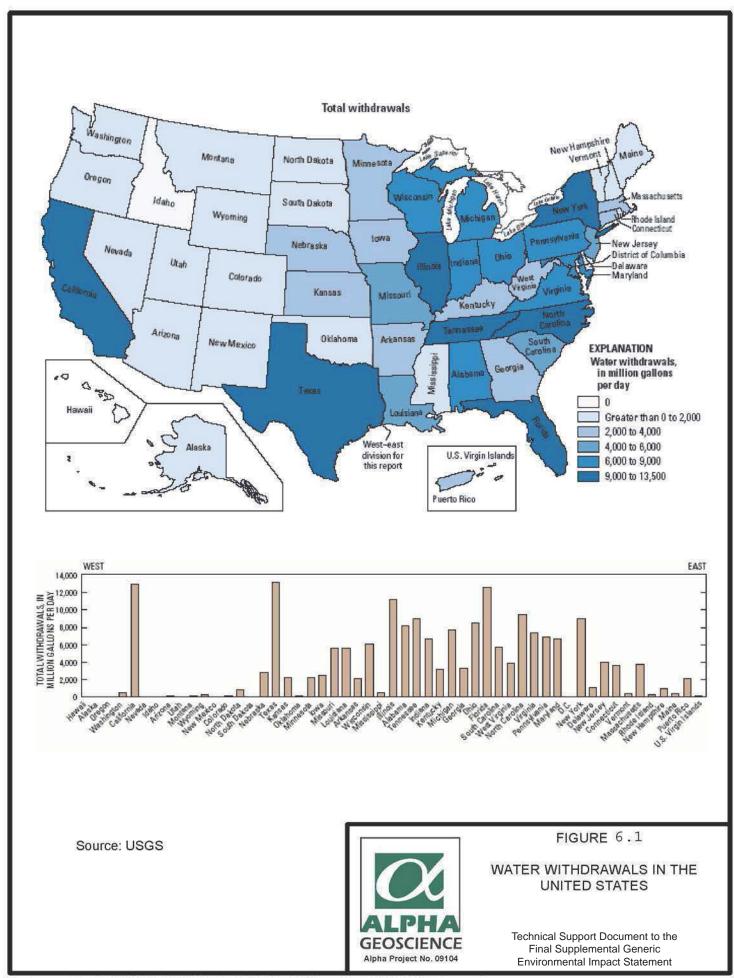
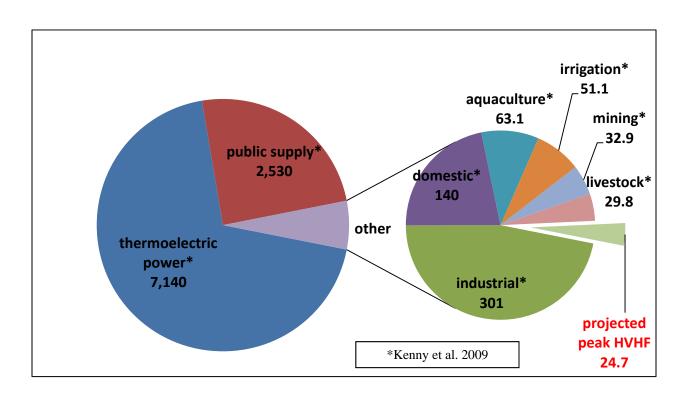
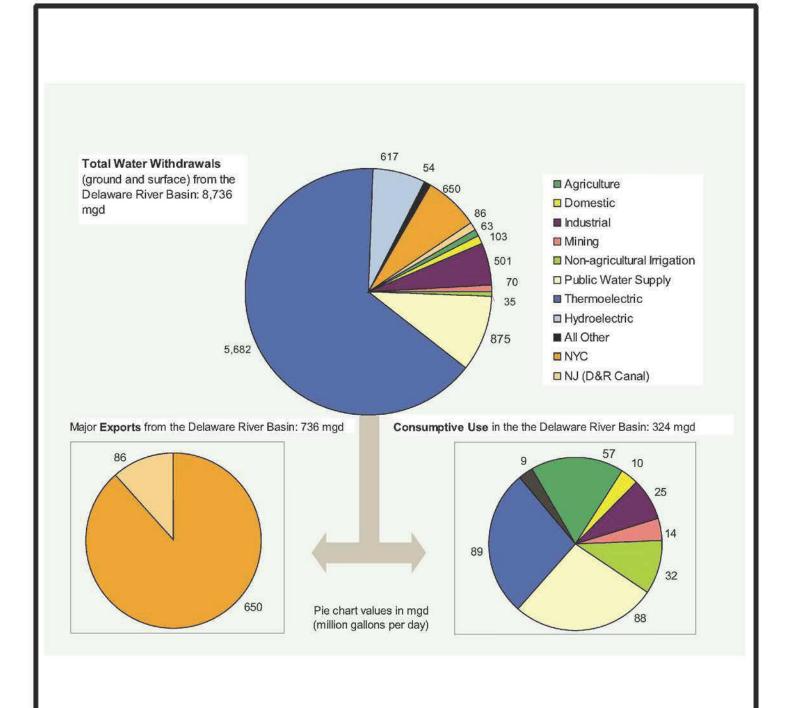


Figure 6.2 - Fresh Water Use in NY (millions of gallons per day) with Projected Peak Water Use for High-Volume Hydraulic Fracturing (New July 2011<sup>275</sup>)



<sup>&</sup>lt;sup>275</sup> This figure is a replacement for Figure 6.2 in the 2009 draft SGEIS which was a bar graph prepared by SRBC showing projected water use in the Susquehanna River Basin.



Source: DRBC



# FIGURE 6.3

DAILY WATER WITHDRAWALS, EXPORTS, AND CONSUMPTIVE USES IN THE DELAWARE RIVER BASIN

> Technical Support Document to the Final Supplemental Generic Environmental Impact Statement

#### 6.1.2 Stormwater Runoff

Stormwater, whether as a result of rainfall or snowmelt, is a valuable resource. It is the source of water for lakes and streams, as well as aquifers. However, stormwater runoff, particularly when it interacts with the human environment, is a pathway for contaminants to be conveyed from the land surface to streams and lakes and groundwater. This is especially true for stormwater runoff from asphalt, concrete, gravel/dirt roads, other impervious surfaces, outdoor industrial activity, and earthen construction sites, where any material collected on the ground is washed into a nearby surface water body. Stormwater runoff may also contribute to heightened peak flows and flooding.

On an undisturbed landscape, precipitation is held by vegetation and pervious soil, allowing it to slowly filter into the ground. This benefits water resources by using natural filtering properties, replenishing groundwater aquifers and feeding lakes and streams through base flow during dry periods. On a disturbed or developed landscape, it is common for the ground surface to be compacted or otherwise made less pervious and for runoff to be shunted away quickly with greater force and significantly higher volumes. Such hydrological modifications result in less groundwater recharge and more rapid runoff to streams, which may cause increased stream erosion and result in water quality degradation, habitat loss and flooding.

All phases of natural gas well development, from initial land clearing for access roads, equipment staging areas and well pads, to drilling and fracturing operations, production and final reclamation, have the potential to cause water resource impacts during rain and snow melt events if stormwater is not properly managed.

Excess sediment can fill or bury the rock cobble of streams that serve as spawning habitat for fish and the macro-invertebrate insects that serve as their food source. Stormwater runoff and heightened sediment loads carry excess levels of nutrient phosphorus and nitrogen that is a major cause of algae bloom, low dissolved oxygen and other water-quality impairments.

Initial land clearing exposes soil to erosion and more rapid runoff. Construction equipment is a potential source of contamination from such things as hydraulic, fuel and lubricating fluids. Equipment and any materials that are spilled, including additive chemicals and fuel, are exposed

to rainfall, so that contaminants may be conveyed off-site during rain events if they are not properly contained. Steep access roads, well pads on hill slopes, and well pads constructed by cut-and-fill operations pose particular challenges, especially if an on-site drilling pit is proposed.

A production site, including access roads, is also a potential source of stormwater runoff impacts discussed above because its hydrologic characteristics, sediment, nutrient, contaminant, and water volumes may be substantially different from the pre-developed condition.

# 6.1.3 Surface Spills and Releases at the Well Pad

Spills or releases can occur as a result of tank ruptures, piping failures, equipment or surface impoundment failures, overfills, vandalism, accidents (including vehicle collisions), ground fires, drilling and production equipment defects, or improper operations. Spilled, leaked or released fluids could flow to a surface water body or infiltrate the ground, reaching subsurface soils and aquifers.

To evaluate potential health impacts from spills or releases of additives, fracturing fluid containing diluted additives or residual diluted additive chemicals in flowback water, the NYSDOH reviewed the composition of additives proposed for high-volume hydraulic fracturing in New York. The NYSDOH concluded that the proposed additives contain similar types of chemical constituents as the products that have been used for many years for hydraulic fracturing of traditional vertical wells in NYS. Some of the same products are used in both well types. The total amount of fracturing additives and water used in hydraulic fracturing of horizontal wells is considerably larger than for traditional vertical wells. This suggests the potential environmental consequences of an upset condition could be proportionally larger for horizontal well drilling and fracturing operations. As mentioned earlier, the 1992 GEIS addressed hydraulic fracturing in Chapter 9, and NYSDOH's review did not identify any potential exposure situations associated with horizontal drilling and high-volume hydraulic fracturing that are qualitatively different from those addressed in the 1992 GEIS.

#### 6.1.3.1 Drilling

Contamination of surface water bodies and groundwater resources during well drilling could occur as a result of failure to maintain stormwater controls, ineffective site management and inadequate surface and subsurface fluid containment practices, poor casing construction, or accidental spills and releases including well blow-outs during drilling or well component failures during completion operations. A release could also occur during a blow-out event if there are not trained personnel on site that are educated in the proper use of the BOP system. Surface spills would involve materials and fluids present at the site during the drilling phase. Pit leakage or failure could also involve well fluids. These issues are discussed in Chapters 8 and 9 of the 1992 GEIS, but are acknowledged here with respect to unique aspects of the proposed multi-well development method. The conclusions regarding pit construction standards and liner specifications presented in the 1992 GEIS were largely based upon the short duration of a pit's use. The greater intensity and duration of surface activities associated with well pads with multiple wells increases the potential for an accidental spill, pit leak or pit failure if engineering controls and other mitigation measures are not sufficient. Concerns are heightened if on-site pits for handling drilling fluids are located in primary and principal aquifer areas, or are constructed on the filled portion of a cut-and-filled well pad.

# 6.1.3.2 Hydraulic Fracturing Additives

As with the drilling phase, contamination of surface water bodies and groundwater resources during well stimulation could occur as a result of failure to maintain stormwater controls, ineffective site management and surface and subsurface fluid containment practices, poor well construction and grouting, or accidental spills and releases including failure of wellhead components during hydraulic fracturing. These issues are discussed in Chapters 8 and 9 of the 1992 GEIS, but are acknowledged here because of the larger volumes of fluids and materials to be managed for high-volume hydraulic fracturing. The potential contaminants are listed in Table 5.7 and grouped into categories recommended by NYSDOH in Table 5.8. URS compared the list of additive chemicals to the parameters regulated via federal and state primary or secondary drinking water standards, SPDES discharge limits (see Section 7.1.8), and DOW Technical and Operational Guidance Series 1.1.1 (TOGS111), Ambient Water Quality Standards and Guidance

*Values and Groundwater Effluent Limitations*. <sup>276,277</sup> In NYS, the state drinking water standards (10 NYCRR 5) apply to all public water supplies and set maximum contaminant levels (MCLs) for essentially all organic chemicals in public drinking water. See Table 6.1.

#### 6.1.3.3 Flowback Water and Production Brine

Gelling agents, surfactants and chlorides are identified in the 1992 GEIS as the flowback water components of greatest environmental concern. Other flowback components can include other dissolved solids, metals, biocides, lubricants, organics and radionuclides. Opportunities for spills, leaks, and operational errors during the flowback water recovery stage are the same as they are during the prior stages with additional potential releases from:

- hoses or pipes used to convey flowback water to tanks or a tanker truck for transportation to a treatment or disposal site; and
- tank leakage.

In general, *flowback water* is water and associated chemical constituents returning from the borehole during or proximate in time to hydraulic fracturing activities. *Production brine*, on the other hand, is fluid that returns from the borehole after completion of drilling operations while natural gas production is underway. The chemical characteristics and volumes of flowback water and production brine are expected to differ in significant respects.

Flowback water composition based on a limited number of out-of-state samples from Marcellus wells is presented in Table 5.9. A comparison of detected flowback parameters, except radionuclides, to regulated parameters is presented in Table 6.1.<sup>279</sup>

Table 5.10 lists parameters found in the flowback analyses, except radionuclides, that are regulated in New York. The number of samples that were analyzed for the particular parameter is shown in Column 3, and the number of samples in which parameters were detected is shown in Column 4. The minimum, median and maximum concentrations detected are indicated in

http://www.dec.ny.gov/regulations/2652.html.

<sup>&</sup>lt;sup>276</sup> URS, 2009, p. 4-18, et seq.

<sup>&</sup>lt;sup>278</sup> NYSDEC, 1992, GEIS, p. 9-37.

<sup>&</sup>lt;sup>279</sup> URS, 2009, p. 4-18, et seq.

Columns 5, 6 and 7. Radionuclides data is presented in Chapter 5, and potential impacts and regulation are discussed in Section 6.7.

Table 5.11 lists parameters found in the flowback analyses that are not regulated in New York. Column 2 shows the number of samples that were analyzed for the particular parameter; column 3 indicates the number of samples in which the parameter was detected.<sup>281</sup>

Information presented in Tables 5.10 and 5.11 are based on limited data from Pennsylvania and West Virginia. Samples were not collected specifically for this type of analysis or under the Department's oversight. Characteristics of flowback from the Marcellus Shale in New York are expected to be similar to flowback from Pennsylvania and West Virginia, but not identical. The raw data for these tables came from several sources, with likely varying degrees of reliability, and the analytical methods used were not all the same for given parameters. Sometimes, laboratories need to use different analytical methods depending on the consistency and quality of the sample; sometimes the laboratories are only required to provide a certain level of accuracy. Therefore, the method detection limits may be different. The quality and composition of flowback from a single well can also change within a few days after the well is fractured. This data does not control for any of these variables. <sup>282</sup>

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<sup>&</sup>lt;sup>280</sup> URS, 2009, pp. 4-10, 4-31 et seq.

<sup>&</sup>lt;sup>281</sup> URS, 2009, pp. 4-10, p. 4-35.

<sup>&</sup>lt;sup>282</sup> URS, 2009, p. 4-31.

Table 6.1 - Comparison of additives used or proposed for use in NY, parameters detected in analytical results of flowback from the Marcellus operations in PA and WV and parameters regulated via primary and secondary drinking water standards, SPDES or TOGS111 (Revised August 2011)<sup>283, 284</sup>

| CAS<br>Number | Parameter Name                      | Used in<br>Additives <sup>285</sup> ,<br>286 | Found in<br>Flowback<br>287 | USEPA<br>MCL or TT<br>(mg/L) <sup>288</sup> ,<br>289 | SPDES<br>Tables 290 | TOGS111<br>Tables | NYS MCL,<br>(mg/L) <sup>291</sup> |
|---------------|-------------------------------------|--|-----------------------------|--|---------------------|-------------------|-----------------------------------|
| 106-24-1      | (2E)-3,7-dimethylocta-2,6-dien-1-ol | Yes  |                             |  |                     |                   | 0.05                              |

Table 6.1 was compiled by URS Corporation, 2011 and revised by the Department in coordination with NYSDOH.

<sup>&</sup>lt;sup>284</sup> This table includes parameters detected in the MSC Study.

<sup>&</sup>lt;sup>285</sup> Information in the "Used in Additives" column is based on the composition of additives used or proposed for use in New York.

<sup>&</sup>lt;sup>286</sup> Parameters marked with ¥ indicates that the compound dissociates, and its components are separately regulated. Not all dissociating compounds are marked.

Information in the "Found in Flowback" column is based on analytical results of flowback from operations in Pennsylvania or West Virginia. There are/may be products used in fracturing operations in Pennsylvania that have not yet been proposed for use in New York for which, therefore, the Department does not have chemical composition data. Blank entries in the "Found in Flowback" column indicate that the parameter was either not sampled for or not detected in the flowback.

USEPA Maximum Contaminant Level (MCL) - The highest level of a contaminant that is allowed in drinking water. MCLs are set as close to MCLGs as feasible using the best available treatment technology and taking cost into consideration. MCLs are enforceable standards. From USEPA Title 40, Part 141--National Primary Drinking Water Regulations.

USEPA Treatment Technique (TT) – A required process intended to reduce the level of a contaminant in drinking water. From USEPA Title 40, Part 141 – National Primary Drinking Water Regulations.

<sup>290</sup> SPDES or TOGS typically regulates or provides guidance for the total substance, (e.g., iron) and rarely regulates or provides guidance for only its dissolved portion (e.g., dissolved iron). The dissolved component is implicitly covered in the total substance. Therefore, the dissolved component is not included in this table. Flowback analyses provided information for the total and dissolved components of metals. Understanding the dissolved vs. suspended portions of a substance is valuable when determining potential treatment techniques.

<sup>&</sup>lt;sup>291</sup> 10 NYCRR Part 5-1.50 through 5-1.52. Under 10 NYCRR Part 5, organic contaminants (with very few exceptions) have either a Specific MCL (28 compounds plus 1 chemical mixture) or a General MCL of 0.05 mg/L for Unspecified Organic Contaminants (UOC) or 0.005 mg/L for Principal Organic Contaminants (POC). A total UOC + POC MCL of 0.1 mg/L also applies to all organic contaminants in drinking water. 10 NYCRR Part 5 also contains 23 MCLs for inorganic contaminants. A section sign (§) indicates that, for organic salts, the free compound (the expected form in drinking water) would be a UOC, but that salts themselves would not be UOC. A double section sign (§§) indicates that, for parameters listed as a group or mixture of related chemicals (e.g., Ethoxylated alcohol (C14-15), petroleum distillates, essential oils) a state MCL does not apply to the group as a whole, but would apply to each individual component of the group if detected in drinking water. A triple section sign (§§§) indicates that, for parameters listed as a polymer, the UOC MCL would apply to the polymer itself, but either the UOC or POC MCL would apply to the individual monomer components. An asterisk (\*) indicates that the total trihalomethane (THM) MCL of 0.08 mg/L also applies.

| CAS<br>Number | Parameter Name  | Used in<br>Additives <sup>285</sup> ,<br>286 | Found in<br>Flowback<br>287 | USEPA<br>MCL or TT<br>(mg/L) <sup>288,</sup><br>289 | SPDES<br>Tables <sup>290</sup> | TOGS111<br>Tables | NYS MCL,<br>(mg/L) <sup>291</sup> |
|---------------|---|--|-----------------------------|---|--------------------------------|-------------------|-----------------------------------|
| 67701-10-4    | (C8-C18) And (C18) Unsaturated<br>Alkylcarboxylic Acid Sodium Salt                        | Yes  |                             |   |                                |                   | §,§§                              |
| 02634-33-5    | 1,2 Benzisothiazolin-2-one / 1,2-<br>benzisothiazolin-3-one                               | Yes  |                             |   |                                |                   | 0.05                              |
| 00087-61-6    | 1,2,3-Trichlorobenzene  |  | Yes                         |   | Table 9                        | Tables 1,5        | 0.005                             |
| 00095-63-6    |   | Yes  | Yes                         |   | Table 9                        | Tables 1,5        | 0.005                             |
| 93858-78-7    | 1,2,4-Butanetricarboxylicacid, 2-phosphono-, potassium salt                               | Yes  |                             |   |                                |                   | 0.05                              |
| 00108-67-8    | 1,3,5-Trimethylbenzene  |  | Yes                         |   | Tables 9,10                    | Tables 1,5        | 0.005                             |
| 00123-91-1    | 1,4 Dioxane   | Yes  |                             |   | Table 8                        |                   | 0.05                              |
| 03452-07-1    | 1-eicosene  | Yes  |                             |   |                                |                   | 0.05                              |
| 00629-73-2    | 1-hexadecene  | Yes  |                             |   |                                |                   | 0.05                              |
| 104-46-1      | 1-Methoxy-4-propenylbenzene   | Yes  |                             |   |                                |                   | 0.05                              |
| 124-28-7      | 1-Octadecanamine, N, N-dimethyl- / N,N-<br>Timethyloctadecylamine                         | Yes  |                             |   |                                |                   | 0.05                              |
| 112-03-8      | 1-Octadecanaminium, N,N,N-Trimethyl-,<br>Chloride /Trimethyloctadecylammonium<br>chloride | Yes  |                             |   |                                |                   | 0.05                              |
| 00112-88-9    | 1-octadecene  | Yes  |                             |   |                                |                   | 0.05                              |
| 40623-73-2    | 1-Propanesulfonic acid  | Yes  |                             |   |                                |                   | 0.05                              |
| 01120-36-1    | 1-tetradecene   | Yes  |                             |   |                                |                   | 0.05                              |
| 98-55-5       | 2-(4-methyl-1-cyclohex-3-enyl)propan-2-ol   | Yes  |                             |   |                                |                   | 0.05                              |
| 10222-01-2    | 2,2 Dibromo-3-nitrilopropionamide   | Yes  |                             |   | Table 9                        | Tables 1,5        |                                   |
| 27776-21-2    | 2,2'-azobis-{2-(imidazlin-2-yl)propane}-dihydrochloride                                   | Yes  |                             |   |                                |                   | 0.05                              |
| 73003-80-2    | 2,2-Dibromomalonamide   | Yes  |                             |   |                                |                   | 0.05                              |
| 00105-67-9    | 2,4-Dimethylphenol  |  | Yes                         |   | Table 6                        | Tables 1,5        | 0.05                              |
| 00087-65-0    | 2,6-Dichlorophenol  |  | Yes                         |   | Table 8                        |                   | 0.005                             |
| 15214-89-8    | 2-Acrylamido-2-methylpropanesulphonic acid sodium salt polymer                            | Yes  |                             |   |                                |                   | 0.05                              |
| 46830-22-2    | 2-acryloyloxyethyl(benzyl)dimethylammonium chloride                                       | Yes  |                             |   |                                |                   | 0.05                              |
| 00052-51-7    | 2-Bromo-2-nitro-1,3-propanediol   | Yes  |                             |   | Table 10                       |                   |                                   |
| 00111-76-2    | 2-Butoxy ethanol /Ethylene glycol monobutyl ether / Butyl Cellusolve                      | Yes  |                             |   |                                |                   | 0.05                              |

| CAS<br>Number | Parameter Name   | Used in<br>Additives <sup>285</sup> ,<br>286 | Found in<br>Flowback<br>287 | USEPA<br>MCL or TT<br>(mg/L) <sup>288,</sup><br>289 | SPDES<br>Tables <sup>290</sup> | TOGS111<br>Tables | NYS MCL, (mg/L) <sup>291</sup> |
|---------------|--|--|-----------------------------|---|--------------------------------|-------------------|--------------------------------|
| 01113-55-9    | 2-Dibromo-3-Nitriloprionamide / 2-   | Yes  |                             |   |                                |                   | 0.05                           |
|               | Monobromo-3-nitrilopropionamide  | ies  |                             |   |                                |                   | 0.03                           |
| 00104-76-7    | 2-Ethyl Hexanol  | Yes  |                             |   |                                |                   | 0.05                           |
| 00091-57-6    | 2-Methylnaphthalene  |  | Yes                         |   | Table 8                        | Tables 1,3        | 0.05                           |
| 00095-48-7    | 2-Methylphenol   |  | Yes                         |   | Table 8                        |                   | 0.05                           |
| 109-06-8      | 2-Picoline (2-methyl pyridine)   |  | Yes                         |   | Table 8                        | Table 3           | 0.05                           |
| 00067-63-0    | 2-Propanol / Isopropyl Alcohol / Isopropanol /<br>Propan-2-ol  | Yes  | Yes                         |   | Table 10                       |                   | 0.05                           |
| 26062-79-3    | 2-Propen-1-aminium, N,N-dimethyl-N-2-<br>propenyl-chloride, homopolymer  | Yes  |                             |   |                                |                   | 0.05                           |
| 95077-68-2    | 2-Propenoic acid, homopolymer sodium salt  | Yes  |                             |   |                                |                   | 0.05                           |
| 09003-03-6    | 2-propenoic acid, homopolymer, ammonium salt   | Yes  |                             |   |                                |                   | 0.05                           |
| 25987-30-8    | 2-Propenoic acid, polymer with 2 p-<br>propenamide, sodium salt / Copolymer of<br>acrylamide and sodium acrylate | Yes  |                             |   |                                |                   | 0.05                           |
| 71050-62-9    | 2-Propenoic acid, polymer with sodium phosphinate (1:1)  | Yes  |                             |   |                                |                   | 0.05                           |
| 66019-18-9    | 2-propenoic acid, telomer with sodium hydrogen sulfite   | Yes  |                             |   |                                |                   | 0.05                           |
| 00107-19-7    | 2-Propyn-1-ol / Progargyl Alcohol  | Yes  |                             |   |                                |                   | 0.05                           |
| 51229-78-8    | 3,5,7-Triaza-1-<br>azoniatricyclo[3.3.1.13,7]decane, 1-(3-chloro-<br>2-propenyl)-chloride,                       | Yes  |                             |   |                                |                   | 0.05                           |
| 106-22-9      | 3,7 - dimethyl-6-octen-1-ol  | Yes  |                             |   |                                |                   | 0.05                           |
| 5392-40-5     | 3,7-dimethyl-2,6-octadienal  | Yes  |                             |   |                                |                   | 0.005                          |
| 00115-19-5    | 3-methyl-1-butyn-3-ol  | Yes  |                             |   |                                |                   | 0.05                           |
| 00108-39-4    | 3-Methylphenol   |  | Yes                         |   | Table 8                        |                   | 0.05                           |
| 104-55-2      | 3-phenyl-2-propenal  | Yes  |                             |   |                                |                   | 0.005                          |
| 127-41-3      | 4-(2,6,6-trimethyl-1-cyclohex-2-enyl)-3-buten-<br>2-one  | Yes  |                             |   |                                |                   | 0.05                           |
| 00072-55-9    | 4,4 DDE  |  | Yes                         |   | Table 6                        | Tables 1,5        | 0.005                          |
| 121-33-5      | 4-hydroxy-3-methoxybenzaldehyde  | Yes  |                             |   |                                |                   | 0.05                           |
| 00106-44-5    | 4-Methylphenol   |  | Yes                         |   | Table 8                        |                   | 0.05                           |
| 127087-87-0   | 4-Nonylphenol Polyethylene Glycol Ether<br>Branched / Nonylphenol ethoxylated /<br>Oxyalkylated Phenol           | Yes  |                             |   |                                |                   | 0.05                           |

| CAS<br>Number | Parameter Name   | Used in<br>Additives <sup>285</sup> ,<br>286 | Found in<br>Flowback<br>287 | USEPA<br>MCL or TT<br>(mg/L) <sup>288,</sup><br>289 | SPDES<br>Tables <sup>290</sup> | TOGS111<br>Tables | NYS MCL,<br>(mg/L) <sup>291</sup> |
|---------------|--|--|-----------------------------|---|--------------------------------|-------------------|-----------------------------------|
| 00057-97-6    | 7,12-Dimethylbenz(a)anthracene   |  | Yes                         |   | Table 8                        | Table 3           | 0.05                              |
| 00064-19-7    | Acetic acid  | Yes  | Yes                         |   | Table 10                       |                   | 0.05                              |
| 68442-62-6    | Acetic acid, hydroxy-, reaction products with triethanolamine  | Yes  |                             |   |                                |                   | 0.05                              |
| 00108-24-7    | Acetic Anhydride   | Yes  |                             |   | Table 10                       |                   | 0.05                              |
| 00067-64-1    | Acetone  | Yes  | Yes                         |   | Table 7                        | Tables 1,5        | 0.05                              |
| 00098-86-2    | Acetophenone   |  | Yes                         |   |                                | Table 3           | 0.05                              |
| 00079-06-1    | Acrylamide   | Yes  |                             | TT  | Table 9                        | Tables 1,5        | 0.005                             |
| 38193-60-1    | Acrylamide - sodium 2-acrylamido-2-<br>methylpropane sulfonate copolymer   | Yes  |                             |   |                                |                   | 0.05                              |
| 25085-02-3    | Acrylamide - Sodium Acrylate Copolymer or<br>Anionic Polyacrylamide  | Yes  |                             |   |                                |                   | 0.05                              |
| 69418-26-4    | Acrylamide polymer with N,N,N-trimethyl-<br>2[1-oxo-2-propenyl]oxy Ethanaminium<br>chloride  | Yes  |                             |   |                                |                   | 0.05                              |
| 15085-02-3    | Acrylamide-sodium acrylate copolymer   | Yes  |                             |   |                                |                   | 0.05                              |
| 00107-13-1    | Acrylonitrile  |  | Yes                         |   | Table 6                        | Tables 1,5        |                                   |
| 68891-29-2    | Alcohols C8-10, ethoxylated, monoether with sulfuric acid, ammonium salt   | Yes  |                             |   |                                |                   | §,§§                              |
| 68526-86-3    | Alcohols, C11-14-iso-, C13-rich  | Yes  |                             |   |                                |                   | §§                                |
| 68551-12-2    | Alcohols, C12-C16, Ethoxylated (a.k.a. Ethoxylated alcohol)  | Yes  |                             |   |                                |                   | §§                                |
| 00309-00-2    | Aldrin   |  | Yes                         |   |                                | Tables 1,5        |                                   |
|               | Aliphatic acids  | Yes  |                             |   |                                |                   | §§                                |
|               | Aliphatic alcohol glycol ether   | Yes  |                             |   |                                |                   | 0.05                              |
| 64742-47-8    | Aliphatic Hydrocarbon / Hydrotreated light<br>distillate / Petroleum Distillates / Isoparaffinic<br>Solvent / Paraffin Solvent / Napthenic Solvent | Yes  |                             |   |                                |                   | §§                                |
|               | Alkalinity, Carbonate, as CaCO <sub>3</sub>  |  | Yes                         |   | Table 10                       |                   |                                   |
| 64743-02-8    | Alkenes  | Yes  |                             |   |                                |                   | §§                                |
| 68439-57-6    | Alkyl (C14-C16) olefin sulfonate, sodium salt  | Yes  |                             |   |                                |                   | 0.05                              |
|               | Alkyl Aryl Polyethoxy Ethanol  | Yes  |                             |   |                                |                   | 0.05                              |
|               | Alkylaryl Sulfonate  | Yes  |                             |   |                                |                   | 0.05                              |
| 09016-45-9    | Alkylphenol ethoxylate surfactants   | Yes  |                             |   |                                |                   | <b>§</b> §                        |
| 07439-90-5    | Aluminum   |  | Yes                         |   | Table 7                        | Tables 1,5        |                                   |
| 01327-41-9    | Aluminum chloride  | Yes (¥)                                      |                             |   |                                |                   |                                   |

| CAS<br>Number | Parameter Name   | Used in<br>Additives <sup>285,</sup><br>286 | Found in<br>Flowback<br>287 | USEPA<br>MCL or TT<br>(mg/L) <sup>288,</sup><br>289 | SPDES<br>Tables <sup>290</sup> | TOGS111<br>Tables | NYS MCL,<br>(mg/L) <sup>291</sup> |
|---------------|--|---|-----------------------------|---|--------------------------------|-------------------|-----------------------------------|
| 68155-07-7    | Amides, C8-18 and C19-Unsatd., N,N-Bis(hydroxyethyl)   | Yes   |                             |   |                                |                   | §§                                |
| 73138-27-9    | Amines, C12-14-tert-alkyl, ethoxylated   | Yes   |                             |   |                                |                   | <b>§</b> §                        |
| 71011-04-6    | Amines, Ditallow alkyl, ethoxylated  | Yes   |                             |   |                                |                   | §§                                |
| 68551-33-7    | Amines, tallow alkyl, ethoxylated, acetates  | Yes   |                             |   |                                |                   | §§                                |
| 01336-21-6    | Ammonia  | Yes   |                             |   | Yes                            |                   | 0.0                               |
| 00631-61-8    | Ammonium acetate   | Yes   |                             |   | Table 10                       |                   | §                                 |
| 68037-05-8    | Ammonium Alcohol Ether Sulfate   | Yes (¥)                                     |                             |   |                                |                   | 0.05                              |
| 07783-20-2    | Ammonium bisulfate   | Yes (¥)                                     |                             |   |                                |                   |                                   |
| 10192-30-0    | Ammonium Bisulphite  | Yes (¥)                                     |                             |   |                                |                   |                                   |
| 12125-02-9    | Ammonium Chloride  | Yes (¥)                                     |                             |   | Table 10                       |                   |                                   |
| 07632-50-0    | Ammonium citrate   | Yes (¥)                                     |                             |   |                                |                   | §                                 |
| 37475-88-0    | Ammonium Cumene Sulfonate  | Yes (¥)                                     |                             |   |                                |                   | §                                 |
| 01341-49-7    | Ammonium hydrogen-difluoride   | Yes (¥)                                     |                             |   |                                |                   |                                   |
| 06484-52-2    | Ammonium nitrate   | Yes (¥)                                     |                             |   |                                |                   |                                   |
| 07727-54-0    | Ammonium Persulfate / Diammonium peroxidisulphate  | Yes (¥)                                     |                             |   |                                |                   |                                   |
| 01762-95-4    | Ammonium Thiocyanate   | Yes   |                             |   | Table 10                       |                   |                                   |
|               | Anionic copolymer  | Yes   |                             |   |                                |                   |                                   |
| 07440-36-0    | Antimony   |   | Yes                         | 0.006   | Table 6                        | Tables 1,5        | 0.006                             |
| 07664-41-7    | Aqueous ammonia  | Yes   | Yes                         |   | Table 7                        | Tables 1,5        |                                   |
| 12672-29-6    | Aroclor 1248   |   | Yes                         |   | Table 6                        |                   | 0.0005                            |
|               | Aromatic hydrocarbons  | Yes   |                             |   |                                |                   | §§                                |
|               | Aromatic ketones   | Yes   |                             |   |                                |                   | <b>§</b> §                        |
| 07440-38-2    | Arsenic  |   | Yes                         | 0.01  | Table 6                        | Tables 1,5        | 0.01                              |
| 12174-11-7    | Attapulgite Clay   | Yes   |                             |   |                                |                   |                                   |
| 07440-39-3    | Barium   |   | Yes                         | 2   | Table 7                        | Tables 1,5        | 2                                 |
|               | Barium Strontium P.S. (mg/L)   |   | Yes                         |   |                                |                   |                                   |
| 121888-68-4   | Bentonite, benzyl(hydrogenated tallow alkyl)<br>dimethylammonium stearate complex /<br>organophilic clay | Yes   |                             |   |                                |                   |                                   |
| 00071-43-2    | Benzene  | Yes   | Yes                         | 0.005   | Table 6                        | Tables 1,5        | 0.005                             |
| 119345-04-9   | Benzene, 1,1'-oxybis, tetratpropylene derivatives, sulfonated, sodium salts                              | Yes   |                             |   |                                |                   | 0.05                              |

| CAS<br>Number | Parameter Name  | Used in<br>Additives <sup>285,</sup><br>286 | Found in<br>Flowback<br>287 | USEPA<br>MCL or TT<br>(mg/L) <sup>288,</sup><br>289 | SPDES<br>Tables <sup>290</sup> | TOGS111<br>Tables | NYS MCL, (mg/L) <sup>291</sup> |
|---------------|---|---|-----------------------------|---|--------------------------------|-------------------|--------------------------------|
| 74153-51-8    | Benzenemethanaminium, N,N-dimethyl-N-[2-<br>[(1-oxo-2-propenyl)oxy]ethyl]-, chloride,<br>polymer with 2-propenamide | Yes   |                             |   |                                |                   | 0.05                           |
| 122-91-8      | Benzenemethanol,4-methoxy-, 1-formate   | Yes   |                             |   |                                |                   | 0.05                           |
| 1300-72-7     | Benzenesulfonic acid, Dimethyl-, Sodium salt (aka Sodium xylene sulfonate)  | Yes   |                             |   |                                |                   | 0.05                           |
| 00050-32-8    | Benzo(a)pyrene  |   | Yes                         |   | Table 6                        |                   | 0.0002                         |
| 00205-99-2    | Benzo(b)fluoranthene  |   | Yes                         |   |                                | Tables 1,5        | 0.05                           |
| 00191-24-2    | Benzo(ghi)perylene  |   | Yes                         |   | Table 6                        | Table 3           | 0.05                           |
| 00207-08-9    | Benzo(k)fluoranthene  |   | Yes                         |   | Table 6                        | Tables 1,5        | 0.05                           |
| 140-11-4      | Benzyl acetate  | Yes   |                             |   |                                |                   | 0.05                           |
| 00100-51-6    | Benzyl alcohol  |   | Yes                         |   | Table 8                        | Table 3           | 0.05                           |
| 07440-41-7    | Beryllium   |   | Yes                         | 0.004   | Table 6                        | Tables 1,5        | 0.004                          |
|               | Bicarbonates (mg/L)   |   | Yes                         |   | Table 10                       |                   |                                |
| 76-22-2       | Bicyclo (2.2.1) heptan-2-one, 1,7,7-trimethyl-  | Yes   |                             |   |                                |                   | 0.05                           |
|               | Biochemical Oxygen Demand   |   | Yes                         |   | Yes                            |                   |                                |
| 00111-44-4    | Bis(2-Chloroethyl) ether  |   | Yes                         |   | Table 6                        | Tables 1,5        | 0.005                          |
| 00117-81-7    | Bis(2-ethylhexyl)phthalate / Di(2-ethylhexyl)phthalate  |   | Yes                         | 0.006   | Table 6                        | Tables 1,5        | 0.006                          |
| 68153-72-0    | Blown lard oil amine  | Yes   |                             |   |                                |                   | <b>§</b> §                     |
| 68876-82-4    | Blown rapeseed amine  | Yes   |                             |   |                                |                   | <b>§</b> §                     |
| 1319-33-1     | Borate Salt   | Yes   |                             |   |                                |                   |                                |
| 10043-35-3    | Boric acid  | Yes   |                             |   |                                |                   |                                |
| 01303-86-2    | Boric oxide / Boric Anhydride   | Yes   |                             |   |                                |                   |                                |
| 07440-42-8    | Boron   |   | Yes                         |   | Table 7                        | Tables 1,5        |                                |
| 24959-67-9    | Bromide   |   | Yes                         |   | Table 7                        | Tables 1,5        |                                |
| 00075-25-2    | Bromoform   |   | Yes                         |   | Table 6                        | Tables 1,5        | 0.005*                         |
| 00071-36-3    | Butan-1-ol  | Yes   |                             |   | Table 10                       | Tables 1,5        |                                |
| 68002-97-1    | C10 - C16 Ethoxylated Alcohol   | Yes   |                             |   |                                |                   | <b>§</b> §                     |
| 68131-39-5    | C12-15 Alcohol, Ethoxylated   | Yes   |                             |   |                                |                   | <b>§</b> §                     |
| 07440-43-9    | Cadmium   |   | Yes                         | 0.005   | Table 6                        | Tables 1,5        | 0.005                          |
| 07440-70-2    | Calcium   |   | Yes                         |   | Table 8                        |                   |                                |
| 1317-65-3     | Calcium Carbonate   | Yes   |                             |   | Table 10                       |                   |                                |
| 10043-52-4    | Calcium chloride  | Yes (¥)                                     |                             |   |                                |                   |                                |

| CAS<br>Number | Parameter Name                   | Used in<br>Additives <sup>285,</sup><br>286 | Found in<br>Flowback<br>287 | USEPA<br>MCL or TT<br>(mg/L) <sup>288,</sup><br>289 | SPDES<br>Tables <sup>290</sup> | TOGS111<br>Tables | NYS MCL, (mg/L) <sup>291</sup> |
|---------------|----------------------------------|---|-----------------------------|---|--------------------------------|-------------------|--------------------------------|
| 1305-62-0     | Calcium Hydroxide                | Yes   |                             |   |                                |                   |                                |
| 1305-79-9     | Calcium Peroxide                 | Yes   |                             |   |                                |                   |                                |
| 00124-38-9    | Carbon Dioxide                   | Yes   | Yes                         |   |                                |                   |                                |
| 00075-15-0    | Carbondisulfide                  |   | Yes                         |   | Table 8                        | Tables 1,5        |                                |
| 68130-15-4    | Carboxymethylhydroxypropyl guar  | Yes   |                             |   |                                |                   | <b>§§§</b>                     |
| 09012-54-8    | Cellulase / Hemicellulase Enzyme | Yes   |                             |   |                                |                   | §§§                            |
| 09004-34-6    | Cellulose                        | Yes   |                             |   |                                |                   | §§§                            |
|               | Cesium 137                       |   | Yes                         | Via beta radiation                                  |                                |                   | Via beta radiation             |
|               | Chemical Oxygen Demand           |   | Yes                         |   | Yes                            |                   |                                |
|               | Chloride                         |   | Yes                         |   | Table 7                        | Tables 1,5        | 250                            |
| 10049-04-4    | Chlorine Dioxide                 | Yes   |                             | MRDL=0.8  | Table 10                       |                   | MRDL=0.8                       |
| 00124-48-1    | Chlorodibromomethane             |   | Yes                         |   | Table 6                        | Tables 1,5        | 0.005*                         |
| 00067-66-3    | Chloroform                       |   | Yes                         |   | Table 6                        | Tables 1,5        | 0.005*                         |
| 78-73-9       | Choline Bicarbonate              | Yes   |                             |   |                                |                   | §                              |
| 67-48-1       | Choline Chloride                 | Yes   |                             |   |                                |                   | §                              |
| 91-64-5       | Chromen-2-one                    | Yes   |                             |   |                                |                   | 0.05                           |
| 07440-47-3    | Chromium                         |   | Yes                         | 0.1   | Table 6                        | Tables 1,5        | 0.1                            |
| 00077-92-9    | Citric Acid                      | Yes   |                             |   |                                |                   | 0.05                           |
| 94266-47-4    | Citrus Terpenes                  | Yes   |                             |   |                                |                   | <b>§</b> §                     |
| 07440-48-4    | Cobalt                           |   | Yes                         |   | Table 7                        | Table 1           |                                |
| 61789-40-0    | Cocamidopropyl Betaine           | Yes   |                             |   |                                |                   | 0.05                           |
| 68155-09-9    | Cocamidopropylamine Oxide        | Yes   |                             |   |                                |                   | 0.05                           |
| 68424-94-2    | Coco-betaine                     | Yes   |                             |   |                                |                   | 0.05                           |
|               | Coliform, Total                  |   | Yes                         | 0.05  | Table 7                        |                   |                                |
|               | Color                            |   | Yes                         |   | Table 7                        |                   |                                |
| 07440-50-8    | Copper                           |   | Yes                         | TT;<br>Action<br>Level=1.3                          | Table 6                        | Tables 1,5        | Action Level = 1.3             |
| 07758-98-7    | Copper (II) Sulfate              | Yes (¥)                                     |                             |   |                                |                   |                                |
| 14808-60-7    | Crystalline Silica (Quartz)      | Yes   |                             | Via solids<br>and TSS                               |                                |                   |                                |
| 07447-39-4    | Cupric chloride dihydrate        | Yes (¥)                                     |                             |   |                                |                   |                                |
| 00057-12-5    | Cyanide                          |   | Yes                         | 0.2   | Table 6                        | Tables 1,5        | 0.2                            |

| CAS<br>Number | Parameter Name  | Used in<br>Additives <sup>285</sup> ,<br>286 | Found in<br>Flowback<br>287 | USEPA<br>MCL or TT<br>(mg/L) <sup>288,</sup><br>289 | SPDES Tables | TOGS111<br>Tables | NYS MCL,<br>(mg/L) <sup>291</sup> |
|---------------|---|--|-----------------------------|---|--------------|-------------------|-----------------------------------|
| 00319-85-7    | Cyclohexane (beta BHC)  |  | Yes                         |   | Table 6      | Tables 1,5        | 0.005                             |
| 00058-89-9    | Cyclohexane (gamma BHC)   |  | Yes                         | 0.0002  | Table 6      | Tables 1,5        | 0.0002                            |
| 1490-04-6     | Cyclohexanol,5-methyl-2-(1-methylethyl)                               | Yes  |                             |   |              |                   | 0.05                              |
| 8007-02-1     | Cymbopogon citratus leaf oil  | Yes  |                             |   |              |                   | <b>§</b> §                        |
| 8000-29-1     | Cymbopogon winterianus jowitt oil                                     | Yes  |                             |   |              |                   | <b>§</b> §                        |
| 01120-24-7    | Decyldimethyl Amine   | Yes (¥)                                      |                             |   |              |                   | 0.05                              |
| 02605-79-0    | Decyl-dimethyl Amine Oxide  | Yes (¥)                                      |                             |   |              |                   | 0.05                              |
| 00055-70-3    | Dibenz(a,h)anthracene   |  | Yes                         |   |              | Table 3           | 0.05                              |
| 03252-43-5    | Dibromoacetonitrile   | Yes  |                             |   | Table 9      | Tables 1          | 0.05                              |
| 00075-27-4    | Dichlorobromomethane  |  | Yes                         |   | Table 6      | Tables 1,5        | 0.005*                            |
| 25340-17-4    | Diethylbenzene  | Yes  |                             |   |              |                   | 0.05                              |
| 00111-46-6    | Diethylene Glycol   | Yes  |                             |   | Table 10     |                   | 0.05                              |
| 22042-96-2    | Diethylenetriamine penta (methylenephonic acid) sodium salt           | Yes  |                             |   |              |                   | 0.05                              |
| 28757-00-8    | Diisopropyl naphthalenesulfonic acid                                  | Yes  |                             |   |              |                   | 0.05                              |
| 68607-28-3    | Dimethylcocoamine, bis(chloroethyl) ether, diquaternary ammonium salt | Yes  |                             |   |              |                   | 0.05                              |
| 07398-69-8    | Dimethyldiallylammonium chloride                                      | Yes  |                             |   |              |                   | 0.05                              |
| 00084-74-2    | Di-n-butyl phthalate  |  | Yes                         |   | Table 6      | Tables 1,5        | 0.05                              |
| 00122-39-4    | Diphenylamine   |  | Yes                         |   | Table 7      | Tables 1,5        | 0.005                             |
| 25265-71-8    | Dipropylene glycol  | Yes  |                             |   |              |                   | 0.05                              |
| 34590-94-8    | Dipropylene glycol methyl ether                                       | Yes  |                             |   |              |                   | 0.05                              |
| 00139-33-3    | Disodium Ethylene Diamine Tetra Acetate                               | Yes  |                             |   |              |                   | 0.05                              |
| 64741-77-1    | Distillates, petroleum, light hydrocracked                            | Yes  |                             |   |              |                   | <b>§</b> §                        |
| 05989-27-5    | D-Limonene  | Yes  |                             |   |              |                   | 0.05                              |
| 00123-01-3    | Dodecylbenzene  | Yes  |                             |   |              |                   | 0.05                              |
| 27176-87-0    | Dodecylbenzene sulfonic acid  | Yes  |                             |   |              |                   | 0.05                              |
| 42504-46-1    | Dodecylbenzenesulfonate isopropanolamine                              | Yes  |                             |   |              |                   | 0.05                              |
| 00050-70-4    | D-Sorbitol / Sorbitol   | Yes  |                             |   |              |                   | 0.05                              |
| 37288-54-3    | Endo-1,4-beta-mannanase, or Hemicellulase                             | Yes  |                             |   |              |                   | 0.05                              |
| 00959-98-8    | Endosulfan I  |  | Yes                         |   | Table 6      | Table 3           | 0.05                              |
| 33213-65-9    | Endosulfan II   |  | Yes                         |   | Table 6      | Table 3           | 0.05                              |
| 07421-93-4    | Endrin aldehyde   |  | Yes                         |   | Table 6      | Tables 1,5        | 0.005                             |
| 149879-98-1   | Erucic Amidopropyl Dimethyl Betaine                                   | Yes  |                             |   |              |                   | 0.05                              |

| CAS<br>Number | Parameter Name  | Used in<br>Additives <sup>285</sup> ,<br>286 | Found in<br>Flowback<br>287 | USEPA<br>MCL or TT<br>(mg/L) <sup>288,</sup><br>289 | SPDES<br>Tables <sup>290</sup> | TOGS111<br>Tables | NYS MCL,<br>(mg/L) <sup>291</sup> |
|---------------|---|--|-----------------------------|---|--------------------------------|-------------------|-----------------------------------|
| 00089-65-6    | Erythorbic acid, anhydrous  | Yes  |                             |   |                                |                   | 0.05                              |
| 54076-97-0    | Ethanaminium, N,N,N-trimethyl-2-[(1-oxo-2-propenyl)oxy]-, chloride, homopolymer | Yes  |                             |   |                                |                   | 0.05                              |
| 00107-21-1    | Ethane-1,2-diol / Ethylene Glycol   | Yes  | Yes                         |   | Table 7                        | Tables 1,5        | 0.05                              |
| 111-42-2      | Ethanol, 2,2-iminobis-  | Yes  |                             |   |                                |                   | 0.05                              |
| 26027-38-3    | Ethoxylated 4-nonylphenol   | Yes  |                             |   |                                |                   | 0.05                              |
| 09002-93-1    | Ethoxylated 4-tert-octylphenol  | Yes  |                             |   |                                |                   | 0.05                              |
| 68439-50-9    | Ethoxylated alcohol   | Yes  |                             |   |                                |                   | §§                                |
| 126950-60-5   | Ethoxylated alcohol   | Yes  |                             |   |                                |                   | 88                                |
| 68951-67-7    | Ethoxylated alcohol (C14-15)  | Yes  |                             |   |                                |                   | §§                                |
| 68439-46-3    | Ethoxylated alcohol (C9-11)   | Yes  |                             |   |                                |                   | 88                                |
| 66455-15-0    | Ethoxylated Alcohols  | Yes  |                             |   |                                |                   | 88                                |
| 67254-71-1    | Ethoxylated Alcohols (C10-12)   | Yes  |                             |   |                                |                   | §§                                |
| 84133-50-6    | Ethoxylated Alcohols (C12-14 Secondary)   | Yes  |                             |   |                                |                   | §§                                |
| 68439-51-0    | Ethoxylated Alcohols (C12-14)   | Yes  |                             |   |                                |                   | §§                                |
| 78330-21-9    | Ethoxylated branch alcohol  | Yes  |                             |   |                                |                   | §§                                |
| 34398-01-1    | Ethoxylated C11 alcohol   | Yes  |                             |   |                                |                   | §§                                |
| 78330-21-8    | Ethoxylated C11-14-iso, C13-rich alcohols                                       | Yes  |                             |   |                                |                   | §§                                |
| 61791-12-6    | Ethoxylated Castor Oil  | Yes  |                             |   |                                |                   | §§                                |
| 61791-29-5    | Ethoxylated fatty acid, coco  | Yes  |                             |   |                                |                   | §§                                |
| 61791-08-0    | Ethoxylated fatty acid, coco, reaction product with ethanolamine                | Yes  |                             |   |                                |                   | §§                                |
| 68439-45-2    | Ethoxylated hexanol   | Yes  |                             |   |                                |                   | §§                                |
| 09036-19-5    | Ethoxylated octylphenol   | Yes  |                             |   |                                |                   | 0.05                              |
| 09005-67-8    | Ethoxylated Sorbitan Monostearate   | Yes  |                             |   |                                |                   | 0.05                              |
| 09005-70-3    | Ethoxylated Sorbitan Trioleate  | Yes  |                             |   |                                |                   | 0.05                              |
| 118-61-6      | Ethyl 2-hydroxybenzoate   | Yes  |                             |   |                                |                   | 0.05                              |
| 00064-17-5    | Ethyl alcohol / ethanol   | Yes  |                             |   |                                |                   | 0.05                              |
| 00100-41-4    | Ethyl Benzene   | Yes  | Yes                         | 0.7   | Table 6                        | Tables 1,5        | 0.005                             |
| 93-89-0       | Ethyl benzoate  | Yes  |                             |   |                                |                   | 0.05                              |
| 00097-64-3    | Ethyl Lactate   | Yes  |                             |   |                                |                   | 0.05                              |

| CAS<br>Number | Parameter Name  | Used in<br>Additives <sup>285,</sup><br>286 | Found in<br>Flowback<br>287 | USEPA<br>MCL or TT<br>(mg/L) <sup>288,</sup><br>289 | SPDES<br>Tables <sup>290</sup> | TOGS111<br>Tables | NYS MCL,<br>(mg/L) <sup>291</sup> |
|---------------|---|---|-----------------------------|---|--------------------------------|-------------------|-----------------------------------|
| 09003-11-6    | Ethylene Glycol-Propylene Glycol Copolymer (Oxirane, methyl-, polymer with oxirane)           | Yes   |                             |   |                                |                   | 0.05                              |
| 00075-21-8    | Ethylene oxide  | Yes   |                             |   | Table 9                        | Tables 1,5        | 0.05                              |
| 05877-42-9    | Ethyloctynol  | Yes   |                             |   |                                |                   | 0.05                              |
| 8000-48-4     | Eucalyptus globulus leaf oil  | Yes   |                             |   |                                |                   | §§                                |
| 61790-12-3    | Fatty Acids   | Yes   |                             |   |                                |                   | <b>§</b> §                        |
| 68604-35-3    | Fatty acids, C 8-18 and C18-unsaturated compounds with diethanolamine                         | Yes   |                             |   |                                |                   | <b>§</b> §                        |
| 68188-40-9    | Fatty acids, tall oil reaction products w/ acetophenone, formaldehyde & thiourea              | Yes   |                             |   |                                |                   | <b>§</b> §                        |
| 09043-30-5    | Fatty alcohol polyglycol ether surfactant   | Yes   |                             |   |                                |                   | §§                                |
| 07705-08-0    | Ferric chloride   | Yes   |                             |   | Table 10                       |                   |                                   |
| 07782-63-0    | Ferrous sulfate, heptahydrate   | Yes   |                             |   |                                |                   |                                   |
| 00206-44-0    | Fluoranthene  |   | Yes                         |   | Table 6                        | Tables 1,5        | 0.05                              |
| 00086-73-7    | Fluorene  |   | Yes                         |   | Table 6                        | Tables 1,5        | 0.05                              |
| 16984-48-8    | Fluoride  |   | Yes                         | 4   | Table 7                        | Tables 1,5        | 2.2                               |
| 00050-00-0    | Formaldehyde  | Yes   |                             |   | Table 8                        | Tables 1,5        |                                   |
| 29316-47-0    | Formaldehyde polymer with 4,1,1-dimethylethyl phenolmethyl oxirane                            | Yes   |                             |   |                                |                   | 0.05                              |
| 153795-76-7   | Formaldehyde, polymers with branched 4-<br>nonylphenol, ethylene oxide and propylene<br>oxide | Yes   |                             |   |                                |                   | 0.05                              |
| 00075-12-7    | Formamide   | Yes   |                             |   |                                |                   | 0.05                              |
| 00064-18-6    | Formic acid   | Yes   |                             |   | Table 10                       |                   | 0.05                              |
| 00110-17-8    | Fumaric acid  | Yes   |                             |   | Table 10                       |                   | 0.05                              |
| 65997-17-3    | Glassy calcium magnesium phosphate  | Yes   |                             |   |                                |                   |                                   |
| 00111-30-8    | Glutaraldehyde  | Yes   |                             |   |                                |                   | 0.05                              |
| 00056-81-5    | Glycerol / glycerine  | Yes   |                             |   |                                |                   | 0.05                              |
| 09000-30-0    | Guar Gum  | Yes   |                             |   |                                |                   | 0.05                              |
| 64742-94-5    | Heavy aromatic petroleum naphtha  | Yes   |                             |   |                                |                   | 0.05                              |
| 09025-56-3    | Hemicellulase   | Yes   |                             |   |                                |                   | 0.05                              |
| 00076-44-8    | Heptachlor  |   | Yes                         | 0.0002  |                                | Tables 1,5        | 0.0004                            |
| 01024-57-3    | Heptachlor epoxide  |   | Yes                         | 0.0002  |                                | Tables 1,5        | 0.0002                            |

| CAS<br>Number | Parameter Name   | Used in<br>Additives <sup>285,</sup><br>286 | Found in<br>Flowback<br>287 | USEPA<br>MCL or TT<br>(mg/L) <sup>288,</sup><br>289 | SPDES Tables 290 | TOGS111<br>Tables | NYS MCL, (mg/L) <sup>291</sup> |
|---------------|--|---|-----------------------------|---|------------------|-------------------|--------------------------------|
|               | Heterotrophic plate count  |   | Yes                         | TT <sup>292</sup>                                   |                  |                   |                                |
| 07647-01-0    | Hydrochloric Acid / Hydrogen Chloride / muriatic acid              | Yes   |                             |   |                  |                   |                                |
| 07722-84-1    | Hydrogen Peroxide  | Yes   |                             |   | Table 10         |                   |                                |
| 64742-52-5    | Hydrotreated heavy napthenic distillate                            | Yes   |                             |   |                  |                   | <b>§</b> §                     |
| 00079-14-1    | Hydroxy acetic acid  | Yes   |                             |   |                  |                   | 0.05                           |
| 35249-89-9    | Hydroxyacetic acid ammonium salt                                   | Yes   |                             |   |                  |                   | 0.05                           |
| 09004-62-0    | Hydroxyethyl cellulose   | Yes   |                             |   |                  |                   | 0.05                           |
| 05470-11-1    | Hydroxylamine hydrochloride  | Yes   |                             |   |                  |                   | 0.05                           |
| 39421-75-5    | Hydroxypropyl guar   | Yes   |                             |   |                  |                   | 0.05                           |
| 00193-39-5    | Indeno(1,2,3-cd)pyrene   |   | Yes                         |   | Table 6          | Tables 1,5        | 0.05                           |
| 07439-89-6    | Iron   |   | Yes                         |   | Table 7          | Tables 1,5        | 0.3                            |
| 35674-56-7    | Isomeric Aromatic Ammonium Salt                                    | Yes   |                             |   |                  |                   | 0.05                           |
| 64742-88-7    | Isoparaffinic Petroleum Hydrocarbons,<br>Synthetic                 | Yes   |                             |   |                  |                   | §§                             |
| 00064-63-0    | Isopropanol  | Yes   |                             |   | Table 10         |                   | 0.05                           |
| 00098-82-8    | Isopropylbenzene (cumene)  | Yes   | Yes                         |   | Table 9          | Tables 1,5        | 0.005                          |
| 68909-80-8    | Isoquinoline, reaction products with benzyl chloride and quinoline | Yes   |                             |   |                  |                   | 0.05                           |
| 08008-20-6    | Kerosene   | Yes   |                             |   |                  |                   | <b>§</b> §                     |
| 64742-81-0    | Kerosine, hydrodesulfurized  | Yes   |                             |   |                  |                   | <b>§</b> §                     |
| 00063-42-3    | Lactose  | Yes   |                             |   |                  |                   |                                |
| 8022-15-9     | Lavandula hybrida abrial herb oil                                  | Yes   |                             |   |                  |                   | §§                             |
| 07439-92-1    | Lead   |   | Yes                         | TT;<br>Action Level<br>0.015                        | Table 6          | Tables 1,5        | Action level = 0.015           |
| 64742-95-6    | Light aromatic solvent naphtha                                     | Yes   |                             |   |                  |                   | <b>§</b> §                     |
| 01120-21-4    | Light Paraffin Oil   | Yes   |                             |   |                  |                   | <b>§</b> §                     |
|               | Lithium  |   | Yes                         |   | Table 10         |                   |                                |
| 07439-95-4    | Magnesium  |   | Yes                         |   | Table 7          | Tables 1,5        |                                |
| 546-93-0      | Magnesium Carbonate  | Yes   |                             |   |                  |                   |                                |
| 1309-48-4     | Magnesium Oxide  | Yes   |                             |   |                  |                   |                                |

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<sup>&</sup>lt;sup>292</sup> Treatment Technology specified.

| CAS<br>Number | Parameter Name   | Used in<br>Additives <sup>285,</sup><br>286 | Found in<br>Flowback<br>287 | USEPA<br>MCL or TT<br>(mg/L) <sup>288,</sup><br>289 | SPDES<br>Tables <sup>290</sup> | TOGS111<br>Tables | NYS MCL,<br>(mg/L) <sup>291</sup> |
|---------------|--|---|-----------------------------|---|--------------------------------|-------------------|-----------------------------------|
| 1335-26-8     | Magnesium Peroxide   | Yes   |                             |   |                                |                   |                                   |
| 14807-96-6    | Magnesium Silicate Hydrate (Talc)                                      | Yes   |                             |   |                                |                   |                                   |
| 07439-96-5    | Manganese  |   | Yes                         |   | Table 7                        | Tables 1,5        | 0.3                               |
| 07439-97-6    | Mercury  |   | Yes                         | 0.002   | Table 6                        | Tables 1,5        | 0.002                             |
| 01184-78-7    | Methanamine, N,N-dimethyl-, N-oxide                                    | Yes   |                             |   |                                |                   | 0.05                              |
| 00067-56-1    | Methanol   | Yes   | Yes                         |   | Table 10                       |                   | 0.05                              |
| 119-36-8      | Methyl 2-hydroxybenzoate   | Yes   |                             |   |                                |                   | 0.05                              |
| 00074-83-9    | Methyl Bromide   |   | Yes                         |   | Table 6                        | Tables 1,5        | 0.005                             |
| 00074-87-3    | Methyl Chloride / chloromethane  |   | Yes                         | 0.005   | Table 6                        | Tables 1,5        | 0.005                             |
| 00078-93-3    | Methyl ethyl ketone / 2-Butanone                                       |   | Yes                         |   | Table 7                        | Tables 1,5        | 0.05                              |
| 68891-11-2    | Methyloxirane polymer with oxirane, mono (nonylphenol) ether, branched | Yes   |                             |   |                                |                   | 0.05                              |
| 08052-41-3    | Mineral spirits / Stoddard Solvent                                     | Yes   |                             |   |                                |                   | §§                                |
| 64742-46-7    | Mixture of severely hydrotreated and hydrocracked base oil             | Yes   |                             |   |                                |                   | §§                                |
| 07439-98-7    | Molybdenum   |   | Yes                         |   | Table 7                        |                   |                                   |
| 00141-43-5    | Monoethanolamine   | Yes   |                             |   |                                |                   | 0.05                              |
| 44992-01-0    | N,N,N-trimethyl-2[1-oxo-2-propenyl]oxy<br>Ethanaminium chloride        | Yes   |                             |   |                                |                   | 0.05                              |
| 64742-48-9    | Naphtha (petroleum), hydrotreated heavy                                | Yes   |                             |   |                                |                   | <b>§</b> §                        |
| 00091-20-3    | Naphthalene  | Yes   | Yes                         |   | Table 6                        | Tables 1,5        | 0.05                              |
| 38640-62-9    | Naphthalene bis(1-methylethyl)   | Yes   |                             |   |                                |                   | 0.05                              |
| 00093-18-5    | Naphthalene, 2-ethoxy-   | Yes   |                             |   |                                |                   | 0.05                              |
| 68909-18-2    | N-benzyl-alkyl-pyridinium chloride                                     | Yes   |                             |   |                                |                   | 0.05                              |
| 68139-30-0    | N-Cocoamidopropyl-N,N-dimethyl-N-2-<br>hydroxypropylsulfobetaine       | Yes   |                             |   |                                |                   | 0.05                              |
| 07440-02-0    | Nickel   |   | Yes                         |   | Table 6                        | Tables 1,5        |                                   |
|               | Nitrate, as N  |   | Yes                         | 10  | Table 7                        | Tables 1,5        | 10                                |
| 07727-37-9    | Nitrogen, Liquid form  | Yes   |                             |   |                                |                   |                                   |
|               | Nitrogen, Total as N   |   | Yes                         |   |                                | Table 5           |                                   |
| 00086-30-6    | N-Nitrosodiphenylamine   |   | Yes                         |   | Table 6                        | Tables 1,5        | 0.05                              |
| 26027-38-3    | Nonylphenol Ethoxylate   | Yes   |                             |   |                                |                   | 0.05                              |

| CAS<br>Number | Parameter Name   | Used in<br>Additives <sup>285</sup> ,<br>286 | Found in<br>Flowback<br>287 | USEPA<br>MCL or TT<br>(mg/L) <sup>288,</sup><br>289 | SPDES<br>Tables <sup>290</sup> | TOGS111<br>Tables | NYS MCL,<br>(mg/L) |
|---------------|--|--|-----------------------------|---|--------------------------------|-------------------|--------------------|
| 68412-54-4    | Nonylphenol Polyethoxylate   | Yes  |                             |   |                                |                   | 0.05               |
|               | Oil and Grease   |  | Yes                         |   |                                | Table 5           |                    |
| 8000-27-9     | Oils, cedarwood  | Yes  |                             |   |                                |                   | <b>§</b> §         |
| 121888-66-2   | Organophilic Clays   | Yes  |                             |   |                                |                   |                    |
|               | Oxyalkylated alkylphenol   | Yes  |                             |   |                                |                   | 0.05               |
| 628-63-7      | Pentyl acetate   | Yes  |                             |   |                                |                   | 0.05               |
| 540-18-1      | Pentyl butanoate   | Yes  |                             |   |                                |                   | 0.05               |
| 8009-03-8     | Petrolatum   | Yes  |                             |   |                                |                   | §§                 |
| 64742-65-0    | Petroleum Base Oil   | Yes  |                             |   |                                |                   | §§                 |
|               | Petroleum distillate blend   | Yes  |                             |   |                                |                   |                    |
| 64742-52-5    | Petroleum Distillates  | Yes  |                             |   |                                |                   | §§                 |
|               | Petroleum hydrocarbons   |  | Yes                         |   |                                |                   |                    |
| 64741-68-0    | Petroleum naphtha  | Yes  |                             |   |                                |                   | 0.05               |
|               | pH   |  | Yes                         |   |                                | Table 5           |                    |
| 00085-01-8    | Phenanthrene   |  | Yes                         |   | Table 6                        | Tables 1,5        | 0.05               |
| 00108-95-2    | Phenol   |  | Yes                         |   | Table 6                        | Tables 1,5        | 0.05               |
|               | Phenols  |  | Yes                         |   | Table 6                        | Tables 1,5        |                    |
| 101-84-8      | Phenoxybenzene   | Yes  |                             |   |                                |                   | 0.05               |
| 70714-66-8    | Phosphonic acid, [[(phosphonomethyl)imino]bis[2,1- ethanediylnitrilobis(methylene)]]tetrakis-, ammonium salt | Yes  |                             |   |                                |                   | §                  |
| 57723-14-0    | Phosphorus   |  | Yes                         |   | Table 7                        | Table 1           |                    |
| 08000-41-7    | Pine Oil   | Yes  |                             |   |                                |                   | <b>§</b> §         |
| 8002-09-3     | Pine oils  | Yes  |                             |   |                                |                   | §§                 |
| 60828-78-6    | Poly(oxy-1,2-ethanediyl), a-[3,5-dimethyl-1-(2-methylpropyl)hexyl]-w-hydroxy-                                | Yes  |                             |   |                                |                   | §§§                |
| 25322-68-3    | Poly(oxy-1,2-ethanediyl), a-hydro-w-hydroxy /<br>Polyethylene Glycol   | Yes  |                             |   |                                |                   | §§§                |
| 24938-91-8    | Poly(oxy-1,2-ethanediyl), α-tridecyl- ω-<br>hydroxy  | Yes  |                             |   |                                |                   | 888                |
| 31726-34-8    | Poly(oxy-1,2-ethanediyl),alpha-hexyl-omega-<br>hydroxy   | Yes  |                             |   |                                |                   | §§§                |

| CAS<br>Number | Parameter Name   | Used in<br>Additives <sup>285</sup> ,<br>286 | Found in<br>Flowback<br>287 | USEPA<br>MCL or TT<br>(mg/L) <sup>288,</sup><br>289 | SPDES<br>Tables <sup>290</sup> | TOGS111<br>Tables      | NYS MCL,<br>(mg/L) <sup>291</sup> |
|---------------|--|--|-----------------------------|---|--------------------------------|------------------------|-----------------------------------|
| 9004-32-4     | Polyanionic Cellulose                                  | Yes  |                             |   |                                |                        | 888                               |
| 51838-31-4    | Polyepichlorohydrin, trimethylamine quaternized        | Yes  |                             |   |                                |                        | §§§                               |
| 56449-46-8    | polyethlene glycol oleate ester                        | Yes  |                             |   |                                |                        | <b>§§§</b>                        |
|               | Polyethoxylated alkanol                                | Yes  |                             |   |                                |                        |                                   |
| 9046-01-9     | Polyethoxylated tridecyl ether phosphate               | Yes  |                             |   |                                |                        | <b>§</b> §                        |
| 63428-86-4    | Polyethylene glycol hexyl ether sulfate, ammonium salt | Yes  |                             |   |                                |                        | §                                 |
| 62649-23-4    | Polymer with 2-propenoic acid and sodium 2-propenoate  | Yes  |                             |   |                                |                        | <b>§§§</b>                        |
|               | Polymeric Hydrocarbons                                 | Yes  |                             |   |                                |                        | <b>§</b> §                        |
| 09005-65-6    | Polyoxyethylene Sorbitan Monooleate                    | Yes  |                             |   |                                |                        | 0.05                              |
| 61791-26-2    | Polyoxylated fatty amine salt                          | Yes  |                             |   |                                |                        | 0.05                              |
| 65997-18-4    | Polyphosphate  | Yes  |                             |   |                                |                        |                                   |
| 07440-09-7    | Potassium  |  | Yes                         |   | Table 8                        |                        |                                   |
| 00127-08-2    | Potassium acetate                                      | Yes  |                             |   |                                |                        | §                                 |
| 1332-77-0     | Potassium borate                                       | Yes  |                             |   |                                |                        |                                   |
| 12712-38-8    | Potassium borate                                       | Yes  |                             |   |                                |                        |                                   |
| 20786-60-1    | Potassium borate                                       | Yes  |                             |   |                                |                        |                                   |
| 00584-08-7    | Potassium carbonate                                    | Yes  |                             |   |                                |                        |                                   |
| 07447-40-7    | Potassium chloride                                     | Yes  |                             |   |                                |                        | §                                 |
| 00590-29-4    | Potassium formate                                      | Yes  |                             |   |                                |                        |                                   |
| 01310-58-3    | Potassium Hydroxide                                    | Yes  |                             |   | Table 10                       |                        |                                   |
| 13709-94-9    | Potassium metaborate                                   | Yes  |                             |   |                                |                        |                                   |
| 24634-61-5    | Potassium Sorbate                                      | Yes  |                             |   |                                |                        | §                                 |
| 112926-00-8   | Precipitated silica / silica gel                       | Yes  |                             |   |                                |                        |                                   |
| 00057-55-6    | Propane-1,2-diol, or Propylene glycol                  | Yes  | Yes                         |   | Table 10                       | Table 3 <sup>293</sup> | 1.0                               |
| 00057-55-6    | Propylene glycol                                       |  |                             |   |                                |                        | 1.0                               |
| 00107-98-2    | Propylene glycol monomethyl ether                      | Yes  |                             |   | Table 10                       |                        | 0.05                              |
| 00110-86-1    | Pyridine   |  | Yes                         |   | Table 7                        | Tables 1,5             | 0.05                              |
| 68953-58-2    | Quaternary Ammonium Compounds                          | Yes  |                             |   | Table 9                        | Tables 1               | §§                                |

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<sup>&</sup>lt;sup>293</sup> TOGS lists this parameter as CAS 58-55-6.

| CAS<br>Number | Parameter Name                                     | Used in<br>Additives <sup>285,</sup><br>286 | Found in<br>Flowback<br>287 | USEPA<br>MCL or TT<br>(mg/L) <sup>288,</sup><br>289 | SPDES<br>Tables <sup>290</sup> | TOGS111<br>Tables | NYS MCL,<br>(mg/L) <sup>291</sup> |
|---------------|--|---|-----------------------------|---|--------------------------------|-------------------|-----------------------------------|
| 62763-89-7    | Quinoline,2-methyl-, hydrochloride                 | Yes   |                             |   |                                |                   | 0.05                              |
| 15619-48-4    | Quinolinium, 1-(phenylmethl),chloride              | Yes   |                             |   |                                |                   | 0.05                              |
| 8000-25-7     | Rosmarinus officinalis l. leaf oil                 | Yes   |                             |   |                                |                   | §§                                |
| 00094-59-7    | Safrole  |   | Yes                         |   | Table 8                        | Table 3           | 0.05                              |
|               | Salt of amine-carbonyl condensate                  | Yes   |                             |   |                                |                   |                                   |
|               | Salt of fatty acid/polyamine reaction product      | Yes   |                             |   |                                |                   |                                   |
|               | Scale Inhibitor (mg/L)                             |   | Yes                         |   |                                |                   |                                   |
| 07782-49-2    | Selenium   |   | Yes                         | 0.05  | Table 6                        | Tables 1,5        | 0.05                              |
| 07631-86-9    | Silica, Dissolved                                  | Yes   |                             |   | Table 8                        |                   |                                   |
| 07440-22-4    | Silver   |   | Yes                         |   | Table 6                        | Tables 1,5        | 0.1                               |
| 07440-23-5    | Sodium   |   | Yes                         |   | Table 7                        | Tables 1,5        |                                   |
| 05324-84-5    | Sodium 1-octanesulfonate                           | Yes   |                             |   |                                |                   | §                                 |
| 00127-09-3    | Sodium acetate                                     | Yes   |                             |   |                                |                   | §                                 |
| 95371-16-7    | Sodium Alpha-olefin Sulfonate                      | Yes   |                             |   |                                |                   | §                                 |
| 00532-32-1    | Sodium Benzoate                                    | Yes   |                             |   |                                |                   | §                                 |
| 00144-55-8    | Sodium bicarbonate                                 | Yes   |                             |   |                                |                   |                                   |
| 07631-90-5    | Sodium bisulfate                                   | Yes   |                             |   |                                |                   |                                   |
| 07647-15-6    | Sodium Bromide                                     | Yes   |                             |   |                                |                   |                                   |
| 00497-19-8    | Sodium carbonate                                   | Yes   |                             |   |                                |                   |                                   |
| 07647-14-5    | Sodium Chloride                                    | Yes   |                             |   |                                |                   |                                   |
| 07758-19-2    | Sodium chlorite                                    | Yes   |                             |   |                                |                   | 1.0 (chlorite)                    |
| 03926-62-3    | Sodium Chloroacetate                               | Yes   |                             |   |                                |                   | §                                 |
| 00068-04-2    | Sodium citrate                                     | Yes   |                             |   |                                |                   | §                                 |
| 06381-77-7    | Sodium erythorbate / isoascorbic acid, sodium salt | Yes   |                             |   |                                |                   | §                                 |
| 02836-32-0    | Sodium Glycolate                                   | Yes   |                             |   |                                |                   | §                                 |
| 1301-73-2     | Sodium hydroxide                                   | Yes   |                             |   |                                |                   |                                   |
| 01310-73-2    | Sodium Hydroxide                                   | Yes   |                             |   | Table 10                       |                   |                                   |
| 07681-52-9    | Sodium hypochlorite                                | Yes   |                             |   | Table 10                       |                   |                                   |
| 07775-19-1    | Sodium Metaborate .8H2O                            | Yes   |                             |   |                                |                   |                                   |
| 10486-00-7    | Sodium perborate tetrahydrate                      | Yes   |                             |   |                                |                   |                                   |
| 07775-27-1    | Sodium persulphate                                 | Yes   |                             |   |                                |                   |                                   |
| 68608-26-4    | Sodium petroleum sulfonate                         | Yes   |                             |   |                                |                   |                                   |

| CAS<br>Number | Parameter Name  | Used in<br>Additives <sup>285</sup> ,<br>286 | Found in<br>Flowback<br>287 | USEPA<br>MCL or TT<br>(mg/L) <sup>288,</sup><br>289 | SPDES<br>Tables <sup>290</sup> | TOGS111<br>Tables | NYS MCL, (mg/L) <sup>291</sup> |
|---------------|---|--|-----------------------------|---|--------------------------------|-------------------|--------------------------------|
| 09003-04-7    | Sodium polyacrylate   | Yes  |                             |   |                                |                   | §                              |
| 07757-82-6    | Sodium sulfate  | Yes  |                             |   | Table 10                       |                   |                                |
| 01303-96-4    | Sodium tetraborate decahydrate                                      | Yes  |                             |   |                                |                   |                                |
| 07772-98-7    | Sodium Thiosulfate  | Yes  |                             |   |                                |                   |                                |
| 01338-43-8    | Sorbitan Monooleate   | Yes  |                             |   |                                |                   | 0.05                           |
|               | Specific Conductivity   |  | Yes                         |   |                                |                   |                                |
| 07440-24-6    | Strontium   |  | Yes                         |   | Table 9                        | Table 1           |                                |
| 00057-50-1    | Sucrose   | Yes  |                             |   |                                |                   |                                |
|               | Sugar   | Yes  |                             |   |                                |                   |                                |
| 05329-14-6    | Sulfamic acid   | Yes  |                             |   |                                |                   |                                |
| 14808-79-8    | Sulfate   |  | Yes                         |   | Table 7                        | Tables 1,5        | 250                            |
|               | Sulfide   |  | Yes                         |   | Table 7                        | Tables 1,5        |                                |
| 14265-45-3    | Sulfite   |  | Yes                         |   | Table 7                        | Table 1           |                                |
|               | Surfactant blend  | Yes  |                             |   |                                |                   |                                |
| 68442-77-3    | Surfactant: Modified Amine  | Yes  |                             |   |                                |                   | §§                             |
|               | Surfactants MBAS  |  | Yes                         |   |                                |                   |                                |
| 112945-52-5   | Syntthetic Amorphous / Pyrogenic Silica /<br>Amorphous Silica       | Yes  |                             |   |                                |                   |                                |
| 68155-20-4    | Tall Oil Fatty Acid Diethanolamine                                  | Yes  |                             |   |                                |                   | <b>§</b> §                     |
| 08052-48-0    | Tallow fatty acids sodium salt                                      | Yes  |                             |   |                                |                   | §,§§                           |
| 72480-70-7    | Tar bases, quinoline derivs., benzyl chloride-<br>quaternized       | Yes  |                             |   |                                |                   | §§                             |
| 68647-72-3    | Terpene and terpenoids  | Yes  |                             |   |                                |                   | §§                             |
| 68956-56-9    | Terpene hydrocarbon byproducts                                      | Yes  |                             |   |                                |                   | §§                             |
| 00127-18-4    | Tetrachloroethylene   |  | Yes                         | 0.005   | Table 6                        | Tables 1,5        | 0.005                          |
| 00533-74-4    | Tetrahydro-3,5-dimethyl-2H-1,3,5-thiadiazine-<br>2-thione / Dazomet | Yes  |                             |   |                                | ,                 | 0.05                           |
| 55566-30-8    | Tetrakis(hydroxymethyl)phosphonium sulfate (THPS)                   | Yes  |                             |   |                                |                   | 0.05                           |
| 00075-57-0    | Tetramethyl ammonium chloride                                       | Yes  |                             |   |                                |                   | §                              |
| 00064-02-8    | Tetrasodium Ethylenediaminetetraacetate                             | Yes  |                             |   |                                |                   | §                              |
| 07440-28-0    | Thallium  |  | Yes                         | 0.002   | Table 6                        | Tables 1,5        | 0.002                          |
| 00068-11-1    | Thioglycolic acid   | Yes  |                             |   |                                |                   | 0.05                           |
| 00062-56-6    | Thiourea  | Yes  |                             |   | Table 10                       |                   | 0.05                           |

| CAS<br>Number | Parameter Name   | Used in<br>Additives <sup>285</sup> ,<br>286 | Found in<br>Flowback<br>287 | USEPA<br>MCL or TT<br>(mg/L) <sup>288,</sup><br>289 | SPDES<br>Tables <sup>290</sup> | TOGS111<br>Tables | NYS MCL,<br>(mg/L) <sup>291</sup> |
|---------------|--|--|-----------------------------|---|--------------------------------|-------------------|-----------------------------------|
| 68527-49-1    | Thiourea, polymer with formaldehyde and 1-phenylethanone | Yes  |                             |   |                                |                   | §§§                               |
| 68917-35-1    | Thuja plicata donn ex. D. don leaf oil                   | Yes  |                             |   |                                |                   | §§                                |
| 07440-32-6    | Titanium   |  | Yes                         |   | Table 7                        |                   |                                   |
| 00108-88-3    | Toluene  | Yes  | Yes                         | 1   | Table 6                        | Tables 1,5        | 0.005                             |
|               | Total Dissolved Solids                                   |  | Yes                         |   |                                | Table 5           |                                   |
|               | Total Kjeldahl Nitrogen                                  |  | Yes                         |   | Yes                            |                   |                                   |
|               | Total Organic Carbon                                     |  | Yes                         |   | Yes                            |                   |                                   |
|               | Total Suspended Solids                                   |  | Yes                         |   | Yes                            |                   |                                   |
| 81741-28-8    | Tributyl tetradecyl phosphonium chloride                 | Yes  |                             |   |                                |                   | §                                 |
|               | Triethanolamine  | Yes  |                             |   |                                |                   | 0.05                              |
| 68299-02-5    | Triethanolamine hydroxyacetate                           | Yes  |                             |   |                                |                   | 0.05                              |
| 00112-27-6    | Triethylene Glycol                                       | Yes  |                             |   |                                |                   | 0.05                              |
| 52624-57-4    | Trimethylolpropane, Ethoxylated,<br>Propoxylated         | Yes  |                             |   |                                |                   | §§                                |
| 00150-38-9    | Trisodium Ethylenediaminetetraacetate                    | Yes  |                             |   |                                |                   | §                                 |
| 05064-31-3    | Trisodium Nitrilotriacetate                              | Yes  |                             |   |                                |                   | §0.05                             |
| 07601-54-9    | Trisodium ortho phosphate                                | Yes  |                             |   |                                |                   |                                   |
| 00057-13-6    | Urea   | Yes  |                             |   |                                |                   | 0.05                              |
| 07440-62-2    | Vanadium   |  | Yes                         |   | Table 7                        | Table 1           |                                   |
| 25038-72-6    | Vinylidene Chloride/Methylacrylate<br>Copolymer          | Yes  |                             |   |                                |                   | <b>§§§</b>                        |
|               | Volatile Acids   |  | Yes                         |   | 294                            |                   |                                   |
| 7732-18-5     | Water  | Yes  |                             |   |                                |                   |                                   |
| 8042-47-5     | White Mineral Oil  | Yes  |                             |   |                                |                   | §§                                |
| 11138-66-2    | Xanthan gum  | Yes  |                             |   |                                |                   | §§§                               |
|               | Xylenes  | Yes  | Yes                         | 10  |                                | Table 1,5         | 0.005                             |
| 13601-19-9    | Yellow Sodium of Prussiate                               | Yes  |                             |   |                                |                   |                                   |
| 07440-66-6    | Zinc   |  | Yes                         |   | Table 6                        | Tables 1,5        | 5.0                               |
|               | Zirconium  |  | Yes                         |   |                                |                   | 0.05                              |

<sup>&</sup>lt;sup>294</sup> Several volatile compounds regulated via SPDES Table 6. Need to evaluate constituents.

| CAS<br>Number | Parameter Name | Used in<br>Additives <sup>285,</sup><br>286 | Found in<br>Flowback<br>287 | USEPA<br>MCL or TT<br>(mg/L) <sup>288</sup> ,<br>289 | SPDES<br>Tables <sup>290</sup> | TOGS111<br>Tables | NYS MCL,<br>(mg/L) <sup>291</sup> |
|---------------|----------------|---|-----------------------------|--|--------------------------------|-------------------|-----------------------------------|
|               |                |   |                             |  |                                |                   | §,§§                              |
|               |                |   |                             |  |                                |                   |                                   |
|               |                |   |                             |  |                                |                   |                                   |
|               |                |   |                             |  |                                |                   |                                   |
|               |                |   |                             |  |                                |                   |                                   |

## 6.1.3.4 Potential Impacts to Primary and Principal Aquifers

An uncontained and unmitigated surface spill could result in rapid contamination of a portion of a Primary or Principal aquifer.

Aside from the NYC Watershed and water supply system, about one half of New Yorkers rely on groundwater as a source of potable water. To enhance regulatory protection in areas where groundwater resources are most highly productive and vulnerable, NYSDOH identified categories of areas for use in geographic targeting. In order of priority, these areas are designated as follows: public water supply wellhead areas; primary water supply aquifer areas; principal aquifer areas; and other areas. The Department's Division of Water Technical & Operational Guidance Series (TOGS) 2.1.3 clarifies the meaning of Primary Water Supply Aquifer (also referred to as a Primary Aquifer) and Principal Aquifer. TOGS 2.1.3 further defines "highly vulnerable" areas as "aquifers which are highly susceptible to contamination from human activities at the land surface over the identified aquifer." This TOGS also further defines "highly productive" aquifers as those "with capability to provide water for public water supply of a quantity and natural background quality which is of regional significance."

NYSDOH identified eighteen Primary Aquifers across New York State, defined in TOGS 2.1.3 as "highly productive aquifers presently utilized as sources of water supply by major municipal water supply systems." Primary Aquifers are generally capable of providing more than 100 gallons of drinking water per minute from an individual well.

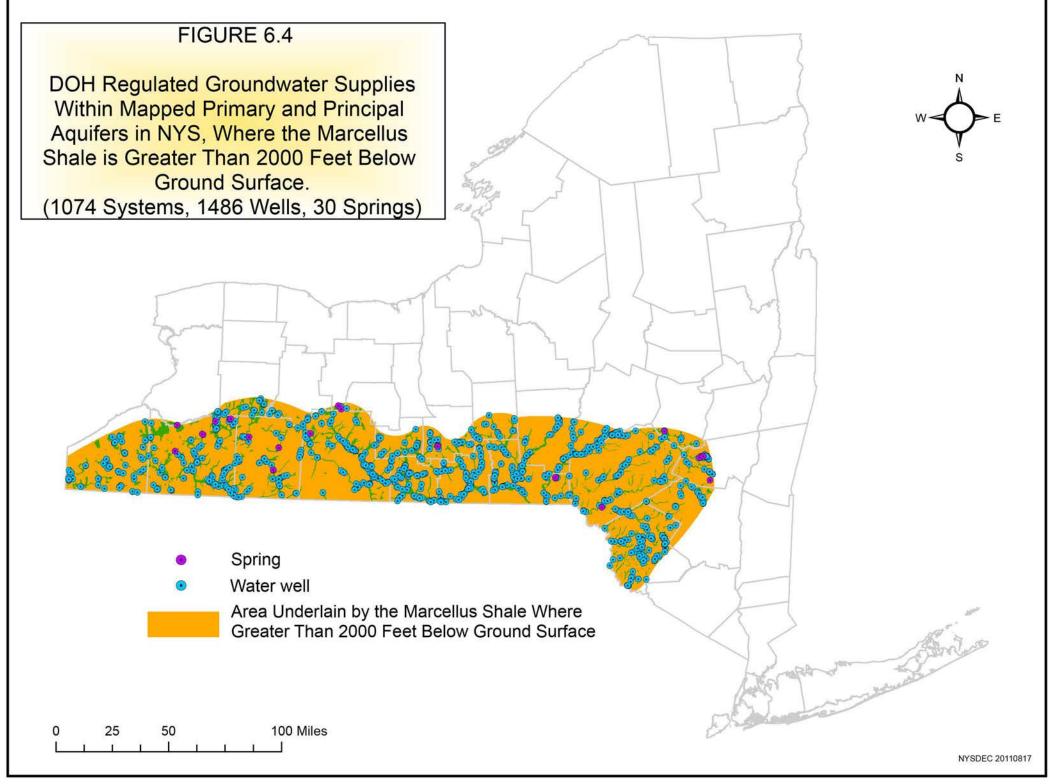
NYSDOH has also identified Principal Aquifers, which are defined in the TOGS as "highly productive but which are not intensively used as sources of water supply by major municipal systems at the present time." The TOGS further states that these areas need special protections, but awards Principal Aquifers a slightly lower priority than that afforded Primary Aquifers. Principal Aquifers are used by individual households, as well as smaller public water supply systems, such as schools or restaurants. However, Principal Aquifers are generally capable of providing 10 to 100 or more gpm of drinking water. Principal Aquifers could become Primary Aquifers depending on future public water supply use.

The groundwater table in the Primary and Principal Aquifers generally ranges from 0 to 20 feet in depth, and is overlain with sands and gravels. Because Primary and Principal Aquifers are

largely located and contained in unconsolidated material (i.e., sand and gravel), the high permeability of soils that overlie these aquifers and the shallow depth to the water table make these aquifers particularly susceptible to contamination from surface activity. TOGS 2.1.3 notes that the aquifer designations provide a rationale for enhancing regulatory protections beyond those provided by existing programs including the SPDES, Chemical Bulk Storage, and Solid and Hazardous Wastes.

The Department has issued regulations prohibiting installation of certain facilities that threaten these aquifers. For example, 6 NYCRR Part 360 "Solid Waste Facilities" provides that landfills are generally not permitted to be constructed above, or within, Primary or Principal Aquifer areas. Likewise, the Department has, since 1982, inserted special conditions into permits for drilling oil, gas and other ECL 23 wells within the boundaries of these aquifers.

As an example of the number and distribution of public supply systems that rely on Primary and Principal Aquifers within areas that could be developed by high-volume hydraulic fracturing, Figure 6.4 depicts public water supply systems that draw from Primary and Principal Aquifers within the area underlain by the Marcellus Shale where the shale occurs at a depth of at least 2,000 feet below the ground surface. The Primary Aquifer areas in this area follow the major river valleys, and serve hundreds of public water supplies, including a number of significantly sized municipalities, such as Binghamton and Endicott, as well as their surrounding areas. There are approximately 1,074 public supply systems that rely on Primary and Principal Aquifers in this area, and the total population served by these combined water supplies is at least 544,740. The total population within the area is approximately 906,000. Therefore, roughly 60.1% of the population in this prospective area is served by community groundwater supplies that draw from Primary and Principal Aquifer areas. The remainder of the population in this area is served by individual private wells or public surface water supplies or community supplies outside of Primary and Principal Aquifer areas.



The Department is chiefly concerned with surface contamination in Primary and Principal Aquifer areas because of the risk that uncontained and unmitigated surface spills could reach the aquifer in a short amount of time, due to the permeable character of the soils above the aquifers, and the shallow depth to the aquifers (generally 0-20 feet below the ground). Water quality management programs for such aquifers focus on preventing contaminants from reaching the waters in the first instance, because once they become contaminated, it is difficult and expensive to reclaim an aquifer as a source of drinking water.

As discussed elsewhere, detailed well pad containment requirements and setbacks proposed for high-volume hydraulic fracturing are likely to effectively contain most surface spills at and in the vicinity of well pads. Nevertheless, despite the best controls, there is a risk of releases to Primary or Principal Aquifers of chemicals, petroleum products and drilling fluids from the well pad.

Therefore, the Department concludes that high-volume hydraulic fracturing operations have the potential to cause a significant adverse impact to the quality of the drinking water resources provided by Primary and Principal Aquifers, even if the risk of such events is relatively small.

# Conclusion

The Department finds that the proposed high-volume hydraulic fracturing operations, although temporary in nature, may pose risks to Primary and Principal Aquifers that are not fully mitigated by the measures identified in this SGEIS.

The proposed activity could result in a degradation of drinking water supplies from accidents, construction activity, runoff and surface spills. Accordingly, the Department concludes that high-volume hydraulic fracturing operations within Primary and Principal Aquifers pose the risk of causing significant adverse impacts to water resources. As discussed in Chapter 7, standard mitigation measures may only partially mitigate such impacts. Such partial mitigation would be unacceptable due to the potential consequences posed by such impacts.

6.1.4 Groundwater Impacts Associated With Well Drilling and Construction

The wellbore being drilled, completed or produced, or a nearby wellbore that is ineffectively sealed, has the potential to provide subsurface pathways for groundwater pollution from well drilling, flowback or production operations. Pollutants could include:

- turbidity;
- fluids pumped into or flowing from rock formations penetrated by the well; and
- natural gas present in the rock formations penetrated by the well.

These potential impacts are not unique to horizontal wells and are described by the 1992 GEIS. The unique aspect of the proposed multi-well development method is that continuous or intermittent activities would occur over a longer period of time at any given well pad. This does not alter the per-well likelihood of impacts from the identified subsurface pathways because existing mitigation measures apply on an individual well basis regardless of how many wells are drilled at the same site. Nevertheless, the potential impacts are acknowledged here and enhanced procedures and mitigation measures are proposed in Chapter 7 because of the concentrated nature of the activity on multi-well pads and the larger fluid volumes and pressures associated with high-volume hydraulic fracturing. As mentioned earlier, the 1992 GEIS addressed hydraulic fracturing in Chapter 9, and NYSDOH's review did not identify any potential exposure situations associated with horizontal drilling and high-volume hydraulic fracturing that are qualitatively different from those addressed in the 1992 GEIS.

### *6.1.4.1 Turbidity*

The 1992 GEIS stated that "review of Department complaint records revealed that the most commonly validated impact from oil and gas drilling activity on private water supplies was a short-term turbidity problem." This remains the case today. Turbidity, or suspension of solids in the water supply, can result from any aquifer penetration (including monitoring wells, water wells, oil and gas wells, mine shafts and construction pilings) if sufficient porosity and permeability or a natural subsurface fracture is present to transmit the disturbance. The majority of these situations correct themselves in a short time.

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<sup>&</sup>lt;sup>295</sup> NYSDEC 1992, GEIS, p. 47.

### 6.1.4.2 Fluids Pumped Into the Well

Fluids for hydraulic fracturing are pumped into the wellbore for a short period of time per fracturing stage, until the rock fractures and the proppant has been placed. For each horizontal well the total pumping time is generally between 40 and 100 hours. ICF International, under its contract with NYSERDA to conduct research in support of SGEIS preparation, provided the following discussion and analysis with respect to the likelihood of groundwater contamination by fluids pumped into a wellbore for hydraulic fracturing (emphasis added): <sup>296</sup>

In the 1980s, the American Petroleum Institute (API) analyzed the risk of contamination from properly constructed Class II injection wells to an Underground Source of Drinking Water (USDW) due to corrosion of the casing and failure of the casing cement seal. Although the API did not address the risks for production wells, production wells would be expected to have a lower risk of groundwater contamination due to casing leakage. Unlike Class II injection wells which operate under sustained or frequent positive pressure, a hydraulically fractured production well experiences pressures below the formation pressure except for the short time when fracturing occurs. During production, the wellbore pressure would be less than the formation pressure in order for formation fluids or gas to flow to the well. Using the API analysis as an upper bound for the risk associated with the injection of hydraulic fracturing fluids, the probability of fracture fluids reaching a USDW due to failures in the casing or casing cement is estimated at less than 2 x 10-8 (fewer than 1 in 50 million wells).

More recently, regulatory officials from 15 states have testified that groundwater contamination as a result of hydraulic fracturing, which includes this pumping process, has not occurred (Appendix 15).

### 6.1.4.3 Natural Gas Migration

As discussed above, turbidity is typically a short-term problem which corrects itself as suspended particles settle. The probability of groundwater contamination from fluids pumped into a properly-constructed well is very low. Natural gas migration is a more reasonably anticipated risk posed by high-volume hydraulic fracturing. The 1992 GEIS, in Chapters 9, 10 and 16, describes the following scenarios related to oil and gas well construction where natural gas could migrate into potable groundwater supplies:

<sup>&</sup>lt;sup>296</sup> ICF Task 1, 2009, p. 21.

- Inadequate depth and integrity of surface casing to isolate potable fresh water supplies from deeper gas-bearing formations;
- Inadequate cement in the annular space around the surface casing, which may be caused by gas channeling or insufficient cement setting time; gas channeling may occur as a result of naturally occurring shallow gas or from installing a long string of surface casing that puts potable water supplies and shallow gas behind the same pipe; and
- Excessive pressure in the annulus between the surface casing and intermediate or
  production casing. Such pressure could break down the formation at the shoe of the
  surface casing and result in the potential creation of subsurface pathways outside the
  surface casing. Excessive pressure could occur if gas infiltrates the annulus because of
  insufficient production casing cement and the annulus is not vented in accordance with
  required casing and cementing practices.

As explained in the 1992 GEIS, potential migration of natural gas to a water well presents a safety hazard because of its combustible and asphyxiant nature, especially if the natural gas builds up in an enclosed space such as a well shed, house or garage. Well construction practices designed to prevent gas migration would also form a barrier to other formation fluids such as oil or brine. Although gas migration may not manifest itself until the production phase, its occurrence would result from well construction (i.e., casing and cement) problems.

The 1992 GEIS acknowledges that migration of naturally-occurring methane from wetlands, landfills and shallow bedrock can also contaminate water supplies independently or in the absence of any nearby oil and gas activities. Section 4.7 of this document explains how the natural occurrence of shallow methane in New York can affect water wells, which needs to be considered when evaluating complaints of methane migration that are perceived to be related to natural gas development.

# 6.1.5 Unfiltered Surface Drinking Water Supplies: NYC and Syracuse

There are two major surface drinking water sources and systems located within New York that have been granted permission by EPA and NYSDOH to operate as unfiltered drinking water supplies pursuant to regulations promulgated under the federal SDWA, known as the Surface Water Treatment Rule (SWTR). These unfiltered systems are the NYC and City of Syracuse water supplies and associated watersheds. For a drinking water system to qualify for filtration avoidance under the SWTR, the system cannot be the source of a waterborne disease outbreak, must meet source water quality limits for coliform and turbidity and meet coliform and total

trihalomethane MCLs in finished water. Disinfectant residual levels and redundant disinfection capability also must be maintained. Filtration avoidance further requires that a watershed control program be implemented to minimize microbial contamination of the source water. This program must characterize the watershed's hydrology, physical features, land use, source water quality and operational capabilities. It must also identify, monitor and control manmade and naturally occurring activities that are detrimental to water quality. The watershed control program must also be able to control activities through land ownership or written agreements.

Heightened public health sensitivities are associated with unfiltered surface water systems because the only treatment that these drinking waters receive before human consumption is basic disinfection through such methods as chlorine addition or ultraviolet light irradiation. In unfiltered systems, there is no application of widely employed treatment measures such as chemical coagulation/flocculation or physical filtration to remove pathogens, sediments, organic matter or other contaminants from the drinking water.

The NYC drinking water supply watershed (NYC Watershed) is located in portions of Delaware, Dutchess, Greene, Putnam, Schoharie, Sullivan, Ulster and Westchester Counties.

Approximately 9.4 million residents rely on the NYC water supply: 8.4 million in NYC and 1 million in portions of Orange, Putnam, Ulster and Westchester Counties. The NYC Watershed contains 19 reservoirs and 3 controlled lakes that supply, on average, 1.1 to 1.3 billion gallons of potable water daily. Historically, 90% of this system's drinking water has been supplied by the "Catskill" and "Delaware" portions of the NYC Watershed, which are located west of the Hudson River (an area that may be described as the "Catskill/Delaware Watershed"). On average, the remaining 10% of the water supply flows from the "Croton" portion of the NYC Watershed that is located in the counties to the east of the Hudson River. An extensive system of aqueducts and tunnels transmit waters by gravity throughout the NYC Watershed and water supply system. The NYC Watershed covers 2,000 square miles, an area that comprises 4.2% of the total land area of New York State.

Eight of the reservoirs located in the Croton portion of the NYC Watershed have been formally determined by the Department, pursuant to Clean Water Act sec. 303(d), to be impaired due to excess nutrient phosphorus (Amawalk, Croton Falls, Diverting, East Branch, Middle Branch, Muscoot, New Croton and Titicus Reservoirs). Designation as "impaired" means that these

reservoirs are in a condition that violates state water quality standards due to a specified pollutant. The Cannonsville Reservoir in Delaware County previously had been declared to be impaired due to excess nutrient phosphorus; however, its status was improved by active water quality remedial management efforts, including wastewater treatment plant upgrades, septic system repairs and replacements, construction of stormwater retrofits, and installation of best management practices on several hundred farms located throughout the Catskill and Delaware Watershed, most notably in Delaware County. As a result of this comprehensive and aggressive watershed protection program, the Department has determined that the Cannonsville Reservoir has been returned to regulatory compliance. The two reservoirs located in the Catskill portion of the NYC Watershed have been determined by the Department to be impaired due to excessive levels of suspended sediment (Ashokan and Schoharie Reservoirs).

The most recent EPA Filtration Avoidance Determination (FAD) was granted to NYC by EPA, in consultation with NYSDOH, in 2007 for the unfiltered use of the Catskill and Delaware systems and interconnected reservoir basins located in watershed communities to the east of the Hudson River. Waters flowing from the Croton portion of the NYC Watershed have been required to be filtered by EPA (at a cost of approximately \$3 billion for construction of the filtration plant). Systems of aqueducts and interchanges, however, allow for Croton waters to be transferred and intermixed with waters from the Catskill and Delaware systems to assure an adequate water supply in stressed or emergency situations, such as significant drought or major infrastructure failure.

The City of Syracuse, with a population of approximately 145,000, has also been granted permission by EPA and NYSDOH to operate an unfiltered drinking water supply. The most recent filtration avoidance determination was issued by NYSDOH to Syracuse in 2004. The unfiltered source water is Skaneateles Lake, a Finger Lake that is located approximately 20 miles to the south and west of Syracuse. The Skaneateles Lake watershed comprises a total area of 59 square miles that includes the lake - which is approximately 14 miles long and 1 mile wide. Reports issued by the Syracuse Department of Water state that Skaneateles Lake generally provides between 32 and 34 million gallons of potable water daily. The most recent NYSDOH source water assessment found that Skaneateles Lake had a moderate susceptibility to contamination, including a level of farm pasture land that results in a high potential for protozoan

contamination. Copper sulfate treatments are at times administered to Skaneateles Lake to control phosphorus-induced algae growth and associated adverse impacts such as poor taste and odor.

## 6.1.5.1 Pollutants of Critical Concern in Unfiltered Drinking Water Supplies

One of the fundamental concepts framing the effective protection of unfiltered drinking water is "source water protection." Management programs in such watershed necessarily focus on systematically preventing contaminants from reaching the waters in the first instance, as there is no mechanism in place (such as a filtration plant) to remove contaminants once they have entered the water. Once polluted, it very difficult and very expensive to return these water supplies back to their original condition. In both the NYC and City of Syracuse watersheds, extensive efforts have been undertaken to stringently treat sewage discharges. Within the Skaneateles Lake watershed, any discharge, whether treated sewage effluent or otherwise, to any surface water is prohibited. Within the NYC Watershed, all sewage treatment plants must achieve an extraordinarily stringent level of treatment consistent with "tertiary treatment, microfiltration and biological phosphorus removal." These are the most technologically advanced sewage treatment plants in New York State. Therefore, the critical remaining potential for impairment of these two unfiltered water supplies stems from human activities that place contaminants on the ground that can then be washed into reservoirs and tributaries via storm water runoff, or flow into them from contaminated groundwater.

The National Research Council of the National Academies of Sciences undertook a detailed assessment of the risks and sensitivities associated with the NYC Watershed and water supply system. This peer-reviewed report provides useful background on the distinctive nature of risks resulting from potential surface pollution in unfiltered drinking water watersheds and supplies.<sup>297</sup> The concerns and management methods discussed in this report are also relevant and applicable to the City of Syracuse drinking water supply.

In general, the pollutants of key concern when managing an unfiltered drinking water system are: (i) nutrient phosphorus; (ii) microbial pathogens; (iii) suspended sediment (or "turbidity"); and

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<sup>&</sup>lt;sup>297</sup> National Research Council, 2000.

(iv) toxic compounds. As explained below, the adverse impacts of these contaminants are substantially heightened in unfiltered drinking water systems.

Phosphorus: Excess phosphorus leads to algae blooms, including increased growth of toxin emitting blue-green algae. Algae blooms lead to high bacteria growth (due to bacterial consumption of algae) that, in turn, deplete the reservoir bottom waters of dissolved oxygen. Low dissolved oxygen suffocates or drives off fish. Low oxygen levels cause a change in the biology of reservoir waters (to anaerobic conditions) that result in impaired water taste, odor, and color. For example, iron, manganese and H<sub>2</sub>S are brought into the water column under these low oxygen conditions. The higher levels of dead algae, bacteria and other chemicals in the water constitute an increase in organic matter that can react with chlorine during the drinking water disinfection process - causing elevated levels of "disinfection by-products"; many of these chlorinated organic compounds are suspected by the EPA of being carcinogens and have been identified in a number of medical studies as a factor linked to early term miscarriage. Finally, the increased material suspended in water, which results from phosphorus-induced algae blooms, can interfere with the effectiveness of chlorination and ultraviolet light irradiation on pathogens, and thereby foster the transport waterborne pathogens to water consumers.

Phosphorus is a naturally-occurring element that is found in human and animal wastes, animal and plant materials, fertilizers and eroded soil particles. While essential for life, excess phosphorus at very low levels can cause the adverse environmental and public health impacts discussed above during the warm weather growing season. Guidance value concentrations, set by the Department to limit adverse impacts from phosphorus in NYC Watershed reservoirs, range between 15 and 20 parts per billion (ppb).

**Microbial Pathogens:** A surface drinking water source may be adversely impacted by a range of disease-causing microorganisms such as bacteria, viruses and protozoa. Such organisms can result from a variety of sources but to a significant extent result from human and animal wastes or possible re-growth in bio-slimes that may form within a drinking water supply system. Both the NYC and Syracuse drinking water supplies are required by EPA and NYSDOH regulations to employ two forms of disinfection in series that, when combined with effective source water protection programs, are highly effective in destroying or de-activating bacteria, viruses and protozoa.

However, there are two disinfection-resistant protozoa that have emerged in recent decades that can cause significant intestinal illness in otherwise healthy humans, and result in severe illness and even death in individuals with compromised immune systems. These protozoa, *Giardia lamblia* and *Cryptosporidium parvum*, both have life stages where they form cysts (or oocysts) that can survive standard disinfection treatments and infect human hosts. The basic public health management response to such organisms is to limit specific human and animal waste transmission pathways to waters on the landscape and to require controls that limit such occurrences as algae blooms and suspended sediments, which can assist in the transmittal of pathogens. As discussed below, inadequately effective controls will likely result in the imposition of a costly filtration requirement by EPA or NYSDOH in accordance with the SDWA and the underlying SWTR.

**Sediment or Turbidity:** Sediment laden, or turbid, water can increase the effective transportation of pathogens, serve as food for pathogens, promote the re-growth of pathogens in the water distribution system, and shelter pathogens from exposure to attack by disinfectants such as chlorine or ultraviolet light. The organic particles that are a cause of turbidity can combine with chlorine to create problematic disinfection by-products that are possible carcinogens and suspected by medical studies of increasing the risk of miscarriage.

EPA, in its SWTR, prohibits raw water turbidity measurements in unfiltered drinking water at the intake to the distribution system in excess of 5 nephelometric turbidity units (essentially, very clear water). More than one violation per year is grounds for EPA or NYSDOH to require construction of a water filtration plant. Such a plant for the Catskill and Delaware portions of the NYC water supply has been estimated to cost between \$8 to \$10 billion with an additional \$200 (plus) million a year in operational and maintenance expenses. An overview of the public health concerns raised by turbidity in drinking water are discussed in greater detail at: *U.S. EPA*, *Guidance Manual for Compliance with the Interim Enhanced Surface Water Treatment Rule: Turbidity Provisions*, Office of Water, EPA 815-R-99-010, April 1999, Chapter 7 (and numerous cited references); *see* also Kistemann, T., *et al.*, *Microbial Load of Drinking Water Reservoir Tributaries During Extreme Rainfall and Runoff*, Applied Environmental Microbiology, Vol. 68, No. 5, pp. 2188-2197 (May 2002); Naumova, E., *et al.*, *The Elderly and Waterborne* 

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<sup>&</sup>lt;sup>298</sup> 40 CFR §141.71(a)(2).

Cryptosporidium Infection: Gastroenteritis Hospitalizations Before and During the 1993 Milwaukee Outbreak, Emerging Infectious Diseases, Vol. 9 No. 4, pp. 418-425 (2003).

Toxic Compounds: Unfiltered drinking water supplies have a heightened sensitivity to chemical discharges as there is no immediately available method to remove contaminants from the drinking water source waters. Well pad containment practices and setbacks are likely to effectively contain most spills at those locations. There is a continuing risk, however, of releases from chemicals, petroleum products and drilling fluids from the well pad as a result of tank ruptures, equipment or surface impoundment failures, overfills, vandalism, accidents (including vehicle collisions), ground fires, or improper operations. Spilled, leaked or released fluids could flow to a surface water body. The intensive level of trucking activity associated with high-volume hydraulic fracturing, including the transport of chemical and petroleum products, presents an additional risk of surface water contamination due to truck accidents and associated releases. Given the topography of much of the NYC and Skaneateles Lake watersheds, many of the roadways are in immediate proximity to tributaries. Such proximity increases the risk that chemical and petroleum spills would not, or could not, be effectively intercepted before entering the drinking water supply.

### 6.1.5.2 Regulatory and Programmatic Framework for Filtration Avoidance

The basic statutory and regulatory framework applicable to unfiltered drinking water supplies is provided by the federal Safe Drinking Water Act (SDWA), 42 U.S.C. sec. 300f, et al. The SDWA directed EPA to adopt regulations requiring public water supplies using surface waters to apply filtration systems to treat their water unless protective "criteria" or "standards" could be met. Pursuant to this grant of authority, EPA issued the SWTR, 40 CFR sec. 141.71, et al. Subject to continuing oversight, EPA has delegated authority to administer the SDWA within New York to the NYSDOH pursuant to State statutory and regulatory authority that is consistent with the federal protocol.

There are numerous "filtration avoidance criteria" specified in the SWTR. These criteria must be met for a drinking water supply system to maintain its unfiltered status. The first two criteria address fecal coliform and turbidity limits in raw water before disinfection. The next four criteria address assuring the effectiveness of disinfection and the maintenance of sufficient levels of disinfection agents in the water distribution system. The next five criteria variously address

landscape control programs for *Giardia lamblia*, water supply system inspections, prohibition on waterborne disease outbreaks, and maximum contaminant level compliance for total coliform and disinfection by-products in drinking water after disinfection.

Another key provision operates to drive overarching watershed planning and protection programs, along with cooperative agreements with individuals and municipalities situated within the unfiltered watershed: "The public water system must demonstrate through ownership and/or written agreements with landowners within the watershed that it can control *all human activities which may have an adverse impact on the microbiological quality of the source water.*" 40 CFR sec. 141.71(b)(2)(iii) (emphasis added). High-volume hydraulic fracturing and associated activities are within the scope of "human activities" covered by this regulatory provision. As discussed above, human activities that increase levels of phosphorus and sediment, or heighten storm water flows that could transmit microbial pathogens into waters, would all have an "impact on the microbiological quality of the source water."

Major efforts have been undertaken to cooperatively assure equitable implementation of programs to protect the NYC Watershed and water supply. In 1997, essentially all stakeholders associated with the NYC Watershed entered into the "1997 New York City Watershed Memorandum of Agreement." This binding three volume agreement specified extensive programs with respect to land acquisition, extra-territorial regulations promulgated by NYC, the establishment of a Watershed Protection and Partnership Council, and an array of specific programs to limit pollution from septic systems, construction excavations, salt storage facilities, runoff from impervious surfaces, timber harvesting, waste water treatment plants, unstable streams and farms. An extensive and updated source water protection program also is detailed in the FAD that was issued to NYC (covering environmental infrastructure, protection and remedial water quality efforts, watershed monitoring and regulatory implementation). Protection programs, as well as programs to equitably address the concerns of local residents, were also detailed in a Department Water Supply Permit that was finalized and issued to NYC in January 2011. It is estimated that at least \$1.6 billion has been invested in NYC Watershed protection programs since 1997.

Syracuse has developed similar programs to prevent contamination of Skaneateles Lake and its watershed. Specific regulations have been developed to address a range of human activities that

could adversely impact water quality – including sewage treatment plants, septic systems, and erosion and sediment controls at construction sites. Syracuse implements a "Watershed Agricultural Program" to cooperatively limit pollution that could result from crop land and animal agricultural activities. A program of conservation easements in certain sensitive lands has also been developed to limit human activity that might harm water quality.

6.1.5.3 Adverse Impacts to Unfiltered Drinking Waters from High-Volume Hydraulic Fracturing Activities associated with high-volume hydraulic fracturing involve a significant amount of land clearing and excavation. New roads, sufficient to reach the well pad and of a design capable of handling a high volume of fully loaded truck traffic, would need to be cleared and cut. The often steep terrain of the NYC and Skaneateles Lake watersheds would necessitate a significant level of cut and fill roadway excavations, as well as soil stockpiles, that would expose soils to erosive activities. The excavation and grading of level well pads (generally ranging from 3 to 5 acres in size) to support drilling activities would create significant additional amounts of exposed soils and cut and fill excavations. Gas transmission pipelines of various sizes would necessarily be cut through the watersheds, often in straight lines and down hills in a manner that can accelerate and channelize water during precipitation events. Both the NYC Watershed and Skaneateles Lake watershed regularly receive high precipitation events that operate to mobilize exposed soil particles.

The clearing of vegetation, and the excavation and compaction of soils, associated with new roads, pipelines and drilling well pads in the NYC and Skaneateles Lake watersheds also will increase the volume and intensity of stormwater runoff, even if subject to stormwater control. While not fully "impervious" this less pervious landscape will increase runoff. Moreover, to support high volumes of truck traffic, narrow existing dirt roads may need to be paved and widened, as has been the experience in Pennsylvania. One acre of impervious surface is estimated to create the same amount of runoff as 16 acres of naturally vegetated meadow or forest. <sup>299</sup> Therefore, new impervious surfaces (as well as the substantially less-pervious surfaces created by the removal of vegetation and compaction of soils associated with construction excavations) can transmit very high volumes of stormwater relative to natural conditions that then operate to destabilize road-side ditches and streams, and cause additional erosion. As

<sup>299</sup> Schuler, 1994, p. 100.

discussed, elevated turbidity or suspended sediment levels present particular public health concerns in an unfiltered drinking water supply, a problem that already significantly affects the Catskill portion of the NYC Watershed, including the Schoharie and Ashokan Reservoirs.

As in other areas of the state, erosion and sediment control measures would significantly limit the adverse impacts of stormwater flow from construction excavations, erosion, soils compaction and increased imperviousness associated with high-volume hydraulic fracturing. However, even with such stormwater controls, the heightened sensitivity of these unfiltered watersheds make the potential for adverse impacts to water quality from sedimentation due to construction excavations significant during levels of projected peak activity. Even with state-of-the art stormwater controls a risk of increased stormwater runoff from accidents or other unplanned events cannot be entirely eliminated. The potential consequences of such events – loss of the FAD – is significant even if the risk of such events occurring is relatively small. Similarly, the risks associated with high volumes of truck traffic transporting chemical and petroleum products associated with high-volume hydraulic fracturing is inconsistent with effective protection of an unfiltered drinking water supply. This is especially so, as a number of factors, discussed above, are already operating to stress the NYC and Syracuse source waters. This concern is exemplified by an extensive study by researchers from SUNY ESF and Yale published in 2008. This peerreviewed report concluded that the current rate of excavations and associated increases in impervious and less pervious surfaces within the NYC Watershed would likely result in the phosphorus impairment of all reservoirs over an approximate 20 year time frame. Hall, M., R. Germain, M. Tyrell, and N. Sampson, Predicting Future Water Quality from Land Use Change Projections in the Catskill-Delaware Watersheds, pp. 217-268 (2008) (available at http://www.esf.edu/es/faculty/hall.asp). This report does not take into consideration the accelerated development associated with high-volume hydraulic fracturing.

### 6.1.5.4 Conclusion

The Department finds that high-volume hydraulic fracturing activity is not consistent with the preservation of the NYC and Syracuse watersheds as unfiltered drinking water supplies. Even with all of the criteria and conditions identified in the revised draft SGEIS, a risk remains that significant high-volume hydraulic fracturing activities in these areas could result in a degradation of drinking water supplies from accidents, surface spills, etc. Moreover, such large scale

industrial activity in these areas, even without spills, could imperil EPA's FADs and result in the affected municipalities incurring substantial costs to filter their drinking water supply.

Accordingly, and for all of the aforementioned reasons, the Department concludes that high-volume hydraulic fracturing operations within the NYC and Syracuse watersheds pose the risk of causing significant adverse impacts to water resources. As discussed in Chapter 7, standard mitigation measures such as stormwater controls would only partially mitigate such impacts. Such partial mitigation is unacceptable due to the potential consequences – adverse impacts to human health and loss of filtration avoidance – posed by such impacts.

# 6.1.6 Hydraulic Fracturing Procedure

Concern has been expressed that potential impacts to groundwater from the high-volume hydraulic fracturing procedure itself could result from:

- wellbore failure as a result of an improperly constructed well; or
- movement of unrecovered fracturing fluid out of the target fracture formation through subsurface pathways such as:
  - o a nearby poorly constructed or improperly plugged wellbore;
  - o fractures created by the hydraulic fracturing process;
  - o natural faults and fractures; and
  - o movement of fracturing fluids through the interconnected pore spaces in the rocks from the fracture zone to a water well or aquifer.

As summarized in Section 8.4.5, regulatory officials from 15 states have recently testified that groundwater contamination from the hydraulic fracturing procedure is not known to have occurred despite the procedure's widespread use in many wells over several decades.

Nevertheless, NYSERDA contracted ICF International to evaluate factors which affect the likelihood of groundwater contamination from high-volume hydraulic fracturing. 300

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<sup>&</sup>lt;sup>300</sup> ICF Task 1, 2009,

#### 6.1.6.1 Wellbore Failure

As described in Section 6.1.4.2, the probability of fracture fluids reaching an underground source of drinking water (USDW) from properly constructed wells due to subsequent failures in the casing or casing cement due to corrosion is estimated at less than 2 x 10<sup>-8</sup> (fewer than 1 in 50 million wells). Hydraulic fracturing is not known to cause wellbore failure in properly constructed wells.

# 6.1.6.2 Subsurface Pathways

Reference is made in Section 5.9 to ICF International's calculations of the rate at which fracturing fluids could move away from the wellbore through fractures and the rock matrix during pumping operations under hypothetical assumptions of a hydraulic connection. Appendix 11 provides ICF's full discussion of the principles governing potential fracture fluid flow under this hypothetical condition. ICF's conclusion is that "hydraulic fracturing does not present a reasonably foreseeable risk of significant adverse environmental impacts to potential freshwater aquifers." <sup>301</sup> Specific conditions or analytical results supporting this conclusion include:

- The developable shale formations are vertically separated from potential freshwater aquifers by at least 1,000 feet of sandstones and shales of moderate to low permeability;
- The amount of time that fluids are pumped under pressure into the target formation is orders of magnitude less than the time that would be required for fluids to travel through 1,000 feet of low-permeability rock;
- The volume of fluid used to fracture a well could only fill a small percentage of the void space between the shale and the aquifer;
- Some of the chemicals in the additives used in hydraulic fracturing fluids would be adsorbed by and bound to the organic-rich shales;
- Diffusion of the chemicals throughout the pore volume between the shale and an aquifer would dilute the concentrations of the chemicals by several orders of magnitude; and
- Any flow of fracturing fluid toward an aquifer through open fractures or an unplugged wellbore would be reversed during flowback, with any residual fluid further flushed by flow from the aquifer to the production zone as pressures decline in the reservoir during production.

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<sup>&</sup>lt;sup>301</sup> ICF Task 1, 2009, p. 34

As noted in Section 2.3.6, a depth of 850 feet to the base of potable water is a commonly used and practical generalization for the maximum depth of potable water in New York. Alpha Environmental, under its contract with NYSERDA, provided the following additional information regarding the Marcellus and Utica Shales:<sup>302</sup>

The Marcellus and Utica Shales dip southward from the respective outcrops of each member, and most of the extents of both shales are found at depths greater than 1,000 feet in New York. There are multiple alternating layers of shale, siltstone, limestone, and other sedimentary rocks overlying the Marcellus and Utica Shales. Shale is a natural, low permeability barrier to vertical movement of fluids and typically is considered a cap rock in petroleum reservoirs (Selley, 1998) and an aquitard to groundwater aquifers (Freeze & Cherry, 1979). The varying layers of rocks of different physical characteristics provide a barrier to the propagation of induced hydraulic fractures from targeted zones to overlying rock units (Arthur et al, 2008). The vertical separation and low permeability provide a physical barrier between the gas producing zones and overlying aquifers.

# Natural Controls on Underground Fluid Migration

As noted by ICF (Subpart 5.11.1.1 and Appendix 11) and Alpha (as cited above), the developable shale formations are vertically separated from potential freshwater aquifers by at least 1,000 feet of sandstones and shales of moderate to low permeability. Figure 4.2 shows that most of the bedrock formations above the Marcellus Shale are other shales. That shales must be hydraulically fractured to produce fluids is evidence that these rocks do not readily transmit fluids. The high salinity of native water in the Marcellus and other Devonian shales is evidence that fluid has been trapped in the pore spaces for a significant length of time, implying that there is no mechanism for discharge.

As previously discussed, hydraulic fracturing is engineered to target the prospective hydrocarbon-producing zone. The induced fractures create a pathway to the intended wellbore, but do not create a discharge mechanism or pathway beyond the fractured zone where none existed before. The pressure differential that pushes fracturing fluid into the formation is diminished once the rock has fractured, and is reversed toward the wellbore during the flowback and production phases.

<sup>&</sup>lt;sup>302</sup> Alpha, 2009, p. 3-3.

Darcy's Law is a universally accepted scientific principle of hydrogeology. It states the relationship that explains fluid flow in porous media. Flow rate, Q, is calculated by

$$Q=KA(P_{high}-P_{low})/\mu L$$

where K= permeability, A= cross sectional area, P=pressure, μ=fluid viscosity and L=length of flow. The factor "Phigh-Plow" describes a pressure differential, and Darcy's Law explains the relationship between pressure and fluid flow. During hydraulic fracturing operations, the pressure in the well is greater than the pressure in the formation and drives the fluid and sand into the rock creating the induced fractures. If induced fractures do intersect an open fault or wellbore that diverts fluid from the target formation during pumping, this would be detected by required pressure monitoring during the fracturing process. Permit conditions will require pumping operations to cease if this occurs, until the anomalous condition is evaluated and addressed. Cessation of pumping will remove the pressure differential and stop further flow away from the target formation. Additionally, the force exerted by lithostatic pressure (i.e., the weight of overlying rocks) tends to close natural fissures at depth, so even when such fissures exist they are not necessarily transmissive. This is the reason that hydraulic fracturing requires the use of proppant to keep induced fractures open to transmit natural gas to the wellbore. Also, even if it is assumed that fractures in overlying strata are transmissive, there is no reason to believe that the fractures of different strata are aligned in a manner that would make hydraulic connections possible.

Once pumping ceases and hydraulic fracturing is accomplished, the well is turned into the production system at the surface which is at a much lower pressure than the formation.

Therefore gas flows to the well and the surface. At this point there is no pressure differential that would cause fluid to move in any direction other than towards the gas well.

All of the above factors that inhibit vertical fracturing fluid migration would also inhibit horizontal migration beyond the fracture zone for the distances required to impact potable water wells in the Marcellus and other shales from high-volume hydraulic fracturing under the conditions specified by ICF. Because of regional dip, the geographic location of any target reservoir where it is more than 1,000 feet below the presumed base of fresh water would be at

least several miles south of any location where water wells are completed in the same rock formation.

## Mapped Marcellus Hydraulic Fracturing Stages

Four hundred Marcellus hydraulic fracturing stages in Pennsylvania, West Virginia and Ohio have been mapped with respect to vertical growth and distance to the deepest water wells in the corresponding areas. Although many of the hydraulic fracturing stages occurred at depths greater than the depths at which the Marcellus occurs in New York, the results across all depth ranges showed that induced fractures did not approach the depth of drinking water aquifers. In addition, as previously discussed, at the shallow end of the target depth range in New York, fracture growth orientation would change from vertical to horizontal.

# 6.1.7 Waste Transport

Drilling and fracturing fluids, mud-drilled cuttings, pit liners, flowback water and production brine are classified as non-hazardous industrial-commercial waste which would be hauled under a New York State Part 364 waste transporter permit issued by the Department. All Part 364 transporters would identify the general category of wastes transported and obtain written authorization from each destination facility, which must be maintained at the place of business and made available to the Department upon request.

Manifesting is not required for non-hazardous industrial-commercial waste, so there is no tracking and verification of disposal destination on an individual load basis. Although the Department's regulations do not classify drilling and production wastes as hazardous, like all wastes they must be handled and disposed of in accordance with all applicable regulatory requirements. One concern is that wastes will not be properly identified or may not be taken to appropriate, permitted facilities. Chapter 7 provides mitigation for this concern in the form of a waste tracking procedure similar to that which is required for medical waste even though the hazards are not equivalent. Another concern relates to potential spills as a result of trucking accidents. It should be noted that the developing practice of treating and reusing flowback water on the same well pad would reduce the number of truck trips for hauling flowback water to other

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<sup>&</sup>lt;sup>303</sup> Fisher, 2010, pp. 30-33.

destinations. Information about traffic management related to high-volume hydraulic fracturing is presented in Section 7.8.

# 6.1.8 Fluid Discharges

Direct discharge of fluids onto the ground or into surface water bodies from the well pad are prohibited. Discharges would be managed at treatment facilities, appropriately recycled, or in permitted disposal wells.

### 6.1.8.1 POTWs

Surface water discharges from water treatment facilities are regulated under the Department's SPDES program. Acceptance by a POTW of a waste stream that upsets its system or exceeds its capacity may result in a SPDES permit effluent violation or a violation of water quality standards within the receiving water. Water pollution degrades surface waters, potentially making them unsafe for drinking, fishing, swimming, and other activities or unsuitable for their classified best uses.

Flowback water may be sent to POTWs. However, treatability of flowback water presents a potential environmental concern because residual fracturing chemicals and naturally-occurring constituents from the rock formation could be present in flowback water and have treatment, sludge disposal, and receiving-water impacts. Salts and dissolved solids may not be sufficiently treated by municipal biological treatment and/or other treatment technologies which are not designed to remove pollutants of this nature. Table 6.1\_provides information on flowback water composition based on a limited number of samples from Pennsylvania and West Virginia.

Appendix 21 is a list of POTWs with approved pretreatment and mini-pretreatment programs. Note that this is not a list of facilities approved to accept wastewater from high-volume hydraulic fracturing. Rather, it is a list of facilities that have SPDES permit conditions and requirements allowing them to accept wastewater from hauled or other significant industrial sources in accordance with 40CFR Part 403. To accept a source of wastewater, the facility must first evaluate the pollutants present in that source of wastewater against an analysis of the capabilities of the individual treatment units and the treatment system as a whole to treat these pollutants; that analysis is known as a Maximum Allowable Headworks Loading analysis (MAHW, or headworks analysis). In addition, any industrial wastewater source, including this source of

wastewater, may only be discharged utilizing all treatment processes within the POTW. Admixture of untreated flowback water or other well development water to the treated effluent of the POTW is not allowed. Improper handling could result in noncompliance with terms of the permit or the ECL and result in formal enforcement actions.

The large volumes of return water from high-volume hydraulic fracturing combined with the diverse mixture of chemicals and high concentrations of TDS that exist in both flowback water and production brine, requires that the permittee submit a headworks analysis specific to the parameters expected present in high-volume hydraulic fracturing wastewater, including TDS and NORM, to both the Department and EPA Region 2 for review in accordance with DOW's Technical and Operational Guidance Series (TOGS) 1.3.8, New Discharges to Publicly Owned Treatment Works. TOGS 1.3.8., was developed to assist Department permit writers in evaluating the potential effect of a new, substantially increased, or changed non-domestic discharge to a POTW on that facility's SPDES permit and pretreatment program. The DOW and EPA must determine whether the POTW has adequately evaluated the effects of the proposed discharge on POTW operation, sludge disposal, effluent quality, and POTW health and safety; whether the discharge will result in the discharge of a substance that will be subject to effluent limits, action levels, or other monitoring requirements in the facility's SPDES permit; and whether the proposed discharge contains any Bioaccumulative Chemicals of Concern or persistent toxic substances that may be subject to SPDES effluent limits or other Departmental permit requirements or controls. Appendix C of TOGS 1.3.8, Guidance for Acceptance of New Discharges, describes the analyses and submittals necessary for a POTW to accept a new source of wastewater. Note that if a facility has a currently approved headworks analysis in place for the parameters and concentrations of those parameters typically found in flowback water and production brine, the permittee may assess the impacts of the proposed discharge against the existing headworks analysis.

The Department proposes to require, as a permit condition, that the permittee demonstrate that it has a source to treat or otherwise legally dispose of wastewater associated with flowback and production <u>brine</u> prior to the issuance of the drilling permit. Disposal and treatment options include publicly owned treatment works, privately owned high volume hydraulic fracturing wastewater treatment and/or reuse facilities, deep-well injection, and out of state disposal.

Flowback water and production brine must be fully characterized prior to acceptance by a POTW for treatment. Note in particular Appendix C. IV of TOGS 1.3.8, Maximum Allowable Headworks Loading. The POTW must perform a MAHW analysis to assure that the flowback water and production brine will not cause a violation of the POTW's effluent limits or sludge disposal criteria, allow pass through of unpermitted substances or inhibit the POTW's treatment processes. As a result, the SPDES permits for POTWs that accept this source of wastewater will be modified to include influent and effluent limits for Radium and TDS, if not already included in the existing SPDES permit, as well as for other parameters as necessary to ensure that the permit correctly and completely characterizes the discharge. In the case of NORM, anyone proposing to discharge flowback water or production brine to a POTW must first determine the concentration of NORM present in those waste streams to determine appropriate treatment and disposal options. POTW operators who accept these waste streams are advised to limit the concentrations of NORM in the influent to their systems to prevent its inadvertent concentration in their sludge. For example, due to the potentially large volumes of these waste waters that could be processed through any given POTW, as well as the current lack of data on the level of NORM concentration that may take place, it will be proposed that POTW influent concentrations of radium-226 (as measured prior to admixture with POTW influent) be limited to 15 pCi/L, or 25% of the 60 pCi/L concentration value listed in 6 NYCRR Part 380-11.7. As more data become available on concentrations in influent vs. sludge it is possible that this concentration limit may be revisited.

Specific information regarding high volume hydraulic fracturing additives, such as chemical makeup and aquatic toxicity, will be required for this analysis. A complete listing of all ingredients in each chemical additive to be used shall be included as part of a headworks analysis, along with aquatic toxicity data for each of the additives. If any confidentiality is allowed under State law based upon the existence of proprietary material, that fact may be noted in the submission. However, in no circumstance shall a fracturing additive be approved or evaluated in a headworks analysis without aquatic toxicity data. Department approval of the headworks analysis, and the modification of the POTW's SPDES permit if necessary, must be received prior to the acceptance of flowback water or production brine from wells permitted pursuant to this Supplement.

In conducting the headworks analysis, the parameters that must be analyzed include, at a minimum:

- pH, range, SU;
- Oil and Grease;
- Solids, Total Suspended;
- Solids, Total Dissolved;
- Chloride;
- Sulfate;
- Alkalinity, Total (CaCO<sub>3</sub>);
- BOD, 5 day;
- Chemical Oxygen Demand (COD);
- Total Kjeldahl Nitrogen (TKN);
- Ammonia, as N;
- Total Organic Carbon;
- Phenols, Total;
- the following scans:
  - o Priority Pollutants Metals;
  - o Priority Pollutants VOC;
  - o Priority Pollutants SVOC Base/Neutral; and
  - Priority Pollutants SVOC Acid Extractable;
- Radiological analysis including:
  - o Gross Alpha EPA Method 900.0, Standard Methods 7110-B;
  - o Gross Beta EPA Method 900.0, Standard Methods 7110-B;

- o Radium EPA Method 903.0, Standard Methods 7500-Ra B;
- o Uranium EPA Method 908, Standard Methods 7500-U; and
- o Thorium EPA Method 910, Standard Methods 7500-Th;
- constituents that were present in the hydraulic fracturing additives.

The high concentrations of TDS present in this source of wastewater may prove to be inhibitory to biological wastewater treatment systems. It has been noted that the concentrations of TDS in the return and process water increase as a higher percentage of native water is produced and then stabilize over the life of the well. The expected concentrations of TDS for both the initial flowback water as well as for the ongoing well operation must therefore be considered in the development of the headworks analysis. It is incumbent upon the POTW to determine whether the volumes and concentrations of chemicals present in the flowback water or production brine would result in adverse impacts to the facility's treatment processes as part of the above headworks analysis.

The Department has performed a very basic analysis to determine the potential available capacity for POTWs to accept high-volume hydraulic fracturing wastewater. The Department estimates that the POTWs within the approximate area of shale development in New York have an aggregate available flow capacity of approximately 300 MGD, which is the difference between existing flow and permitted flow. Based on this capacity, an estimate was developed to determine the existing total treatment capacity based on the actual flows, existing TDS levels and allowable TDS discharge limits. This estimate was based on a conservative assumption of influent TDS from production brine. This estimate assumes that all of these POTWs would be willing to accept this wastewater to their maximum available capacity, and that no other increased discharges or other growth in the service area are expected. A TDS level of 350,000 mg/L will be used, as this is on the upper end of expected concentrations. Discharge levels from POTWs would be limited to 1,000 mg/L. Typical influent levels of TDS at a POTW are approximately 300 mg/L. Therefore, a typical POTW can be expected to have a disposal capacity of approximately 700 mg/L (1,000 – 300mg/L) of TDS. Again assuming an influent level of 350,000 mg/L of TDS and a disposal capacity of 700 mg/L at an existing POTW, the dilution ratio of existing POTW flow to allowable high-volume hydraulic fracturing wastewater influent flow is 500:1 (350,000 divided by 700). Based on this analysis, the maximum total

capacity for disposal of high-volume hydraulic fracturing wastewater is estimated to be less than 1 MGD. The estimated production <u>brine</u> per well may range from 400 gpd to 3,400 gpd depending on the life of the well.

The above analysis is subject to a number of assumptions which, when actual conditions are factored in, will limit the available capacity to much less than 1 MGD. The analysis assumes that the treatment facilities are willing to accept this source of wastewater; following its December 2008 letter to POTWs outlining the requirements to accept high-volume hydraulic fracturing wastewater, the Division of Water has yet to receive any requests from any POTW in the State to accept this source of wastewater. The analysis assumes that POTWs are equipped to take this source of wastewater and that haulers are willing to pump the waste into the POTW at the rate that will be required to protect the POTW; no POTWs in New York State currently have TDS-specific treatment technologies, so the ability to accept this wastewater is limited by influent concentration and flow rates. The analysis assumes that the receiving water has assimilative capacity to accept additional TDS loadings from POTWs and that the background TDS in the receiving water is less than the in-stream water quality standard of 500 mg/L; there are several streams in New York State which cannot accept additional TDS loads. Based on the above, there is questionable available capacity for POTWs in New York State to accept high-volume hydraulic fracturing wastewater.

Case Study: One wellpad is expected to have 8 wells. Each well is expected to produce 3,000 gallons of production <u>brine</u>. Assuming 3,000 gpd x 8 wells = 24,000 gpd. With a 500:1 ratio needed for disposal, a POTW with an existing flow of 12 mgd would be needed to dispose of the production <u>brine</u> from this single wellpad.

Further, because of the inability of biological treatment systems to remove certain high-volume hydraulic fracturing additives in flowback water, as previously described, POTWs are not usually equipped to accept influent containing these contaminants. The potential for inhibition of biological activity and sludge settling and the potential for radionuclide concentration in the sludge impacts sludge disposal options.

As noted previously, acceptance of wastewater from high-volume hydraulic fracturing operations must consider the impacts to POTW operation, sludge disposal, effluent quality, and POTW

health and safety. Concentrations of NORM, specifically radium, in natural gas drilling wastewater have the potential to impact POTW sludge disposal. At this time there is a lack of detailed information on levels of NORM in POTW sludge and to what extent NORM that is introduced to a POTW is concentrated in the sludge. Therefore, to ensure that POTW sludge disposal is not affected, an influent radium-226 limit of 15 pCi/L for high-volume hydraulic fracturing wastewater, to be determined prior to admixture with other POTW influents, would be required in SPDES permits for any POTW that proposes to accept high-volume hydraulic fracturing wastewater. It is noted that there are a number of water bodies in NY where the ambient levels of TDS already exceed the water quality standard or where TDS has already been fully allocated in existing SPDES permits. This may further limit the ability of POTWs to accept these discharges.

# 6.1.8.2 Private Off-site Wastewater Treatment and/or Reuse Facilities

Privately owned facilities built specifically for the reuse and/or treatment and disposal of industrial wastewater from high-volume hydraulic fracturing operate in other states, including Pennsylvania. Similar facilities that might be constructed in New York would require a SPDES permit if the operator of the facility intends to discharge treated effluent to surface or groundwater. The treatment methods that would be applicable to these facilities are discussed in Chapter 5. A number of adverse impacts are possible resulting from improper maintenance or overloading of these systems, resulting in either surface or water discharges that do not comply with applicable standards. However, properly maintained and regulated systems, along with waste tracking and SPDES permitting control measures as described in Chapter 7 would mitigate the potential for these impacts. The same limitations and impacts noted regarding the effects of discharges from POTWs to the waters of the State, including the ability of the receiving water to accept additional TDS loads, as described in Section 6.1.8.1 above, also apply to privately-owned off-site treatment works.

#### 6.1.8.3 Private On-site Wastewater Treatment and/or Reuse Facilities

As noted in Chapter 5 of this Draft SGEIS, on-site treatment of flowback water for purposes of reuse is currently being used in Pennsylvania and other states. The treated water is blended with fresh water at the well site and reused for hydraulic fracturing, with the treatment system residue hauled off-site. A number of adverse impacts are possible resulting from improper maintenance

or overloading of these systems, resulting in either surface or water discharges that do not comply with applicable standards. However, properly maintained and operated treatment and/or reuse systems, along with the waste tracking measures described in Chapter 7, would mitigate the potential for these impacts. Because all applicable technology-based requirements must be applied in NPDES/SPDES permits under the Clean Water Act section 402(a) and implementing regulations at 40 CFR 125.3, an NPDES/SPDES permit issued for drilling activity would need to be consistent with 40 CFR Part 435, Subpart C, which states that "there shall be no discharge of wastewater pollutants into navigable waters from any source associated with production, field exploration, drilling, well completion, or well treatment (i.e. production brine, drilling muds, drill cuttings, and produced sand."

## 6.1.8.4 Disposal Wells

As stated in the 1992 GEIS, the primary environmental consideration with respect to disposal wells is the potential for movement of injected fluids into or between potential underground sources of drinking water. The Department is not proposing to alter its 1992 Finding that proposed disposal wells require individual site-specific review. Therefore, the potential for significant adverse environmental impacts from any proposal to inject flowback water from high-volume hydraulic fracturing into a disposal well would be reviewed on a site-specific basis with consideration to local geology (including faults and seismicity), hydrogeology, nearby wellbores or other potential conduits for fluid migration and other pertinent site-specific factors.

#### 6.1.8.5 Other Means of Wastewater Disposal

Wastewater generated by high-volume hydraulic fracturing would be able to be treated and disposed of to the extent that available capacity exists using the disposal options referenced in Section 6.1.8.4 above. Should wastewater be generated in volumes exceeding available capacity within the State, the wastewater would require transport and disposal at facilities not located in New York State, or additional treatment facilities to be constructed. Potential impacts that may result from insufficient wastewater treatment capacity would include either storage of wastewater and associated potential for leaks or spillage, illegal discharge of wastewater to the ground surface or directly to waters of the State, and increased truck traffic resulting from transport of wastewater to out of state treatment and disposal facilities.

## 6.1.9 Solids Disposal

Most waste generated at a well site is in liquid form. Rock cuttings and the reserve pit liner are the significant exception. The 1992 GEIS describes potential adverse impacts to agricultural operations if materials are buried at too shallow a depth or work their way back up to the surface. Concerns unique to Marcellus development and multi-well pad drilling are discussed below.

## 6.1.9.1 NORM Considerations - Cuttings

Gamma ray logs from deep wells drilled in New York over the past several decades show the Marcellus Shale to be higher in radioactivity than other bedrock formations including other potential reservoirs that could be developed by high-volume hydraulic fracturing. However, based on the analytical results from field-screening and gamma ray spectroscopy performed on samples of Marcellus Shale, NORM levels in cuttings are not likely to pose a problem because – as set forth in Section 5.2.4.2 – the levels are similar to those naturally encountered in the surrounding environment.

## 6.1.9.2 Cuttings Volume

As explained in Chapter 5, the total volume of drill cuttings produced from drilling a horizontal well may be about 40% greater than that for a conventional, vertical well to the same target depth. For multi-well pads, cuttings volume would be multiplied by the number of wells on the pad. The potential water resources impact associated with the greater volume of drill cuttings from multiple horizontal well drilling operations would arise from the retention of cuttings during drilling, necessitating a larger reserve pit that may be present for a longer period of time, unless the cuttings are directed into tanks as part of a closed-loop tank system. The geotechnical stability and bearing capacity of buried cuttings, if left in a common pit, may need to be reviewed prior to pit closure. 304

## 6.1.9.3 Cuttings and Liner Associated With Mud-Drilling

Operators have not proposed on-site burial of mud-drilled cuttings, which would be equivalent to burial or direct ground discharge of the drilling mud itself. Contaminants in the mud or in contact with the liner if buried on-site could adversely impact soil or leach into shallow groundwater.

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<sup>&</sup>lt;sup>304</sup> Alpha, 2009, p. 6-7.

# 6.2 Floodplains

Flooding is hazardous to life, property and structures. Chapter 2 describes Flood Damage Prevention Laws implemented by local communities to govern development in floodplains and floodways and also provides information about recent flooding events in the Susquehanna and Delaware River Basins. The GEIS summarizes the potential impacts of flood damage relative to mud or reserve pits, production brine and oil tanks, other fluid tanks, brush debris, erosion and topsoil, bulk supplies (including additives) and accidents. Severe flooding is described as "one of the few ways" that bulk supplies such as additives "might accidentally enter the environment in large quantities." Accordingly, construction of drill pads within flood plains raises serious and significant environmental issues and risks.

#### **6.3** Freshwater Wetlands

State regulation of wetlands is described in Chapter 2. The 1992 GEIS summarizes the potential impacts to wetlands associated with interruption of natural drainage, flooding, erosion and sedimentation, brush disposal, increased access and pit location, and those potential impacts are applicable to high-volume hydraulic fracturing. Potential impacts to downstream wetlands as a result of surface water withdrawal are discussed in Section 6.1.1.4 of this Supplement. Other concerns described herein relative to stormwater runoff and surface spills and releases, also extend to wetlands.

## 6.4 Ecosystems and Wildlife

The 1992 GEIS discusses the significant habitats known to exist at the time in or near then-existing oil and gas fields (heronries, deer wintering areas, and uncommon, rare and endangered plants). Significant habitats are defined as areas that provide one or more of the key factors required for survival, variety, or abundance of wildlife, and/or for human recreation associated with such wildlife. This section considers the potential impact of high-volume hydraulic fracturing on all terrestrial habitat types, including forests, grasslands (including old fields managed for grasslands, and pasture and hay fields) and shrublands. Four areas of concern related to high-volume hydraulic fracturing are:

1) fragmentation of habitat;

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<sup>&</sup>lt;sup>305</sup> NYSDEC, 1992, GEIS, p. 8-44

- 2) potential transfer of invasive species;
- 3) potential impacts on endangered and threatened species; and
- 4) use of certain State-owned lands.

When the 1992 GEIS was developed, the scale and scope of the anticipated impact of oil and gas drilling in New York State was much different than it is today. Development of low-permeability reservoirs by high-volume hydraulic fracturing have the potential to draw substantial development into New York, which is reasonably anticipated to result in potential impacts to habitats (fragmentation, loss of connectivity, degradation, etc.), species distributions and populations, and overall natural resource biodiversity.

The development of Marcellus Shale gas will have a large footprint. In addition to direct loss of habitat, constant activity on each well pad from construction, drilling, and waste removal can be expected for 4 to 10 months, further affecting species. If a pad has multiple wells, it might be active for several years. More land is disturbed for multi-well pads, but fewer access roads, infrastructure, and total pads would be needed. Well pad sites are partially restored after drilling, but 1-3 acres is typically left open for the life of the well (as are access roads and pipelines), which is expected to be 20 to 40 years.

## 6.4.1 Impacts of Fragmentation to Terrestrial Habitats and Wildlife

Fragmentation is an alteration of habitats resulting in changes in area, configuration, or spatial patterns from a previous state of greater continuity, and usually includes the following:

<sup>&</sup>lt;sup>306</sup> Environmental Law Clinic, 2010.

- Reduction in the total area of the habitat;
- Decrease of the interior to edge ratio;
- Isolation of one habitat fragment from other areas of habitat;
- Breaking up of one patch of habitat into several smaller patches; and
- Decrease in the average size of each patch of habitat.

# General Direct, Indirect, and Cumulative Impacts:

Habitat loss, conversion, and fragmentation (both short-term and long-term) would result from land grading and clearing, and the construction of well pads, roads, pipelines, and other infrastructure associated with gas drilling.<sup>307</sup>

Habitat loss is the direct conversion of surface area to uses not compatible with the needs of wildlife, and can be measured by calculating the physical dimensions of well pads, roads, and other infrastructure. In addition to loss of habitat, other potential direct impacts on wildlife from drilling in the Marcellus Shale include increased mortality, increase of edge habitats, altered microclimates, and increased traffic, noise, lighting, and well flares. Existing regulation of wellhead and compressor station noise levels is designed to protect human noise receptors. Little definitive work has been done on the effects of noise on wildlife. 308

Habitat degradation is the diminishment of habitat value or functionality; its indirect and cumulative effects on wildlife are often assessed through analysis of landscape metrics. Indirect and cumulative impacts may include a loss of genetic diversity, species isolation, population declines in species that are sensitive to human noise and activity or dependent on large blocks of habitat, increased predation, and an increase of invasive species. Certain life-history characteristics, including typically long life spans, slow reproductive rates, and specific habitat requirements for nesting and foraging, make raptor (birds of prey) populations especially vulnerable to disturbances. Direct habitat loss has less impact than habitat degradation through

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<sup>&</sup>lt;sup>307</sup> Environmental Law Clinic, 2010.

<sup>&</sup>lt;sup>308</sup> New Mexico Dept. Game & Fish, 2007.

fragmentation and loss of connectivity due to widespread activities like oil and gas development. 309

Biological systems are exceedingly complex, and there can be serious cascading ecological consequences when these systems are disturbed. Little baseline data are available with which comparisons can later be made in the attempt to document changes, or lack thereof, due to oil and gas development. In cases where serious adverse consequences may reasonably be expected, it is prudent to err on the side of caution. 310

Habitat fragmentation from human infrastructure has been identified as one of the greatest threats to biological diversity. Research on habitat fragmentation impacts from oil and gas development specific to New York is lacking. However, the two following studies from the western United States are presented here to illustrate qualitatively the potential impacts to terrestrial habitats that could occur in New York. A quantitative comparison between these studies and potential impacts in New York is not possible because these studies were conducted under a regulatory structure that resulted in well spacing that differs from those anticipated for high-volume hydraulic fracturing in New York. Additional research would be necessary to determine the precise impacts to species and wildlife expected from such drilling in New York's Marcellus Shale.

While fragmentation of all habitats is of conservation concern, the fragmentation of grasslands and interior forest habitats are of utmost concern in New York. Some of the bird species that depend on these habitat types are declining. This decline is particularly dramatic for grasslands where 68% of the grassland-dependent birds in New York are declining.<sup>311</sup>

**Projected Direct Impacts** 

Study 1, General Discussion: The Wilderness Society conducted a study in 2008<sup>312</sup> that provided both an analytical framework for examining habitat fragmentation and results from a hypothetical GIS analysis simulating the incremental development of an oil and gas field to

<sup>309</sup> New Mexico Dept. Game & Fish, 2007.

<sup>&</sup>lt;sup>310</sup> New Mexico Dept. Game & Fish, 2007.

<sup>311</sup> Post 2006.

<sup>312</sup> Wilbert et al., 2008.

progressively higher well pad numbers over time. Results of the sample analysis gave a preliminary estimate of the minimum potential fragmentation impacts of oil and gas development on wildlife and their habitats; the results were not intended to be a substitute for site-specific analyses.

The study identified a method to measure fragmentation (landscape metrics), and a way to tie various degrees of fragmentation to their impacts on wildlife (from literature). Two fragmentation indicator values (road density and distance-to-nearest-road or well pad) were analyzed for impacts to a few important wildlife species present in oil and gas development areas across the western U.S.

**Study 1, Findings:** The total area of direct disturbance from well pads and roads used in oil and gas development was identified for a hypothetical undeveloped 120-acre site, with seven separate well-pad densities - one pad per 640 acres, 320 acres, 160 acres, 80 acres, 40 acres, 20 acres, and 10 acres:

- 1. Well pads: the disturbance area increased approximately linearly as pad density increased:
- 2. Total road length: the disturbance area increased more rapidly in the early stages of development;
- 3. Mean road density: the rate of increase was higher at earlier stages of development. The size of the pre-development road system had an effect on the magnitude of change between subsequent development stages, but the effect decreased as development density increased;
- 4. Distance-to-nearest-road (or well pad): the rate of decrease was higher at earlier stages of development than at later stages; and
- 5. Significant negative effects on wildlife were predicted to occur over a substantial portion of a landscape, even at the lower well pad densities characteristic of the early stages of development in gas or oil fields.

This suggests that landscape-level planning for infrastructure development and analysis of wildlife impacts need to be done prior to initial development of a field. Where development has already occurred, the study authors recommend that existing impacts on local wildlife species be measured and acknowledged, and the cumulative impacts from additional development be assessed.

**Study 1, Implications for New York:** The study results emphasize the importance of maintaining undeveloped areas. Note that the degree of habitat fragmentation and the associated impacts on wildlife from such development in real landscapes would be even greater than those found in the study, which used conservative estimates of road networks (no closed loops, shorter roads, and few roads pre-development) and did not include pipelines and other infrastructure.

## Projected Indirect and Cumulative Impacts

Study 2, General Discussion: The Wilderness Society conducted a study in 2002<sup>313</sup> that analyzed the landscape of an existing gas and oil field in Wyoming to identify habitat fragmentation impacts. As fragmentation of the habitat occurred over a wide area, cumulative and indirect impacts could not be adequately addressed at the individual well pad site level. Rather, analyzing the overall ecological impacts of fragmentation on the composition, structure, and function of the landscape required a GIS spatial analysis. A variety of metrics were developed to measure the condition of the landscape and its level of fragmentation, including: density of roads and linear features; acreage of habitat in close proximity to infrastructure; and acreage of continuous uniform blocks of habitat or core areas.

**Study 2, Findings:** The study area covered 166 square miles, and contained 1864 wells, equaling a density of 11 wells per square mile. The direct physical footprint of oil and gas infrastructure was only 4% of the study area; however, the ecological impact of that infrastructure was much greater. The entire study area was within one-half mile of a road, pipeline corridor, well head, or other infrastructure, while 97% fell within one-quarter mile. Study results also showed the total number, total acreage, and the percent of study area remaining in core areas decreased as the width of the infrastructure impact increased. No core areas remained within one-half mile of infrastructure, and only 27% remained within 500 feet of infrastructure. These results, combined with a review of the scientific literature for fragmentation impacts to western focal species, indicated there was little to no place in the study area where wildlife would not be impacted.

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<sup>&</sup>lt;sup>313</sup> Weller et al. 2002.

Note that this density is between that of single horizontal wells (9 per square mile) and vertical wells (16 per square mile) expected in New York (section 5.1.3.2).

**Study 2, Implications for New York:** This study demonstrated that impacts to wildlife extended beyond the direct effects from the land physically altered by oil and gas fields. Note that the overall impacts predicted in the study were likely conservative as the data were only assessed at the individual gas field scale, not the broader landscape. While well densities from multiple horizontal wells from a common pad (a minimum of 1 well pad per square mile) would be less than in this study, all three drilling scenarios might result in negative impacts to wildlife in New York, as the impacts predicted to the complement of species in Wyoming were so extreme.

## 6.4.1.1 Impacts of Grassland Fragmentation

Grassland birds have been declining faster than any other habitat-species suite in the northeastern United States. The primary cause of these declines is the fragmentation of habitat caused by the abandonment of agricultural lands, causing habitat loss due to reversion to later successional stages or due to sprawl development. Remaining potential habitat is also being lost or severely degraded by intensification of agricultural practices (e.g., conversion to row crops or early and frequent mowing of hayfields).

Stabilizing the declines of populations of grassland birds has been identified as a conservation priority by virtually all of the bird conservation initiatives, groups, and agencies in the northeastern US, as well as across the continent, due to concern over how precipitous their population declines have been across portions of their ranges (for the list of species of concern and their population trends, see Table 6.2). In New York, grassland bird population declines are linked strongly to the loss of agricultural grasslands, primarily hayfields and pastures; it is therefore critical to conserve priority grasslands in order to stabilize or reverse these declining trends.

<sup>&</sup>lt;sup>315</sup> Morgan and Burger 2008.

Table 6.2 - Grassland Bird Population Trends at Three Scales from 1966 to 2005. 316 (New July 2011)

|                                  | New               | V York                         | USFWS             | Region 5                       | Surve             | ey-wide                        |
|----------------------------------|-------------------|--------------------------------|-------------------|--------------------------------|-------------------|--------------------------------|
| Species                          | trend<br>(%/year) | population<br>remaining<br>(%) | trend<br>(%/year) | population<br>remaining<br>(%) | trend<br>(%/year) | population<br>remaining<br>(%) |
| Northern Harrier <sup>1</sup>    | -3.4              | 25.9                           | 1.1               | 153.2                          | -1.7              | 51.2                           |
| Upland Sandpiper <sup>1</sup>    | -6.9              | 6.2                            | -0.7              | 76.0                           | 0.5               | 121.5                          |
| Short-eared Owl1                 |                   |                                |                   |                                | -4.6              | 15.9                           |
| Sedge Wren <sup>1</sup>          | -11.5             | 0.9                            | 0.5               | 121.5                          | 1.8               | 200.5                          |
| Henslow's Sparrow                | -13.8             | 0.3                            | -12.6             | 0.5                            | -7.9              | 4.0                            |
| Grasshopper Sparrow <sup>1</sup> | -9.4              | 2.1                            | -5.2              | 12.5                           | -3.8              | 22.1                           |
| Bobolink <sup>1</sup>            | -0.5              | 82.2                           | -0.3              | 88.9                           | -1.8              | 49.2                           |
| Loggerhead Shrike <sup>1</sup>   |                   | -                              | -11.4             | 0.9                            | -3.7              | 23.0                           |
| Horned Lark <sup>2</sup>         | -4.7              | 15.3                           | -2.1              | 43.7                           | -2.1              | 43.7                           |
| Vesper Sparrow <sup>2</sup>      | -7.9              | 4.0                            | -5.4              | 11.5                           | -1.0              | 67.6                           |
| Eastern Meadowlark <sup>2</sup>  | -4.9              | 14.1                           | -4.3              | 18.0                           | -2.9              | 31.7                           |
| Savannah Sparrow <sup>2</sup>    | -2.6              | 35.8                           | -2.3              | 40.4                           | -0.9              | 70.3                           |

<sup>&</sup>lt;sup>1</sup>Highest priority or <sup>2</sup>High priority for conservation

Note: Background colors correspond with "regional credibility measures" for the data as provided by the authors. <u>Blue</u> indicates no deficiencies, <u>Yellow</u> (yellow) indicates a deficiency, and <u>Red</u> indicates an important deficiency.

## Bold indicates significant trends (P<0.05).

Some of New York's grassland birds have experienced steeper declines than others, or have a smaller population size and/or distribution across the state or region, and are therefore included in the highest priority tier in Table 6.2: northern harrier (*Circus cyaneus*), upland sandpiper (*Bartramia longicauda*), short-eared owl (*Asio flammeus*), sedge wren (*Cistothorus platensis*), Henslow's sparrow (*Ammodramus henslowii*), grasshopper sparrow (*Ammodramus savannarum*), bobolink (*Dolichonyx oryzivorus*), and loggerhead shrike (*Lanius ludovicianus*). Species included in the high priority tier are those that have been given relatively lower priority, but whose populations are also declining and are in need of conservation. The high priority tier in

<sup>&</sup>lt;sup>316</sup> Morgan and Burger, 2008.

Table 6.2 includes: horned lark (*Eremophila alpestris*), vesper sparrow (*Pooecetes gramineus*), eastern meadowlark (*Sturnella magna*), and savannah sparrow (*Passerculus sandwichensis*).

While these birds rely on grasslands in New York as breeding habitat (in general), two of these species (northern harrier and short-eared owl) and several other raptor species also rely on grasslands for wintering habitat. For this reason, a third target group of birds are those species that rely on grassland habitats while they over-winter (or are year-round residents) in New York, and include: snowy owl (Bubo scandiacus), rough-legged hawk (Buteo lagopus), red-tailed hawk (Buteo jamaicensis), American kestrel (Falco sparverius), and northern shrike (Lanius excubitor).

The specific effects of drilling for natural gas on nesting grassland birds are not well studied. However, the level of development expected for multi-pad horizontal drilling and minimum patch sizes of habitat necessary for bird reproduction, unless mitigated, will result in substantial impacts from the fragmentation of existing grassland habitats. Minimum patch sizes would vary by species and by surrounding land uses, but studies have shown that a minimum patch size of between 30-100 acres is necessary to protect a wide assemblage of grassland-dependent species. <sup>317</sup>

#### 6.4.1.2 Impacts of Forest Fragmentation

Forest fragmentation issues were the subject of two assessments referenced below which are specific to the East and address multiple horizontal well drilling from common pads. These studies, therefore, are more directly applicable to New York than previously mentioned western studies of vertical drilling. The Multi-Resolution Land Characteristic Dataset ("MRLC") (2004) indicates the following ratios of habitat types in the area underlain by the Marcellus shale in New York: 57% forested; 28% grassland/agricultural lands; and 3% scrub/shrub. The other 12% is divided evenly between developed land and open water/wetlands. As forests are the most common cover type, it is reasonable to assume that development of the Marcellus Shale would have a substantial impact on forest habitats and species.

<sup>&</sup>lt;sup>317</sup> USFWS n.d., Sample and Mossman 1997, Mitchell et al, 2000.

Today, New York is 63% (18.95 million acres) forested<sup>318</sup> and is unlikely to substantially increase. Current forest parcelization and fragmentation trends will likely result in future losses of large, contiguous forested areas.<sup>319</sup> Therefore, protecting these remaining areas is very important for maintaining the diversity of wildlife in New York.

The forest complex provides key ecosystem services that provide substantial ecological, economic, and social benefits (water quality protection, clean air, flood protection, pollination, pest predation, wildlife habitat and diversity, recreational opportunities, etc.) that extend far beyond the boundaries of any individual forested area.

Large contiguous forest patches are especially valuable because they sustain wide-ranging forest species, and provide more habitat for forest interior species. They are also more resistant to the spread of invasive species, suffer less tree damage from wind and ice storms, and provide more ecosystem services – from carbon storage to water filtration – than small patches, <sup>320</sup>

Lands adjacent to well pads and infrastructure can also be affected, even if they are not directly cleared. This is most notable in forest settings where clearings fragment contiguous forest patches, create new edges, and change habitat conditions for sensitive wildlife and plant species that depend on interior forest conditions.

Forest ecologists call this the edge effect. While the effect is somewhat different for each species, research has shown measurable impacts often extend at least 330 feet (100 meters) into forest adjacent to an edge. 321 Interior forest species avoid edges for different reasons. Blackthroated blue warblers and other interior forest birds, for example, avoid areas near edges during nesting season because of the increased risk of predation. Tree frogs, flying squirrels and certain woodland flowers are sensitive to forest fragmentation because of changes in canopy cover, humidity and light levels. Some species, such as white-tailed deer and cowbirds, are attracted to forest edges – often resulting in increased competition, predation, parasitism, and herbivory. Invasive plant species, such as tree of heaven, stilt grass, and Japanese barberry, often thrive on

<sup>&</sup>lt;sup>318</sup> NYSDEC, Forest Resource Assessment and Strategy, 2010.

<sup>&</sup>lt;sup>319</sup> NYSDEC, Forest Resource Assessment and Strategy, 2010.

<sup>&</sup>lt;sup>320</sup> Johnson, 2010, p. 19.

<sup>&</sup>lt;sup>321</sup> Johnson, 2010, p. 11.

forest edges and can displace native forest species. As large forest patches become progressively cut into smaller patches, populations of forest interior species decline.

# Lessons Learned from Pennsylvania

Assessment 1, General Discussion: The Nature Conservancy (TNC) conducted an assessment in 2010<sup>322</sup> to develop credible energy development projections for horizontal hydraulic fracturing in Pennsylvania's Marcellus Shale by 2030, and how those projections might affect high priority conservation areas, including forests. The projections were informed scenarios, not predictions, for how much energy development might take place and where it was more and less probable. Project impacts, however, were based on measurements of actual spatial footprints for hundreds of well pads.

Potential Direct Impacts, Methodology and Assessment Findings: Projections of future Marcellus gas development impacts depended on robust spatial measurements for existing Marcellus well pads and infrastructure. This assessment compared aerial photos of Pennsylvania Department of Environmental Protection (PADEP) Marcellus well permit locations taken before and after development and precisely documented the spatial foot print of 242 Marcellus well pads (totaling 435 drilling permits) in Pennsylvania.

Well pads in Pennsylvania occupy 3.1 acres on average while the associated infrastructure (roads, water impoundments, pipelines) takes up an additional 5.7 acres, or a total of nearly 9 acres per well pad (Figure 6.5). 323

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<sup>&</sup>lt;sup>322</sup> Johnson, 2010.

<sup>&</sup>lt;sup>323</sup> This is larger than the 7.4 acres predicted by IOGA to be disturbed in New York (section 6.4b).

| Average Spatial Disturbance for Marcellus Shale<br>Well Pads in Forested Context (acres)  |      |     |  |
|---|------|-----|--|
| Forest cleared for Marcellus Shale well pad   | 3.1  | 8.8 |  |
| Forest cleared for associated infrastructure (roads, pipelines, water impoundments, etc.) | 5.7  | 0.0 |  |
| Indirect forest impact from new edges   | 21.2 |     |  |
| TOTAL DIRECT AND INDIRECT IMPACTS   | 30   |     |  |

Figure 6.5 - Average Spatial Disturbance for Marcellus Shale Well Pads in Forested Context<sup>324</sup> (New July 2011)

Another key variable for determining land-use and habitat impacts in this assessment was the number of wells on each pad; more wells per pad translated to less disturbance and infrastructure on the landscape. It is technically possible to put a dozen or more Marcellus wells on one pad. For the 242 well pads assessed in this study, the average in Pennsylvania has been 2 wells per pad to date (IOGA estimates the same for New York) as companies quickly moved on to drill other leases to test productivity and to secure as many potentially productive leases as possible (leases typically expire after 5 years if there is no drilling activity). TNC assumed that in many cases, the gas company would return to these pads later and drill additional wells. This assumption may not be valid in New York where there is a three-year limit on well development (ECL 23-0501).

The TNC assessment developed low, medium, and high scenarios for the amount of energy development that might take place in Pennsylvania. The projections included a conservative estimate of 250 horizontal drilling rigs, each of which could drill one well per month, resulting in

<sup>&</sup>lt;sup>324</sup> Taken from Johnson, 2010, p. 10.

an estimated 3,000 wells drilled annually. Estimates in New York predict less activity than this, but activity could result in approximately 40,000 wells by 2040.

The low scenario (6,000 well pads) assumed that each pad on average would have 10 wells, or 1 well pad per 620 acres. Because many leases are irregularly shaped, in mixed ownership, or their topography and geology impose constraints, TNC concluded that it is unlikely this scenario would develop in Pennsylvania. It would take relatively consolidated leaseholds and few logistical constraints for this scenario to occur. 325

The medium scenario for well pads assumed 6 wells on average would be drilled from each pad (10,000 well pads), or 1 pad per 386 acres. Industry generally agreed that 6 is the most likely number of wells they would be developing per pad for most of their leaseholds in Pennsylvania. 326

The high scenario assumed each pad would have 4 wells drilled on average (15,000 well pads), or 1 pad per 258 acres. This scenario is more likely if there is relatively little consolidation of lease holds between companies in the next several years. While this scenario would result in a loss of less than 1% of Pennsylvania's total forest acreage, areas with intensive Marcellus gas development could see a loss of 2-3% of local forest habitats.

In summary, 60,000 wells could be drilled by 2030 in the area underlain by the Marcellus Shale in Pennsylvania on between 6,000 and 15,000 new well pads (there are currently about 1,000), depending on how many wells are placed on each pad.

A majority (64%) of projected well locations were found in a forest setting for all three scenarios. By 2030, a range of between 34,000 and 82,000 acres of forest cover could be cleared by new Marcellus gas development in Pennsylvania. Some part of the cleared forest area would become reforested after drilling is completed, but there has not been enough time to establish a trend since the Marcellus development started.

Note that while no definitive number is provided in section 5.1.3.2, this is expected to be the most common spacing for horizontal drilling in New York's Marcellus Shale.

Note that IOGA assumes that 6 horizontal wells would be drilled per pad in New York.

**Potential Direct Impacts, Implications for New York:** Direct land disturbance from horizontal hydraulic fracturing of Marcellus Shale in New York is expected to result in 7.4 acres of direct impacts from each well pad and associated infrastructure. This is different from the experiences in Pennsylvania where nearly 9 acres of habitat was removed for each well pad and its associated infrastructure. Under either scenario, the direct impacts are substantial.

The most likely drilling scenario in Pennsylvania would result in a density of 1 pad per 386 acres. However, given New York's regulatory structure, a spacing of 1 pad per 640 acres is anticipated. If spacing units are less than 640 acres, or if there are less than 6-8 horizontal wells per pad, the percentage of land disturbance could be greater. Again, using the set of currently pending applications as an example, the 47 proposed horizontal wells would be drilled on eleven separate well pads, with between 2 and 6 wells for each pad. Therefore, greater than 1.2% land disturbance per pad estimated by industry can be expected in New York.

Potential Indirect Impacts, Methodology and Assessment Findings: To assess the potential interior forest habitat impact, a 100-meter buffer was created into forest patches from new edges created by well pad and associated infrastructure development (Figure 6.6). For those well sites developed in forest areas or along forest edges (about half of the assessed sites), TNC calculated an average of 21 acres of interior forest habitat was lost. Thus, the total combined loss of habitat was 30 acres per well pad due to direct and indirect impacts. Figure 6.5 summarizes these data.

In addition to the direct clearing of between 34,000 to 82,000 acres of forest cover in Pennsylvania, forest interior species could be negatively impacted within an additional 85,000 to 190,000 forest acres adjacent to Marcellus development. Forest impacts would be concentrated where many of Pennsylvania's largest and most intact forest patches occur, resulting in fragmentation into smaller patches by well pads, roads, and other infrastructure. In contrast to overall forest loss, projected Marcellus gas development scenarios in Pennsylvania indicate a more pronounced impact on large forest patches. Impacts to forest interior species would vary depending on their geographic distribution and density. Some species, such as the black-throated blue warbler, could see widespread impacts to their relatively restricted breeding habitats in the state, while widely distributed species such as the scarlet tanager, would be relatively less affected.

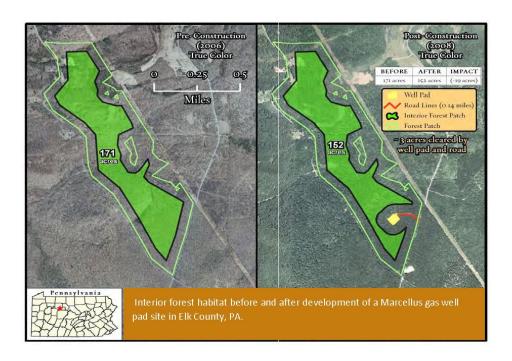


Figure 6.6 – Interior Forest Habitat Before & After Development of a Marcellus Gas Well Pad, Elk County PA<sup>327</sup> (New July 2011)

This study went on to find that locating energy infrastructure in open areas or toward the outer edges of large patches can significantly reduce impacts to important forest areas. To address this finding and explore potential ways in which conservation impacts could be minimized, TNC examined how projected Marcellus gas pads could be relocated to avoid forest patches in a specific region of Pennsylvania. To reduce the impacts to forest habitats, the wells were hypothetically relocated, where practicable, to nearby existing openings maintained by human activity (e.g., old fields, agricultural fields). If nearby open areas did not exist, the locations of the well pads were moved toward the edges of forest patches to minimize impacts to forest interior habitats. This exercise did not eliminate forest impacts in this heavily forested Pennsylvania landscape, but there was a significant reduction in impacts. Total forest loss declined almost 40% while impacts to interior forest habitats adjacent to new clearings declined by one-third (Figure 6.7). The study authors recommend that information about Pennsylvania's important natural habitats be an important part of the calculus about trade-offs and optimization as energy development proceeds.

<sup>&</sup>lt;sup>327</sup> Taken from Johnson, 2010, p. 11.

**Total Forested Acres Converted** 4500 4,138 4000 3500 2.830 3000 2500 ■ Modeled Well/Road 2000 Locations 1500 ■ Relocated 1000 Well/Road 400 Locations 500 244 0 Direct Impacts Indirect Impacts

Figure 6.7 - Total Forest Areas Converted 328 (New July 2011)

**Potential Indirect Impacts, Implications for New York:** For each acre of forest directly cleared for well pads and infrastructure in New York, an additional 2.5 acres can be expected to be indirectly impacted. Interior forest bird species with restricted breeding habitats, such as the black-throated blue and cerulean warblers, might be highly impacted.

Additional assessment work conducted for New York based on estimates and locations of well pad densities across the Marcellus landscape could better quantify expected impacts to forest interior habitats and wildlife.

New York Forest Matrix and Landscape Connectivity

Forest matrix blocks contain mature forests with old trees, understories, and soils that guarantee increased structural diversity and habitat important to many species. They include important stabilizing features such as large, decaying trunks on the forest floor and big, standing snags. Set within these matrix forests are smaller ecosystems offering a wide range of habitat (wetlands,

<sup>328</sup> Taken from Johnson, 2010, p. 27

streams, and riparian areas) that depend on the surrounding forested landscape for their long-term persistence and health. These large, contiguous areas are viable examples of the dominant forest types that, if protected, and in some cases allowed to regain their natural condition, serve as critical source areas for all species requiring interior forest conditions. Few remnants of such matrix blocks remain in the Northeast; it is therefore critical to conserve these priority areas to ensure long-term conservation of biodiversity.<sup>329</sup>

Assessment 2, General Discussion: The New York Natural Heritage program in 2010<sup>330</sup> identified New York's forest matrix blocks and predicted corresponding forest connectivity areas. Securing connections between major forested landscapes and their imbedded matrix forest blocks is important for the maintenance of viable populations of species, especially those that are wide-ranging and highly mobile, and ecological processes such as dispersal and pollination over the long term. Identifying, maintaining, and enhancing these connections represents a critical adaptation strategy if species are to shift their ranges in response to climate change and other landscape changes.

Assessment 2, Findings. Figure 6.8 depicts the large forested landscapes within New York and predicts the linkages between them, called least-cost path (LCP). A least-cost path corridor represents the most favorable dispersal path for forest species based on a combination of percent natural forest cover in a defined area, barriers to movement, and distance traveled. Thus, as many species that live in forests generally prefer to travel through a landscape with less human development (i.e., fewer impediments to transit) as well as in a relatively direct line, the predicted routes depict a balance of these sometimes opposing needs.

Assessment 2, Implications for New York: The area underlain by the Marcellus Shale in New York is 57% forested with about 7% of that forest cover occurring on State-owned lands. It is reasonable to assume high-volume horizontal hydraulic fracturing would have negative impacts to forest habitats similar to those predicted in Pennsylvania (Section 6.4.1.2).

In order to minimize habitat fragmentation and resulting restrictions to species movement in the area underlain by the Marcellus, it is recommended that forest matrix blocks be managed to

<sup>329</sup> TNC 2004.

<sup>&</sup>lt;sup>330</sup> NYSDEC, Strategic Plan for State Forest Management, 2010.

create, maintain, and enhance the forest cover characteristics that are most beneficial to the priority species that may use them.

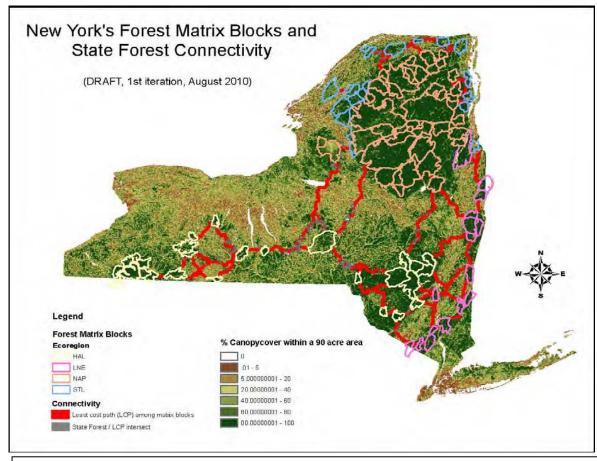


Figure 6.8 - New York's Forest Matrix Blocks and State Connectivity<sup>331</sup> (New July 2011)

HAL = High Allegheny Plateau; LNE = Lower New England/Northern Piedmont; NAP= Northern Appalachian/ Acadian; STL= St. Lawrence-Champlain Valley

## 6.4.2 Invasive Species

An invasive species, as defined by ECL §9-1703, is a species that is nonnative to the ecosystem under consideration and whose introduction causes or is likely to cause economic or environmental harm or harm to human health. Invasive species can be plants, animals, and other organisms such as microbes, and can impact both terrestrial and aquatic ecosystems.

<sup>&</sup>lt;sup>331</sup> Taken from NYSDEC, Strategic Plan for State Forest Management, 2010.

While natural means such as water currents, weather patterns and migratory animals can transport invasive species, human actions - both intentional and accidental - are the primary means of invasive species introductions to new ecosystems. Once introduced, invasive species usually spread profusely because they often have no native predators or diseases to limit their reproduction and control their population size. As a result, invasive species out-compete native species that have these controls in place, thus diminishing biological diversity, altering natural community structure and, in some cases, changing ecosystem processes. These environmental impacts can further impose economic impacts as well, particularly in the water supply, agricultural and recreational sectors. 332

The number of vehicle trips associated with high-volume hydraulic fracturing, particularly at multi-well sites, has been identified as an activity which presents the opportunity to transfer invasive terrestrial species. Surface water withdrawals also have the potential to transfer invasive aquatic species.

#### 6.4.2.1 Terrestrial

Terrestrial plant species which are widely recognized as invasive<sup>333</sup> or potentially-invasive in New York State, and are therefore of concern, are listed in Table 6.3 below.

Table 6.3 - Terrestrial Invasive Plant Species In New York State (Interim List) 334,335

| Terrestrial – Herbaceous       |                                  |  |  |
|--------------------------------|----------------------------------|--|--|
| Common Name                    | Scientific Name                  |  |  |
| Garlic Mustard                 | Alliaria petiolata               |  |  |
| Mugwort                        | Artemisia vulgaris               |  |  |
| Brown Knapweed                 | Centaurea jacea                  |  |  |
| Black Knapweed                 | Centaurea nigra                  |  |  |
| Spotted Knapweed               | Centaurea stoebe ssp. micranthos |  |  |
| Canada Thistle Cirsium arvense |                                  |  |  |
| Bull Thistle Cirsium vulgare   |                                  |  |  |

<sup>&</sup>lt;sup>332</sup> ECL §9-1701.

<sup>&</sup>lt;sup>333</sup> As per ECL §9-1703.

<sup>&</sup>lt;sup>334</sup> NYSDEC, DFWMR March 13, 2009. Interim List of Invasive Plant Species in New York State

This list was prepared pursuant to ECL §9-1705(5)(b) and ECL §9-1709(2)(d), but is not the so-called "four-Tier lists" referenced in ECL §9-1705(5)(h). As such the interim list is expected to be supplanted by the "four-Tier list" at such time that it becomes available.

| Terrestrial – Herbaceous                 |                             |  |  |  |
|--|-----------------------------|--|--|--|
| Common Name                              | Scientific Name             |  |  |  |
| Crown vetch                              | Coronilla varia             |  |  |  |
| Black swallow-wort                       | Cynanchum louiseae (nigrum) |  |  |  |
| European Swallow-wort                    | Cynanchum rossicum          |  |  |  |
| Fuller's Teasel                          | Dipsacus fullonum           |  |  |  |
| Cutleaf Teasel                           | Dipsacus laciniatus         |  |  |  |
| Giant Hogweed                            | Heracleum mantegazzianum    |  |  |  |
| Japanese Stilt Grass                     | Microstegium vimineum       |  |  |  |
| Terrestrial - Vines                      |                             |  |  |  |
| <b>Common Name</b>                       | Scientific Name             |  |  |  |
| Porcelain Berry                          | Ampelopsis brevipedunculata |  |  |  |
| Oriental Bittersweet                     | Celastrus orbiculatus       |  |  |  |
| Japanese Honeysuckle                     | Lonicera japonica           |  |  |  |
| Mile-a-minute Weed Persicaria perfoliata |                             |  |  |  |
|  |                             |  |  |  |

# **Terrestrial – Shrubs & Trees**

Pueraria montana var. lobata

Kudzu

| Common Name        | Scientific Name                    |  |
|--------------------|------------------------------------|--|
| Norway Maple       | Acer platanoides                   |  |
| Tree of Heaven     | Ailanthus altissima                |  |
| Japanese Barberry  | Berberis thunbergii                |  |
| Russian Olive      | Elaeagnus angustifolia             |  |
| Autumn Olive       | Elaeagnus umbellata                |  |
| Glossy Buckthorn   | Frangula alnus                     |  |
| Border Privet      | Ligustrum obtusifolium             |  |
| Amur Honeysuckle   | Lonicera maackii                   |  |
| Shrub Honeysuckles | Lonicera morrowii/tatarica/x bella |  |
| Bradford Pear      | Pyrus calleryana                   |  |
| Common Buckthorn   | Rhamnus cathartica                 |  |
| Black Locust       | Robinia pseudoacacia               |  |
| Multiflora Rose    | Rosa multiflora                    |  |

Operations involving land disturbance such as the construction of well pads, access roads, and engineered surface impoundments for fresh water storage have the potential to both introduce and transfer invasive species populations. Machinery and equipment used to remove vegetation and soil may come in contact with invasive plant species that exist at the site and may inadvertently transfer those species' seeds, roots, or other viable plant parts via tires, treads/tracks, buckets, etc. to another location on site, to a separate project site, or to any location in between.

The top soil that is stripped from the surface of the site during construction and set aside for reuse during reclamation also presents an opportunity for the establishment of an invasive species population if it is left exposed. Additionally, fill sources (e.g., gravel, crushed stone) brought to the well site for construction purposes also have the potential to act as a pathway for invasive species transfer if the fill source itself contains viable plant parts, seeds, or roots.

#### *6.4.2.2 Aquatic*

The presence of non-indigenous aquatic invasive species in New York State waters is recognized, and, therefore, operations associated with the withdrawal, transport, and use of water for horizontal well drilling and high volume hydraulic fracturing operations have the potential to transfer invasive species. Species of concern include, but are not necessarily limited to; zebra mussels, eurasian watermilfoil, alewife, water chestnut, fanwort, curly-leaf pondweed, round goby, white perch, didymo, and the spiny water flea. Other aquatic, wetland and littoral plant species that are of concern due to their status as invasive <sup>336</sup> or potentially-invasive in New York State are listed in Table 6.4.

<sup>&</sup>lt;sup>336</sup> As per ECL §9-1703.

Table 6.4 - Aquatic, Wetland & Littoral Invasive Plant Species in New York State (Interim List)<sup>337,338</sup>

| Floating & Su                  | bmerged Aquatic                     |  |
|--------------------------------|-------------------------------------|--|
| Common Name                    | Scientific Name                     |  |
| Carolina Fanwort               | Cabomba caroliniana                 |  |
| Rock Snot (didymo)             | Didymosphenia geminata              |  |
| Brazilian Elodea               | Egeria densa                        |  |
| Water thyme                    | Hydrilla verticillata               |  |
| European Frog's Bit            | Hydrocharis morus-ranae             |  |
| Floating Water Primrose        | Ludwigia peploides                  |  |
| Parrot-feather                 | Myriophyllum aquaticum              |  |
| Variable Watermilfoil          | Myriophyllum heterophyllum          |  |
| Eurasian Watermilfoil          | Myriophyllum spicatum               |  |
| Brittle Naiad                  | Najas minor                         |  |
| Starry Stonewort (green alga)  | Nitellopsis obtusa                  |  |
| Yellow Floating Heart          | Nymphoides peltata                  |  |
| Water-lettuce                  | Pistia stratiotes                   |  |
| Curly-leaf Pondweed            | Potamogeton crispus                 |  |
| Water Chestnut                 | Trapa natans                        |  |
|                                | etland & Littoral                   |  |
| Common Name                    | Scientific Name                     |  |
| Flowering Rush                 | Butomus umbellatus                  |  |
| Japanese Knotweed              | Fallopia japonica                   |  |
| Giant Knotweed                 | Fallopia sachalinensis              |  |
| Yellow Iris                    | Iris pseudacorus                    |  |
| Purple Loosestrife             | Lythrum salicaria                   |  |
| Reed Canarygrass               | Phalaris arundinacea                |  |
| Common Reed- nonnative variety | Phragmites australis var. australis |  |

<sup>337</sup> NYSDEC, DRWMR March 13, 2009 Interim List of Invasive Plant Species in New York State

This list was prepared pursuant to ECL §9-1705(5)(b) and ECL §9-1709(2)(d) ), but is not the so-called "four-Tier lists" referenced in ECL §9-1705(5)(h). As such the interim list is expected to be supplanted by the "four-Tier list" at such time that it becomes available.

Invasive species may be transported with the fresh water withdrawn for, but not used for drilling or hydraulic fracturing. Invasive species may potentially be transferred to a new area or watershed if unused water containing such species is later discharged at another location. Other potential mechanisms for the possible transfer of invasive aquatic species may include trucks, hoses, pipelines and other equipment used for water withdrawal and transport.

## 6.4.3 Impacts to Endangered and Threatened Species

The area underlain by the Marcellus Shale includes both terrestrial and aquatic habitat for 18 animal species listed as endangered or threatened in New York State (Table 6.5 and Figure 6.9) protected under the State Endangered Species Law (ECL 11-0535) and associated regulations (6 NYCRR Part 182). Some species, such as the northern harrier and upland sandpiper, are dependent upon grassland habitat for breeding and foraging and can be found in many counties within the project area. Species such as the rayed bean mussel and mooneye fish are aquatic species limited to only two counties on the western edge of the project area. Other species are associated with woodlands, with bald eagles nesting in woodlands adjacent to lakes, rivers and ponds throughout many counties within the project area. The area also includes habitat for cerulean warblers and eastern hellbenders, two species currently under consideration for listing by both the State and the federal government.

Endangered and threatened wildlife may be adversely impacted through project actions such as clearing, grading and road building that occur within the habitats that they occupy. Certain species are unable to avoid direct impact due to their inherent poor mobility (e.g., Blanding's turtle, club shell mussel). Certain actions, such as clearing of vegetation or alteration of stream beds, can also result in the loss of nesting and spawning areas. If these actions occur during the time of year that species are breeding, there can be a direct loss of eggs and/or young. For species that are limited to specific habitat types for breeding, the loss of the breeding area can result in a loss of productivity in future years as adults are forced into less suitable habitat. Any road construction through streams or wetlands within habitats occupied by these species can result in the creation of impermeable barriers to movement for aquatic species and reduce dispersal for some terrestrial species. Other impacts from the project, such as increased vehicle traffic, can result in direct mortality of adult animals. In general, the loss of habitat in areas

occupied by listed species can result in reduced numbers of breeding pairs and lowered productivity.

Table 6.5 - Endangered & Threatened Animal Species within the Area Underlain by the Marcellus Shale <sup>339</sup>(New July 2011)

| Common Name             | Scientific name             | NYS Listing | Primary Habitats         |
|-------------------------|-----------------------------|-------------|--------------------------|
| Henslow's Sparrow       | Ammodramus<br>henslowii     | Threatened  | Grassland                |
| Short-eared Owl         | Asio flammeus               | Endangered  | Grassland                |
| Upland Sandpiper        | Bartramia longicauda        | Threatened  | Grassland                |
| Northern Harrier        | Circus cyaneus              | Threatened  | Grassland, wetlands      |
| Sedge Wren              | Cistothorus platensis       | Threatened  | Grassland                |
| Peregrine Falcon        | Falco peregrinus            | Endangered  | Cliff faces              |
| Bald Eagle              | Haliaeetus<br>leucocephalus | Threatened  | Forest, open water       |
| Least Bittern           | Ixobrychus exilis           | Threatened  | Wetlands                 |
| Pie-billed Grebe        | Podilymbus podiceps         | Threatened  | Wetlands                 |
| Eastern Sand Darter     | Ammocrypta<br>pellucida     | Threatened  | Streams                  |
| Mooneye                 | Hiodon tergisus             | Endangered  | Large Lakes, Rivers      |
| Longhead Darter         | Percina<br>macrocephala     | Threatened  | Large Streams, Rivers    |
| Brook Floater           | Alasmidonta varicosa        | Threatened  | Streams and Rivers       |
| Wavyrayed<br>Lampmussel | Lampsilis fasciola          | Threatened  | Small, Medium<br>Streams |
| Green Floater           | Lasmigona subviridis        | Threatened  | Small, Medium<br>Streams |
| Clubshell               | Pleurobema clava            | Endangered  | Small, Medium<br>Streams |
| Rayed Bean              | Villosa fabalis             | Endangered  | Small Streams            |
| Timber rattlesnake      | Crotalus horridus           | Threatened  | Forest                   |

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<sup>&</sup>lt;sup>339</sup> November 3, 2010

Areas of Concern for Endangered and Threatened Animals within New York State

Lake Ontario

Lake Erie

PENNSYLVANIA

LEGEND

Threatened and Endangered Animals
Map prepared by NYS HIU, 6/8/2011; 3/13/2012; ktb

County Boundary for New York State

Figure 6.9 - Areas of Concern for Endangered and Threatened Animal Species, March 31, 2011 (New July 2011)

## 6.4.4 Impacts to State-Owned Lands

State-owned lands play a unique role in New York's landscape because they are managed under public ownership to allow for sustainable use of natural resources, provide recreational opportunities for all New Yorkers, and provide important wildlife habitat and open space. They represent the most significant portions of large contiguous forest patch in the study area. Industrial development on these lands is, for the most part, prohibited, and any type of clearing and development on these lands is limited and managed. Given the level of development expected for multi-pad horizontal drilling, it is anticipated that there would be additional pressure for surface disturbance on state-owned lands. Surface disturbance associated with gas extraction could have a significant adverse impact on habitats contained on the state-owned lands, and recreational use of those lands.

#### Forest Habitat Fragmentation

As described earlier, large contiguous forest patches are especially valuable because they sustain wide-ranging forest species, and provide more habitat for forest interior species. State-owned lands, by their very nature, consist of large contiguous forest patches. While some fragmentation has occurred, the level of activity associated with multi-well horizontal drilling (e.g., well pad construction, access roads, pipelines, etc.) would negatively impact the state's ability to maintain the existing large contiguous patches of forest.

The Department has stated that protecting these areas from further fragmentation is a high priority. One of the objectives stated in the Strategic Plan for State Forest Management is to "emphasize closed canopy and interior forest conditions to maintain and enhance" forest matrix blocks. It is critical therefore, that any additional road, pipeline and well pad construction be carefully assessed in order to avoid further reducing this habitat (see also Section 6.4.1). Given the State's responsibility to protect these lands as steward of the public trust, the State has a heightened responsibility, as compared to its role with respect to private lands, to ensure that any State permitted action does not adversely impact the ecosystems and habitat on these public lands so that they may be enjoyed by future generations.

## Public Recreation

State-owned lands have been acquired over the past century to provide compatible public recreation opportunities, protect watersheds, and provide sustainable timber harvesting. Drilling and trucking activities disturb the tranquility found on these lands and can cause significant visual impacts. Also, many State Forest roads serve as recreational trails for bicyclists, horseback riders, snowmobilers and others. The level of truck traffic associated with horizontal drilling and high-volume hydraulic fracturing presents safety issues, and would significantly degrade the experience for users of these roads, if not altogether during the drilling and construction phases of development.

# Legal Considerations

State Forests have an identity that is distinct from private lands, prescribed by the NYS Constitution, the ECL and the Environmental Quality Bond Acts of 1972 and 1986, under the

provisions of which they were acquired. New York State Constitution Article XIV, Section 3(1) states:

"Forest and wild life conservation are hereby declared to be policies of the state. For the purposes of carrying out such policies the legislature may appropriate moneys for the acquisition by the state of land, outside of the Adirondack and Catskill parks as now fixed by law, for the practice of forest or wild life conservation."

ECL Section 9-0501(1), in keeping with the above constitutional provision, authorizes the state to acquire reforestation areas, "which are adapted for reforestation and the establishment and maintenance thereon of forests for watershed protection, the production of timber and other forests products, and for recreation and kindred purposes,... which shall be forever devoted to the planting, growth and harvesting of such trees..."

Similarly, ECL Section 11-2103(1) authorizes the state to acquire "lands, waters or lands and waters...for the purpose of establishing and maintaining public hunting, trapping and fishing grounds."

ECL Section 9-0507 provides the Department discretionary authority to lease oil and gas rights on reforestation areas, provided that "such leasehold rights shall not interfere with the operation of such reforestation areas for the purposes for which they were acquired and as defined in Section 3 of Article XIV of the Constitution." The expected volume of truck traffic, the expected acreage that would be converted to non-forest use in the form of well pads, roads and pipelines, and noise and other impacts, raise serious questions as to how the surface activities anticipated with horizontal drilling and high-volume hydraulic fracturing could be viewed as consistent with this provision of the ECL.

For Wildlife Management Areas (WMAs) there are additional legal considerations stemming from the use of federal funds. Many WMAs were purchased using Federal Aid in Wildlife Restoration (Pittman-Robertson) funds and all are managed/maintained using Pittman-Robertson funds. Under these provisions, any surface use of the land must not be in conflict with the intended use as a WMA. These areas are managed for natural habitats to benefit wildlife, and disturbance associated with multi-pad wells raises questions about compatibility with essential

wildlife behaviors such as breeding, raising young, and preparation for migration. Also, selling or leasing of minerals rights must be approved by the U.S. Fish and Wildlife Service, and may require reimbursement of the federal government for revenue generated. In addition, siting well pads on WMAs purchased with Conservation Fund monies may require additional mitigation under federal statutes and/or compensation.

# 6.5 Air Quality

## 6.5.1 Regulatory Overview

This section provides a comprehensive list of federal and New York State regulations which could potentially be applicable to air emissions and air quality impacts associated with the drilling, completion (hydraulic fracturing and flowback) and production phases (processing, transmission and storage). At each of these phases, there are a number of air emission sources that may be subject to regulation. These general regulatory requirements are then followed by specific information regarding emission sources that have potential regulatory implications, as presented below in Sections 6.5.1.1 to 6.5.1.8. Certain discussions reflect new industry information provided in response to Department requests, as well as finalization, clarification, and revision to EPA regulations and policy. For example, the definition of what constitutes a stationary source or "facility" has been refined for criteria pollutants. These discussions are then followed with Department rule-applicability determinations on in instances where such decisions can be made as part of the SGEIS, as well as how the Department envisions the permitting of specific operations should proceed (Section 6.5.1.8).

## Applicable Federal Regulations

Prevention of Significant Deterioration of Air Quality (PSD): Under the PSD program, a federally-enforceable permit is required in order to restrict emissions from new major or major modification to existing sources (e.g., power plants and manufacturing facilities which emit criteria air pollutants in quantities above 100 tons per year) located in areas classified as attainment or unclassifiable with respect to the <a href="http://www.epa.gov/ttn/naaqs/">http://www.epa.gov/ttn/naaqs/</a>. That is, PSD requirements apply to all pollutants that do not exceed the NAAQS in the source location area. The NAAQS are numerical maximum pollution levels set to protect public health and welfare which have been established for ozone (O<sub>3</sub>), nitrogen dioxide (NO<sub>2</sub>), sulfur dioxide (SO<sub>2</sub>), fine particulate matter (PM10 and PM2.5), carbon monoxide (CO) and lead. The federal PSD

program is contained in 40 CFR Section 52.21 and the federally approved State program is found at 6 NYCRR Part 231.

Nonattainment New Source Review (NNSR): This federal program applies to new major or modified existing major sources in areas where the NAAQS are exceeded. The requirements for source emissions and potential impacts are more restrictive than through the PSD program. The federal program is found at 40 CFR Section 51.165 and the federally approved State program is found at 6 NYCRR Part 231. In New York State, nonattainment requirements are currently applicable to major sources of O<sub>3</sub> precursors (NO<sub>x</sub> and VOC) and direct PM2.5 and its precursor emissions (SO<sub>2</sub> and NOx). EPA has approved 6 NYCRR Part 231 into the State Implementation Plan. The regulation is described further under "Applicable State Regulations" below.

New Source Performance Standards (NSPS): Section 111 of the Clean Air Act (CAA) requires EPA to adopt emissions standards that are applicable to new, modified, and reconstructed sources. The requirements are meant to force new facilities to perform as well as or better than the best existing facilities (commonly known as "best demonstrated technology"). As new technology advances are made, EPA is required to revise and update NSPS applicable to designated sources. The following federal NSPS may apply:

- 40 CFR Part 60, Subpart JJJJ, Standards of Performance for Stationary Spark Ignition (SI) Internal Combustion Engines (ICE). Subpart JJJJ applies to manufacturers, owners and operators of SI ICE which affects new, modified, and reconstructed stationary SI ICE (i.e., generators, pumps and compressors), combusting any fuel (i.e., gasoline, natural gas, LPG, landfill gas, digester gas etc.), except combustion turbines. The applicable emissions standards are based on engine type, fuel type, and manufacturing date. The regulated pollutants are NO<sub>x</sub>, CO and VOC and there is a sulfur limit on gasoline. Subpart JJJJ would apply to facilities operating spark ignition engines at compressor stations;
- 40 CFR Part 60, Subpart IIII Standards of Performance for Stationary Compression Ignition (CI) ICEs. Subpart IIII applies to manufacturers, owners and operators of CI ICE (diesel) which affects new, modified, and reconstructed (commencing after July 11, 2005) stationary CI ICE (i.e., generators, pumps and compressors), except combustion turbines. The applicable emissions standards (phased in Tiers with increasing levels of stringency) are based on engine type and model year. The regulated pollutants are NO<sub>x</sub>, PM, CO, non-methane hydrocarbons (NMHC), while the emissions of sulfur oxides

- (SO<sub>x</sub>) are reduced through the use of low sulfur fuel. Particulate emissions are also reduced by standards. Subpart IIII would apply to facilities operating compression ignition engines at compressor stations;
- 40 CFR Part 60, Subpart KKK Standards of Performance for Equipment Leaks of VOC from Onshore Natural Gas Processing Plants. Subpart KKK applies to gas processing plants that are engaged in the extraction of natural gas liquids from field gas and contains provisions for VOC leak detection and repair (LDAR);
- 40 CFR Part 60, Subpart LLL Standards of Performance for Onshore Natural Gas Processing: SO<sub>2</sub> Emissions. Subpart LLL governs emissions of SO<sub>2</sub> from gas processing plants, specifically gas sweetening units (remove H<sub>2</sub>S and CO<sub>2</sub> from sour gas) and sulfur recovery units (recover elemental sulfur); and
- 40 CFR Part 60 Subpart Kb Standards of Performance for Volatile Organic Liquid Storage Vessels (Including Petroleum Liquid Storage Vessels) for which Construction, Reconstruction, or Modification Commenced after July 23, 1984.

National Emission Standards for Hazardous Air Pollutants (NESHAPs): Section 112 of the CAA requires EPA to adopt standards to control emissions of hazardous air pollutants (HAPs). NESHAPs are applicable to both new and existing sources of HAPs, and there are NESHAPs for both "major" sources of HAPs and "area" sources of HAPs. A major source of HAPs is one with the potential to emit in excess of 10 Tpy of any single HAP or 25 Tpy of all HAPs, combined. An area source of HAPs is a stationary source of HAPs that is not major. The aim is to develop technology-based standards which require levels met by the best existing facilities. The pollutants of concern in the oil and gas sector primarily are the following: BTEX, formaldehyde, and n-hexane. The following federal NESHAPs may apply:

- 40 CFR Part 63, Subpart ZZZZ National Emission Standards for Hazardous Air Pollutants for Reciprocating Internal Combustion Engines (RICE). Appendix 17 has been revised from the initial analysis to reflect the requirements in the final EPA rule;
- 40 CFR Part 63, Subpart H National Emission Standards for Organic Hazardous Air Pollutants for Equipment Leaks. Subpart H applies to equipment that contacts fluids with a HAP concentration of 5%;
- 40 CFR Part 63, Subpart HH NESHAPs from Oil and Natural Gas Production Facilities. Subpart HH controls air toxics from oil and natural gas production operations and

contains provisions for both major sources and area sources of HAPs. Emission sources affected by this regulation are tanks with flash emissions (major sources only), equipment leaks (major sources only), and glycol dehydrators (major and area sources). Further details on this subpart are presented in section 6.5.1.2;

- 40 CFR Part 63, Subpart HHH NESHAPs from Natural Gas Transmission and Storage Facilities. Subpart HHH controls air toxics from natural gas transmission and storage operations. It affects glycol dehydrators located at major sources of HAPs; and
- 40 CFR Part 61, Subpart V National Emission Standard for Equipment Leaks (Fugitive Emission Sources). Subpart V applies to equipment that contacts fluids with a volatile HAP concentration of 10%.

# Applicable New York State Regulations

New York State Air Regulations are codified at 6 NYCRR Part 200 *et seq*, and can be obtained from the Department's web site at <a href="http://www.dec.ny.gov/regs/2492.html">http://www.dec.ny.gov/regs/2492.html</a>. Some of the applicable regulations are briefly described below.

- Part 200 General Provisions;
  - Section 200.1 Definitions (relevant subsections);
    - (cd) *Stationary source*. Any building, structure, facility or installation, excluding nonroad engines, that emits or may emit any air pollutant;
    - (aw) *Nonroad engine*. (1) Except as specified in paragraph (2) of this subdivision, a nonroad engine is an internal combustion engine:
    - (iii) that, by itself or in or on a piece of equipment, is portable or transportable, meaning designed to be and capable of being carried or moved from one location to another. Indicators of transportability include, but are not limited to, wheels, skids, carrying handles, dolly, trailer, or platform.
    - (2) An internal combustion engine is not a nonroad engine if:
    - (iii) the engine otherwise included in subparagraph (1)(iii) of this subdivision remains or would remain at a location for more than 12 consecutive months or a shorter period of time for an engine located at a seasonal source. A *location* is any single site at a building, structure, facility, or installation. Any engine (or engines) that replaces an engine at a location and that is intended to perform the

same or similar function as the engine replaced would be included in calculating the consecutive time period. An engine located at a seasonal source is an engine that remains at a seasonal source during the full annual operating period of the seasonal source. A seasonal source is a stationary source that remains in a single location on a permanent basis (*i.e.* at least two years) and that operates at that single location approximately three months (or more) each year. This paragraph does not apply to an engine after the engine is removed from the location;

- Section 200.6 Acceptable Ambient Air Quality. Section 200.6 states, "notwithstanding the provisions of this Subchapter, no person shall allow or permit any air contamination source to emit air contaminants in quantities which alone or in combination with emissions from other air contamination sources would contravene any applicable ambient air quality standard and/or cause air pollution. In such cases where contravention occurs or may occur, the commissioner shall specify the degree and/or method of emission control required". This regulation prohibiting air pollution, allowing the Department to evaluate ambient impacts from emission sources; and
- Section 200.7 Maintenance of Equipment. Section 200.7 states, "any person who owns or operates an air contamination source which is equipped with an emission control device shall operate such device and keep it in a satisfactory state of maintenance and repair in accordance with ordinary and necessary practices, standards and procedures, inclusive of manufacturer's specifications, required to operate such device effectively.

#### • Part 201 - Permits and Registrations;

- o 201-2.1 Definitions.
  - (21) Major stationary source or major source or major facility (see further details and discussions below);
- o 201-5 State Facility Permits. Subpart 201-5 contains the criteria to issue "state facility permits" to facilities that are not considered to be major. These are generally facilities with the following characteristics: (1) Their actual emissions exceed 50% of the level that would make them major, but their potential to emit as defined in 6 NYCRR Part 200 does not place them in the major category, (2) They require the use of permit conditions to limit emissions below thresholds that would make them subject to certain state or federal requirements, or (3) They have been granted variances under the Department's air regulations;

- O 201-6 Title V Facility Permits. Subpart 201-6 contains the requirements and procedures for CAA "Title V Permits". These include facilities that are judged to be major under the Department's regulations, or that are subject to NSPSs, to a standard or other requirements regulating HAPs or to federal acid rain program requirements; and
- O 201-7 Federally Enforceable Emission Caps. Subpart 201-7 provides the ability to accept federally enforceable permit terms and conditions which restrict or cap emissions from a stationary source or emission unit in order to avoid being subject to one or more applicable requirements.
- Part 212 General Process Emission Sources. In general, Part 212 regulates emissions of particulate, opacity, VOCs (from major sources), NO<sub>x</sub> (from major sources) and is mainly used to control air toxics from industries not regulated in other specific 6 NYCRR Parts;
- Part 227- Stationary Combustion Installations (see Appendix 16 for more details):
  - o 227-1- <u>Stationary Combustion Installations</u>. Subpart 227-1 regulates emissions from stationary combustion installations.
  - o 227-2 Reasonably Available Control Technology (RACT) For Major Facilities of Oxides Of Nitrogen (NO<sub>x</sub>). Subpart 227-2 imposes NO<sub>x</sub> limits on major sources (with a potential to emit 100 tons of NO<sub>x</sub> per year) located in the attainment areas of the northeast ozone transport region;
- Part 229 <u>Petroleum and Volatile Organic Liquid Storage and Transfer</u>. Part 229
  regulates petroleum and volatile organic liquid storage and transfer (i.e., gasoline bulk
  plants, gasoline loading terminals, marine loading vessels, petroleum liquid storage tanks
  or volatile organic liquid storage tanks); and
- Part 231- New Source Review (NSR) for New and Modified Facilities. Part 231 addresses both the federal NSR and PSD requirements for sources located in nonattainment or attainment areas and the relevant program requirements. For new major facilities or modification of existing major facilities, Part 231 applies to those NSR pollutants with proposed emissions increases greater than the major facility or significant project threshold, as applicable. The applicable PSD major facility threshold (100 or 250 tons per year) is determined by whether the facility belongs to one of the source categories listed in 6 NYCRR §201-2.1(b)(21)(iii). Reciprocating internal combustion engines are not on the list, making the major source threshold 250 tons per year (instead of 100 tons/year) for PSD applicable pollutants. For the nonattainment pollutants, the threshold levels are lower, and depend on the location of the proposed new facility or

modification. For the Marcellus Shale area, which is located within the Ozone Transport Region (OTR), for regulatory purposes, the area is treated as moderate ozone nonattainment. The major facility thresholds are 50 tons per year for VOC and 100 tons per year for NOx.

The following sections discuss what regulatory determinations the Department has made with respect to operations associated with drilling and completion activities and how the regulatory process would be used for further permitting determinations related to the offsite compressor stations and its association with the well pad operations.

# 6.5.1.1 Emission Analysis $NO_x$ - Internal Combustion Engine Emissions Compressor Engine Exhausts

Internal combustion engines provide the power to run compressors that assist in the production of natural gas from wells and pressurize natural gas from wells to the pressure of lateral lines that move natural gas in large pipelines to and from processing plants and through the interstate pipeline network. The engines are often fired with raw or processed natural gas, and the combustion of the natural gas in these engines results in air emissions.

# Well Drilling and Hydraulic Fracturing Operations

Oil and gas drilling rigs require substantial power to drill and case wellbores to their target formations. For the development of the Marcellus Shale, this power would typically be provided by transportable diesel engines, which generate exhaust from the burning of diesel fuel. After the wellbore is drilled to the target formation, additional power is needed to operate the pumps that move large quantities of water, sand, or chemicals into the target formation at high pressure to hydraulically fracture the shale.

The preferred method for calculating engine emissions is to use emission factors provided by the engine manufacturer. If these cannot be obtained, a preliminary emissions estimate can be made using EPA AP-42 emission factors. The most commonly used tables appear as Table 6.6 below.

| EPA AP-42 Table 3.2-1: Emission Factors for Uncontrolled Natural Gas-Fired Engines |                          |                          |                          |                          |                          |                          |
|--|--------------------------|--------------------------|--------------------------|--------------------------|--------------------------|--------------------------|
|  | 2-cycle l                | ean burn                 | 4-cycle le               | ean burn                 | 4-cycle rich burn        |                          |
| Pollutant  | g/Hp-hr<br>(power input) | lb/MMBtu<br>(fuel input) | g/Hp-hr<br>(power input) | lb/MMBtu<br>(fuel input) | g/Hp-hr<br>(power input) | lb/MMBtu<br>(fuel input) |
| NO <sub>x</sub>  | 10.9                     | 2.7                      | 11.8                     | 3.2                      | 10.0                     | 2.3                      |
| CO   | 1.5                      | 0.38                     | 1.6                      | 0.42                     | 8.6                      | 1.6                      |
| TOC 1  | 5.9                      | 1.5                      | 5.0                      | 1.3                      | 1.2                      | 0.27                     |

TOC is total organic compounds (sometimes referred to as THC). To determine VOC emissions calculate TOC emissions and multiply the value by the VOC weight fraction of the fuel gas.

| EPA AP-42 Table 3.3-1: Emission Factors for Uncontrolled Gasoline and Diesel Industrial Engines |                           |                          |                           |                          |  |
|---|---------------------------|--------------------------|---------------------------|--------------------------|--|
|   | Gasoline                  | Fuel                     | Diesel Fuel               |                          |  |
| Pollutant   | g/Hp-hr<br>(power output) | lb/MMBtu<br>(fuel input) | g/Hp-hr<br>(power output) | lb/MMBtu<br>(fuel input) |  |
| NO <sub>x</sub>   | 5.0                       | 1.63                     | 14.1                      | 4.41                     |  |
| СО  | 3.16                      | 0.99                     | 3.03                      | 0.95                     |  |
| Exhaust (TOC)   | 6.8                       | 2.10                     | 1.12                      | 0.35                     |  |
| Evaporative (TOC)   | 0.30                      | 0.09                     | 0.00                      | 0.00                     |  |
| Crankcase (TOC)   | 2.2                       | 0.69                     | 0.02                      | 0.01                     |  |

# Engine Emissions Example Calculations

0.15

0.00

0.00

A characterization of the significant  $NO_x$  emission sources during the three operational phases of horizontally drilled, hydraulically fractured natural gas wells is as follows:

# 1. Horizontally Drilled/ High-Volume Hydraulically Fractured Wells - Drilling Phase

0.5

For a diesel engine drive total of 5400 Hp drilling rig power, <sup>340</sup> using NO<sub>x</sub> emission factor data from engine specification data received from natural gas production companies currently operating in the Marcellus Shale formation outside New York State, a representative NO<sub>x</sub> emission factor of 6.4 g/Hp-hr is used in this example. For purposes of estimating the Potential

Refueling (TOC)

<sup>&</sup>lt;sup>340</sup> Engine information provided by Chesapeake Energy

to Emit (PTE) for the engines, continuous year-round operation is assumed. The estimated NO<sub>x</sub> emission would be:

 $NO_{y}$  emissions =  $(6.4 \text{ g/Hp-hr}) \times (5400 \text{ Hp}) \times (8760 \text{ hr/yr}) \times (\text{ton/2000 lb}) \times (1 \text{ lb/453.6 g}) = 333.7 \text{ Tpy}$ 

The actual emissions from the engines would be much lower than the above PTE estimate, depending on the number of wells drilled and the time it takes to drill the wells at a well site in a given year.

2. Horizontally Drilled/ High-Volume Hydraulically Fractured Wells - Completion Phase

For diesel-drive 2333 Hp fracturing pump engine(s),  $^{341}$  using NO<sub>x</sub> emission factor data from engine specification data received from natural gas production companies currently operating in the Marcellus Shale formation outside New York State, a representative NO<sub>x</sub> emission factor of 6.4 g/Hp-hr is used in this example. For purposes of estimating the Potential to Emit (PTE) for the engines, continuous year-round operation is assumed. The estimated NO<sub>x</sub> emission would be:

$$NO_{\chi}$$
 emissions =  $(6.4 \text{ g/Hp-hr}) \times (2333 \text{ Hp}) \times (8760 \text{ hr/yr}) \times (\text{ton/2000 lb}) \times (1 \text{ lb/453.6 g}) = 144.1 \text{ Tpy}$ 

The actual emissions from the engines would be lower than the above PTE estimate, depending on the time it takes to hydraulically fracture each well and the number of wells hydraulically fractured at a well site in a given year.

3. Horizontally Drilled/High-Volume Hydraulically Fractured Wells - Production Phase

Using recent permit application information from a natural gas compressor station in the Department's Region 8, a NO<sub>x</sub> emission factor 2.0 g/Hp-hr was chosen as more reasonable (yet still conservative) than AP-42 emission data. The maximum site-rated horsepower is 2500 Hp. <sup>342</sup> The engine(s) is expected to run year round (8760 hr/yr).

$$NO_x$$
 emissions =  $(2.0 \text{ g/Hp-hr}) \times (2500 \text{ Hp}) \times (8760 \text{ hr/yr}) \times (\text{ton/}2000 \text{ lb}) \times (1 \text{ lb/}453.6 \text{ g}) = 48.3 \text{ TPY}$ 

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<sup>&</sup>lt;sup>341</sup> Engine information provided by Chesapeake Energy.

<sup>&</sup>lt;sup>342</sup> Engine information provided by Chesapeake Energy.

Since the engines in the example comply with the NO<sub>x</sub> RACT emission limits, non-applicability of the rule implies merely avoiding the monitoring requirements that were designed for permanently located engines. In addition to NO<sub>x</sub> RACT requirements, Title V permitting requirements could also apply to other air pollutants such as CO, SO<sub>2</sub>, particulate matter (PM), ozone (as VOCs), and elemental lead, with the same emission thresholds as for NO<sub>x</sub>. An initial review of other emission information for these engines, such as CO and PM emission factor data, reveals an unlikely possibility of reaching major source thresholds triggering Title V permitting requirements for these facilities as discussed further in Section 6.5.1.8.

# 6.5.1.2 Natural Gas Production Facilities NESHAP 40 CFR Part 63, Subpart HH (Glycol Dehydrators)

Natural gas produced from wells is a mixture of a large number of gases and vapors. Wellhead natural gas is often delivered to processing plants where higher molecular weight hydrocarbons, water, nitrogen, and other compounds are largely removed if they are present. Processing results in a gas stream that is enriched in methane at concentrations of usually more than 80%. Not all natural gas requires processing, and gas that is already low in higher hydrocarbons, water, and other compounds can bypass processing.

Processing plants typically include one or more glycol dehydrators, process units that dry the natural gas. Glycol, usually TEG, is used in dehydration units to absorb water from wet produced gas. "Lean" TEG contacts the wet gas and absorbs water. The TEG is then considered "rich." As the rich TEG is passed through a flash separator and/or reboiler for regeneration, steam containing hydrocarbon vapors is released from it. The vapors are then vented from the dehydration unit flash separator and/or reboiler still vent.

Dehydration units with a natural gas throughput below 3 MMscf per day or benzene emissions below 1 Tpy are exempted from the control, monitoring and recordkeeping requirements of Subpart HH. Although the natural gas throughput of some Marcellus horizontal shale wells in New York State could conceivably be above 3 MMscf, preliminary analysis of gas produced at Marcellus horizontal shale gas well sites in Pennsylvania indicates a benzene-content below the exemption threshold of 1 Tpy, for the anticipated range of annual gas production for wells in the Marcellus. However, the affected natural gas production facilities would still likely be required

to maintain records of the exemption determination as outlined in 40 CFR §63.774(d) (1) (ii). Sources with a throughput of 3 MMscf/day or greater and benzene emissions of 1.0 Tpy or greater are subject to the rule's emission reduction requirements. This does not necessarily mean control, depending on the location of the affected emission sources relative to "urbanized areas (UA) plus offset" or to "urban clusters (UC) with a population of 10,000 or greater" as defined in the rule.

# 6.5.1.3 Flaring Versus Venting of Wellsite Air Emissions

Well completion activities include hydraulic fracturing of the well and a flowback period to clean the well of flowback water and any excess sand (fracturing proppant) that may return out of the well. Flowback water is routed through separation equipment to separate water, gas, and sand. Initially, only a small amount of gas is vented for a period of time. Once the flow rate of gas is sufficient to sustain combustion in a flare, the gas is flared for a short period of time for testing purposes. Recovering the gas to a sales gas line is called a reduced emissions completion (REC). See Section 6.6.8 for further discussion of RECs.

Normally the flowback gas is flared when there is insufficient pressure to enter a sales line, or if a sales line is not available. There is no current requirement for REC, and the Public Service Commission (PSC) has not historically authorized construction of sales lines before the first well is drilled on a pad (see Section 8.1.2.1 for a discussion of the PSC's role and a presentation of reasons why pre-authorization of gathering lines have been suggested under certain circumstances), therefore, estimates of emissions from both flaring and venting of flowback gas are included in the emissions tables in Section 6.5.1.6. Unless PSC revisits this policy in the future in order to allow for REC, the well pad activities would be required to minimize these emissions due to the potential for relatively high short-term VOC and CO emissions, as estimated by the Industry Information Report. The modeling and regional emission assessments, as well as regulatory applicability discussions, have incorporated industry's quantifications of the short term operations associated with flaring and venting. Thus, the well permitting process would be constrained by the assumed amount of gas to be vented or flared (or the corresponding average maximum hours of operations).

Also, during drilling, gaseous zones can sometimes be encountered such that some gas is returned with the drilling fluid, which is referred to as a gas "kick." For safety reasons, the drilling fluid is circulated through a "mud-gas separator" as the gas kick is circulated out of the wellbore. Circulating the kick through the mud-gas separator diverts the gas away from the rig personnel. Any gas from such a kick is vented to the main vent line or a separate line normally run adjacent to the main vent line.

Drilling in a shale formation does not result in significant gas adsorption into the drilling fluid as the shale has not yet been fractured. Experience in the Marcellus thus far has shown few, if any, encounters with gas kicks during drilling. However, to account for the potential of a gas kick where a "wet" gas from another formation might result in some gas being emitted from the mudgas separator, an assumed wet-gas composition was used to estimate emissions.

Gas from the Marcellus Shale in New York is expected to be "dry", i.e., have little or no VOC content, and "sweet", i.e., have little or no H<sub>2</sub>S. Except for drilling emissions, two sets of emissions estimates are made to enable comparison of emissions of VOC and HAP from both dry gas production and wet gas production.

# 6.5.1.4 Number of Wells Per Pad Site

Drilling as many wells as possible from a single well pad provides for substantial environmental benefits from less road construction, surface disturbance, etc. Also, experience shows that average drilling time can be improved as more experience is gained in a shale play. Based on industry information submitted in response to Department requests, it is expected that no more than four wells could be drilled, completed, and hooked up to production in any 12-month period. Therefore, the annual emission estimates presented in Section 6.5.1.7 are based on an assumed maximum of four wells per site per year.

# 6.5.1.5 Natural Gas Condensate Tanks

Fluids that are brought to the surface during production at natural gas wells are a mixture of natural gas, other gases, water, and hydrocarbon liquids (known as condensate). Some gas wells produce little or no condensate, while others produce large quantities. The mixture typically is sent first to a separator unit, which reduces the pressure of the fluids and separates the natural gas

and other gases from any entrained water and hydrocarbon liquids. The gases are collected off the top of the separator, while the water and hydrocarbon liquids fall to the bottom and are then stored on-site in storage tanks. Hydrocarbons vapors from the condensate tanks can be emitted to the atmosphere through vents on the tanks. Condensate liquid is periodically collected by truck and transported to refineries for incorporation into liquid fuels, or to other processors.

Initial analysis of natural gas produced at Marcellus Shale horizontal gas well sites in Pennsylvania's Marcellus Shale area indicates insufficient BTEX and other liquid hydrocarbon content to justify installation of collection and storage equipment for natural gas liquids. However, in the instances where "wet" gas is encountered and there is a need to store the condensate in tanks either at the well pad or at the compressor station, potential VOC and HAP (e.g., benzene) emissions should be minimized to the maximum extent practicable and controlled where necessary. The ALL report notes that it is difficult to properly quantify the loss of vapors from these tanks, but notes that in states where substantial quantities of condensate are recovered, either a vapor recovery system or flaring is used to control emissions. If such condensate tanks are to be used in New York, a vapor recovery system would be required to be installed instead of flaring the emissions since the latter creates additional combustion emissions and other potential issues.

#### 6.5.1.6 Emissions Tables

Estimated annual emissions from drilling, completion and production activities are based on industry's response to the Department's information requests<sup>343</sup> (hereafter Industry Information Report) that a maximum number of four wells would be drilled at a given pad in any year (see further discussion in the modeling section). These estimates are presented in Table 6.7, Table 6.8, Table 6.9, and Table 6.10 below.

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<sup>&</sup>lt;sup>343</sup> ALL Consultant Information Request Report on behalf of IOGANY, dated September 16, 2010.

Table 6.7 - Estimated Wellsite Emissions (Dry Gas) - Flowback Gas Flaring (Tpy)(Updated July 2011)

|            | Drilling | Completion | Production | Subtotal | Flowback Gas | Total |
|------------|----------|------------|------------|----------|--------------|-------|
| PM         | 0.5      | 0.2        | 0.2        | 0.9      | 1.4          | 2.3   |
| $NO_x$     | 15.1     | 5.8        | 3.8        | 24.7     | 4.9          | 29.6  |
| CO         | 8.3      | 3.2        | 9.2        | 20.7     | 24.5         | 45.2  |
| VOC        | 0.8      | 0.2        | 2.4        | 3.4      | 0.7          | 4.1   |
| $SO_2$     | 0.02     | 0.01       | 0.07       | 0.1      | 0.0          | 0.1   |
|            |          |            |            |          |              |       |
| Total HAPs | 0.09     | 0.02       | 0.03       | 0.14     | 0.08         | 0.22  |

Table 6.8 - Estimated Wellsite Emissions (Dry Gas) - Flowback Gas Venting (Tpy)(Updated July 2011)

|            | Drilling | Completion | Production | Subtotal | Flowback Gas | Total |
|------------|----------|------------|------------|----------|--------------|-------|
| PM         | 0.5      | 0.2        | 0.2        | 0.9      | 0.0          | 0.9   |
| $NO_x$     | 15.1     | 5.8        | 3.8        | 24.7     | 0.0          | 24.7  |
| СО         | 8.3      | 3.2        | 9.2        | 20.7     | 0.0          | 20.7  |
| VOC        | 0.8      | 0.2        | 2.4        | 3.4      | 0.6          | 4.0   |
| $SO_2$     | 0.02     | 0.01       | 0.07       | 0.1      | 0.0          | 0.1   |
|            |          |            |            |          |              |       |
| Total HAPs | 0.09     | 0.02       | 0.03       | 0.14     | 0.0          | 0.14  |

Table 6.9 - Estimated Wellsite Emissions (Wet Gas) - Flowback Gas Flaring (Tpy) (Updated July 2011)

|            | Drilling | Completion | Production | Subtotal | Flowback Gas | Total |
|------------|----------|------------|------------|----------|--------------|-------|
| PM         | 0.5      | 0.2        | 0.2        | 0.9      | 1.4          | 2.3   |
| $NO_x$     | 15.1     | 5.8        | 3.8        | 24.7     | 4.9          | 29.6  |
| CO         | 8.3      | 3.2        | 9.2        | 20.7     | 24.5         | 45.2  |
| VOC        | 0.8      | 0.2        | 2.4        | 3.4      | 0.7          | 4.1   |
| $SO_2$     | 0.02     | 0.01       | 0.07       | 0.1      | 0.22         | 0.31  |
| Total HAPs | 0.09     | 0.02       | 0.31       | 0.42     | 0.69         | 1.11  |

Table 6.10 - Estimated Wellsite Emissions (Wet Gas) - Flowback Gas Venting (Tpy) (Updated July 2011)

|            | Drilling | Completion | Production | Subtotal | Flowback Gas | Total |
|------------|----------|------------|------------|----------|--------------|-------|
| PM         | 0.5      | 0.2        | 0.2        | 0.9      | 0.0          | 0.9   |
| $NO_x$     | 15.1     | 5.8        | 3.8        | 24.7     | 0.0          | 24.7  |
| СО         | 8.3      | 3.2        | 9.2        | 20.7     | 0.0          | 20.7  |
| VOC        | 0.8      | 0.2        | 2.4        | 3.4      | 21.9         | 25.3  |
| $SO_2$     | 0.02     | 0.01       | 0.07       | 0.1      | 0.0          | 0.1   |
| Total HAPs | 0.09     | 0.02       | 0.31       | 0.42     | 0.002        | 0.422 |

It is important to understand that the "totals" columns in these tables are not meant to be compared to the major source thresholds discussed in section 6.5.1.2 for the purpose of determining source applicability to the various regulations. This is because these estimates include emissions from activities which are not considered stationary sources, as detailed in the discussions in Section 6.5.1.8. These estimates should be looked upon merely as giving a relative sense of the expected well pad emissions and what the relation is to major source thresholds.

## 6.5.1.7 Offsite Gas Gathering Station Engine

For gas gathering compression, it is anticipated that most operators would select a large 4-stroke lean-burn engine because of its fuel efficiency. A typical compressor engine is the 1,775-hp Caterpillar G3606, which is the engine model used for the analysis.

The final revision to NESHAPs Subpart ZZZZ has placed very strict limits on formaldehyde emissions from reciprocating internal combustion engines (see Appendix 17). Future, 4-stroke lean-burn engines would be required to have an oxidation catalyst that would reduce formaldehyde emissions by approximately 90%.

The annual emissions data for a typical gas gathering compressor engine is given in Table 6.11 below.<sup>344</sup>

Table 6.11 - Estimated Off-Site Compressor Station Emissions (Tpy)

| Component | Controlled 4-Stroke Lean Burn Engine |
|-----------|--------------------------------------|
| PM        | 0.5                                  |
| $NO_x$    | 33.3                                 |
| CO        | 6.6                                  |
| $SO_2$    | 0.0                                  |
| Total VOC | 5.0                                  |
| Total HAP | 2.7                                  |

<sup>&</sup>lt;sup>344</sup> A<u>LL Consulting</u> August 26, 2009.

6.5.1.8 Department Determinations on the Air Permitting Process Relative to Marcellus Shale High-Volume Hydraulic Fracturing Development Activities.

A determination would first be made as to whether these internal combustion engines (ICEs) would qualify for the definition of non-road or stationary sources. This, in turn, determines whether the engines are subject to requirements such as NSPS or NESHAPs.

When considering applicability of these rules, engines can fall into three general classes: stationary, mobile, or nonroad. The applicable NSPS regulations (40 CFR Part 60, Subpart IIII and Subpart JJJJ) and NESHAP (40 CFR Part 63 Subpart ZZZZ) define stationary internal combustion engines as excluding mobile engines and nonroad engines. The New York State definition of stationary sources given in 6.5.1 also notes the non-road engine exclusion. The latter engines are defined at 40 CFR Part 1068 (General Compliance Provisions for Nonroad Program), which is virtually the same as it appears in 40 CFR Part 89 (Control of Emissions from New and In-Use Nonroad Compression-Ignition Engines) as well as in New York's regulations at NYCRR Part 200.1, as given in Section 6.5.1. Paragraph (1)(iii) of the definition describes a nonroad engine that would be portable or would be part of equipment that would be considered portable, with the exception given in paragraph 2(iii) if the engines are to remain at the same location for more than 12 months.

It is clear from the Industry Information Report that the engines used to power the drilling and well development equipment would be used at a given well pad for maximum of less than half a year (see discussions in ALL, 8/26/09 and the modeling section on the timeframes of engine use), even if the maximum of four wells per pad were to be completed in a year. Thus, these engines are considered as nonroad engines and are not subject to the NSPS, NESHAP or permitting requirements.

However, as detailed in the following section, the environmental consequences of these engines are fully analyzed and mitigated where necessary in keeping with SEQRA. For example, the use of ULSF with a 15 ppm sulfur content would be required for use in all drilling and well development equipment engines. This limit is required for stationary engines in the final NESHAPS Subpart ZZZZ rule as discussed in Appendix 17. In addition, a set of control measures would be required on most of these engines in order to meet NAAQS, as fully

addressed in the modeling analysis section. The permitting of the various activities associated with drilling and development activities in the Marcellus Shale would be consistent with regulatory scheme in 6 NYCRR Part 200, et. seq. for regulating emissions of air pollutants. Thus, the Department would not subject the nonroad engines to the regulatory requirements applicable to stationary source, such as the determination of what constitutes a major source per Part 201. In instances throughout the country reviewed by the Department in terms of permitting gas drilling and production activities, the determination of a stationary source or facility has relied on the association of the compressor stations and nearby well emissions, but in none of these were the nonroad engine emissions included in the permitting emission calculations. This approach would also be followed in New York as the appropriate regulatory scheme.

Thus, in accounting for the well site operation emissions in the permitting process, the emissions from Tables 1 to 4 above would only include the remaining activities at the site which are essentially a small line heater (1 million Btu) a small compressor (150 horsepower), and possibly a flare. Tables 1 to 4 indicate that for the three higher emission pollutants, NO<sub>x</sub>, CO and VOCs, these sources would add up to a maximum of 8.7, 33.7, and 3.1 Tpy, respectively, under the normal dry gas scenario for each pad. In the unlikely event of encountering "wet" gas, the VOC emissions could be 24.3 Tpy. However, these CO and VOC emissions are associated with the transient sources, the flare and gas venting, respectively, which are to be minimized, as would be apparent in the discussions to follow. In addition, in the unlikely event that a glycol dehydration would be located at a well site instead of the compressor station, the strict regulatory requirement noted in Section 6.5.1 would limit the VOC (benzene) emissions to below 1 Tpy. Thus, total HAPs emissions from a well pad would be much less than even the major source threshold of 10 Tpy for a single HAP.

Therefore, the process which the Department would follow in permitting the air emissions from Marcellus Shale activities would start with the compressor station permit application review. As noted in Section 8.1.2.1, this SGEIS for drilling wells is not meant to address the full extent of the compressor station permitting and the environmental consequences, which falls under the purview of the PSC and would be dealt with on a case by case basis. The applicable Public Service Law, Article VII, would be followed in which PSC would be the lead agency for the environmental review, however the Department would remain the agency responsible for

reviewing and acting on the air permit application. In this review, the Department would incorporate all of the applicable regulations, including the determination of what constitutes a source or facility. The air quality analysis has considered the impacts of a potential compressor station which is hypothetically placed next to the well pad in the modeling assessment of standards and other compliance thresholds.

Section 112(n) of the CAA (Section 112) applies specifically to HAPs. The EPA, on September 22, 2009, clarified that for the purposes of New Source Review (NSR) and Title V applicability review, the process of facility determination should include a detailed consideration of the traditional set of three criteria used by EPA in past actions. In this determination, a set of related and adjacent activities could be "aggregated" if they meet the requirements of the criteria.

The Department would follow EPA's process for the determination of a stationary source or facility for criteria pollutants, as also guided by recent applicability determinations by EPA and other states. Details of the Department's approach are presented in Appendix 18. The process would involve requesting information during the compressor station permit application phase using a set of questions framed from previous EPA determinations. A sentinel aspect of EPA's regulation and policy, which New York's approach is adapting, is the use of case-by-case information to make an informed decision. That process would also consider information requested on drilling wells which could be associated with the compressor stations.

### 6.5.2 Air Quality Impact Assessment

#### 6.5.2.1 Introduction

As part of the Department's effort to address the potential air quality impacts of horizontal drilling and hydraulic fracturing activities in the Marcellus Shale and other low-permeability gas reservoirs, an air quality modeling analysis was undertaken by the Department's Division of Air Resources (DAR). The original modeling analysis was carried out to determine whether the various expected operations at a "typical" multi-well site would have the potential for any adverse air quality impacts, and it addressed a number of issues raised in public comments during the SGEIS scoping process. The analysis also incorporated subsequently-developed information on operational scenarios specific to multi-well horizontal drilling and hydraulic fracturing, to help determine possible air permitting requirements.

The initial modeling analysis has been updated based on information from both the Industry Information Report and related public information which has become available since September 2009. In particular, industry has indicated that: 1) simultaneous drilling and completion operations at a single pad would not occur; 2) the maximum number of wells to be drilled at a pad would be four in any 12-month period; and 3) flowback impoundments are not contemplated. The effects of these operational changes are discussed where appropriate. It is to be noted that the revision from maximum of ten wells down to four wells per pad per year affects only the annual emissions and the modeled annual impacts and not the short term impacts. Therefore, the annual impacts were revisited to determine if the reduced emissions had an effect on the previous conclusions reached on standards compliance. In instances where previous impacts due to emissions using ten wells did not pose an exceedance, the annual impacts have not been recalculated since these represent conservative concentrations versus the revised maximum of four well operations. Instances where this approach is used are noted in the subsequent discussions.

Due to remaining issues with exceedances of the 24-hour PM2.5 ambient standard and the adoption of new 1-hour SO<sub>2</sub> and NO<sub>2</sub> standards by EPA since the initial modeling analysis, a supplemental modeling analysis was performed. The approach to this assessment and the consequent results are presented in a separate section which follows this section. That assessment has incorporated the discussions from an industry modeling exercise for PM2.5 and PM10, as well as more recent EPA guidance documents on modeling for these pollutants.

This section presents the initial air quality analysis undertaken by DAR staff based on operational and emissions information supplied mainly by industry and its consultant in a submission hereafter referred to as the Industry Information Report. To a limited extent, certain supplemental information from ICF International's report to NYSERDA was also used. The applicability determinations of the Department's air permitting regulations and the verification approach to the emission calculations are contained in Section 6.5.2.

345 ALL Consulting, 2009,

<sup>&</sup>lt;sup>346</sup> ICF Task 2, 2009,

To the extent that the information being used was for the modeling of a generic multi-well site and its operations, it was necessary to reconcile and define a "worst case" scenario for the various activities in terms of expected impacts. Certain assumptions were made on the type and sizes of equipment to be used, the potential for simultaneous operation of the equipment on a short-term basis (i.e., hourly and daily), and the duration of these activities over a period of a year in order to be able to compare impacts to the corresponding ambient thresholds. The supplemental modeling analysis indicates that, although the operational time frame for certain equipment (e.g., engines) over a given year would be reduced according to the Industry Information Report, 347 the consequences of these reduced annual emissions are only qualitatively addressed in the following sections since these do not affect any of the initial conclusions reached on annual impacts. That is, the reduced annual emissions from certain operations which were initially demonstrated to meet the corresponding standards and thresholds would only be lowered by this new information.

The air quality analysis relied upon recommended EPA and the Department's air dispersion modeling procedures to determine "worst case" impacts of the various operations and activities identified for the horizontal multi-well sites. Dispersion modeling is an acceptable tool, and at times the only option, to determine the impacts of many source types in permitting activities and environmental impact statements. Where necessary, the analysis approach relied on assumed worst case emissions and operations scenarios due to not only the nature of this generic assessment, but also because detailed model input data for the sources and their relative locations on a typical well pad cannot be simply identified or analyzed. Modeling was performed for various criteria pollutants (those with NAAQS) and a set of non-criteria pollutants (including toxics) for which New York has established a standard or other ambient threshold levels. Some of these toxic pollutants were identified in public comments during the SGEIS scoping process and were quantified to the extent possible for both the modeling and applicability determinations.

The following sections describe the basic source categories and operations at a typical multi-well site with hydraulic fracturing, the modeling procedures and necessary input data, the resultant impacts, and a set of conclusions drawn from these results. These conclusions are meant to

<sup>&</sup>lt;sup>347</sup> A<u>LL</u> Consulting, 2010.

guide the set of conditions under which a site specific assessment might or might not be necessary. Based on information in the Industry Information Report and an update to EPA's dispersion model, the initial PM10/PM2.5 modeling approach and conclusions have been updated.

# 6.5.2.2 Sources of Air Emissions and Operational Scenarios

In order to properly estimate the air quality impacts of the set of sources at a single pad with multiple horizontal wells, the operating scenarios and associated air emission sources would be correctly represented. Since these operations have a number of interdependent as well as independent components, the Department has defined both the short-term and long term emission scenarios from the various source types in order to predict conservative, yet realistic impacts. The information used to determine the emission sources and their operating scenarios and constraints, as well as the associated emission rates and parameters, were provided by the Industry Information Report, while certain operational scenario restrictions were presented in the ICF report, which reflects information obtained from industry with drilling activities in other states. Where necessary, further data supplied by industry or determined appropriate by DMN was used to fill in data gaps or to make assumptions. In some of these instances, the lack of specific information necessitated a worst-case assumption be made for the purposes of the modeling exercise. Examples of the latter include defining "ambient air" based on the proximity of public access to the well pad and the likely structure dimensions to calculate their influence on the stack plumes.

The Industry and ICF Reports indicate three distinct operation stages and four distinct source types of air emissions for developing a representative horizontally-drilled multi-well pad. The phases are drilling, completion, and gas production, each of which has either similar or distinct sources of air emissions. These phases and the potential air pollution sources are presented in the Industry Information Report, Section 2.1.5 and Exhibit 2.2.1 of the ICF report, and in Chapter 5 of the SGEIS, and would only be briefly noted herein. Of the various potential sources of air emissions, a number have distinct quantifiable and continuous emissions which lend themselves to modeling. On the other hand, the ICF report also identifies other generic sources of minor fugitive emissions (e.g., mud return lines) or of emergency release type (e.g., BOP stack), or of a pollutant which is quantified only as of "generic" nature (total VOCs for

tanks) which cannot be modeled to any reliable extent without a well-defined source. The best approach to address these sources is to apply best minimization techniques, as recommended in Section 6.5.1.5 for condensate tanks. However, in instances where speciated VOCs or HAPs are available and provided by industry, such as for the glycol dehydrator and flowback venting of gas, the modeling was used to predict impacts which were then compared to available ambient thresholds.

The total operations associated with well drilling can be assigned to three "types" of potential sources: 1) combustion from engines, compressors, line heaters, and flares; 2) short-term venting of gas constituents which are not flared; and 3) emissions from truck activities near the well pad. Each of these source categories have limitations in terms of the size and number of the needed equipment, their possible simultaneous operations over a short-term period (e.g., 24-hour), and the time frames over which these equipment or activities could occur over a period of one year, which effects the corresponding annual impacts. Some of these limitations are described in the Industry Information Report. These limitations and further assumptions were taken into account in the modeling analysis, as further discussed in Section 6.5.2.3.

Many of the sources for which the Industry Information Report tabulates the drilling, completion and production activities are depicted in the typical site layout represented schematically in Exhibit 2.1.3 of the ICF report. The single pad for multi-horizontal wells is confined to an area of about 150 meters (m) by 150 m as a worst case size of the operations. From this single pad, wells are drilled in horizontal direction to develop an area of about one square mile. The initial industry report noted the possibility of up to ten horizontal wells being eventually drilled and completed per pad over a year's time, while the ICF report notes that simultaneous drilling and completion on the same pad would be limited to a single operation for each. This limitation was determined appropriate by DMN for analysis of short-term impacts. Thus, the simultaneous operations on a pad for the assessment of impacts of 24 hours or less is limited to the equipment necessary to drill one well and complete another. In addition, according to DMN, there is a potential that a third well's emissions could be flared at the same time as these latter operations. Thus, this source was also included in the simultaneous operation scenario for criteria pollutants. The Industry Information Report indicates that the number of wells drilled in a year at a given well pad would be four and asserts that there would not be any simultaneous operations of the

well drilling and completion equipment engines. These revisions are incorporated in the supplemental modeling analysis section. Their influence on the results in this section is addressed in places where deemed of consequence.

It should be noted that no emissions of criteria pollutants resulting from uncontrolled venting of the gas are expected. The other sources which could emit criteria pollutants are associated with the production phase operations; that is, the off-site compressors and line heaters could be operating simultaneously with the single pad drilling, completion and flaring operations. The Industry Information Report provides data for a possible "on-site" line heater instead of at the compressor station and this source was placed on the pad area and provides for a more conservative impact.

The Industry Information Report also provides emission data for the non-criteria pollutants as species of VOCs or HAPs associated with both combustion and gas venting. Review of this information indicates two essentially different sets of sources which can be treated independently in the modeling analysis. The first set is the gas venting sources: the mud-gas separator, the flowback gas venting, and the glycol dehydrator. These sources emit a distinct set of pollutants associated with the "wet" gas scenario, defined in the Industry Information Report as containing "heavier" hydrocarbons such as benzene. The industry and ICF reports note that gas samples in the Marcellus Shale have detected neither these heavier species of VOCs, nor H<sub>2</sub>S. However, the Industry Information Report also notes the possibility of gas pockets with "wet" gas and provides associated emissions. To be comprehensive, the modeling analysis has calculated the impacts of these species which could be realized in the westernmost part of New York according to DMN.

The Industry Information Report also notes that gas venting is a relatively short-term phenomenon, especially during the flowback period where the vented gas is preferentially flared after a few hours of venting. Since there are essentially no simultaneous short-term emissions expected of the same pollutants at the pad from processes other than flowback venting, coupled with the clear dominance of the flowback venting emissions of these pollutants, the modeling was simplified for this scenario and only the short-term impacts were determined, as described in more detail in Section 6.5.1.3. The second set of non-criteria pollutant emissions presented in the Industry Information Report is associated mainly with combustion sources. These non-

criteria pollutants could be emitted over much longer time periods, considering these sources are operated over these longer periods, both per-well drilling activity and potential multi-well operations over a given year. Thus, for these pollutants, both short-term and annual impacts were calculated. It should be noted that, since the glycol dehydrator could operate for a full year also, its emissions of the same pollutants as those due to combustion were also included in this assessment of both short-term and annual toxic impacts. Furthermore, the flare emissions are included in the combustion scenario (and not in the venting), as the flaring of flowback gas results in over 95% destruction of these pollutants.

In addition, due to the conversion of H<sub>2</sub>S to SO<sub>2</sub> during flaring, the flare was included in the criteria pollutant simultaneous operations scenario modeling. Table 6.12 summarizes the set of sources and the pollutants which have been modeled for the various simultaneous operations for short-term impacts. The specific modeling configuration and emissions data of the various sources are discussed in Section 6.5.2.3.

The last type of emission source associated with the multi-well operations is truck traffic. An estimate of the number of trucks needed for the various activities at a single well pad, including movement of ancillary equipment, delivery of fresh water and proppant/additives, and the hauling of flowback is presented in Section 6.11. It should be first noted that direct emissions from mobile sources are controlled under Title II of the CAA and are specifically exempt from permitting activities. Thus, these emissions are also not addressed in general in a modeling analysis, with two exceptions. At times, the indirect emissions of fugitive particulate matter are modeled when estimates of emissions are large. The latter occurs mainly due to poor dust control measures and the best approach to mitigate these emissions is to have a dust control plan. In addition, emissions of PM2.5 from mobile sources associated with a project and which occur on-site are to be addressed by the Department's Commissioner's Policy CP-33. Again, if these emissions are large enough, a modeling analysis is performed for an EIS. For the assessment of PM2.5 per CP-33, the emission calculations are not to include those associated with incidental roadway traffic away from the onsite operations.

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<sup>348</sup> http://www.dec.ny.gov/chemical/8912.html.

Emissions of both PM10 and PM2.5 due to truck operations at the well pad were initially calculated by DAR's Mobile Source Panning Section based on the movement of total number of trucks on-site for the drilling of one well. These emissions were then multiplied by the 10 potential wells which might be drilled over a year, and resulted in relatively minor quantities of 0.2 Tpy maximum PM2.5 emissions. This is consistent with the limited use of trucks at the well pad. These emissions are well below the CP-33 threshold of 15 Tpy. Thus, no modeling was performed for these pollutants and any necessary mitigation scheme for these would be the application of an appropriate dust control methods and similar limitations on truck usage, such as inordinate idling.

In order to address on-road truck traffic movement and emissions in the area underlain by the Marcellus Shale, estimates of regional emissions have been calculated based on information provided in the Industry Information Report. These regional emissions and their consequence are discussed in the section to follow. In addition, at the well pad, EPA's updated emission model MOVES (Motor Vehicle Emission Simulator) was used instead of the MOBILE 6e model used in the initial analysis. The MOVES model was also applied to generate regional emissions of on-road mobile sources associated with Marcellus Shale well development and included PM2.5 emissions. These estimates have been incorporated in the discussions of regional annual emissions. Results from the MOVES model indicate that the very low PM2.5 emissions initially estimated for a single pad are unchanged.

# 6.5.2.3 Modeling Procedures

EPA<sup>349</sup> and Department<sup>350</sup> guidelines on air dispersion modeling recommend a set of models and associated procedures for assessing impacts for a given application. For stationary sources with "non-reactive" pollutants and near-field impacts, the refined AERMOD model (latest version, 07026) and its meteorological and terrain preprocessors is best suited to simulate the impacts of the sources and pollutants identified in the Marcellus Shale and other gas reservoir operations. This model is capable of providing impacts for various averaging times using point, volume or area source characteristics, using hourly meteorological data and a set of receptor locations in the

Appendix W to 40 CFR Part 51. http://www.epa.gov/ttn/scram/guidance\_permit.htm.

<sup>350</sup> http://www.dec.ny.gov/chemical/8923.html.

surrounding area as inputs. The model simulates the impact of "inert" pollutants such as SO<sub>2</sub>, NO<sub>2</sub>, CO, and particulates without taking into account any removal or chemical conversions in air, which provides for conservative ambient impacts. However, these effects are of minor consequences within the context of plume travel time and downwind distances associated with the maximum ambient impact of pollutants discussed in this section.

AERMOD also does not treat secondary formation of pollutants such as  $O_3$  from  $NO_x$  and VOCs, but it can model the non-criteria and toxic pollutant components of gas or VOC emissions in relation to established ambient thresholds. There does not exist a recommended EPA or Department "single" source modeling scheme to simulate  $O_3$  formation from its precursors. This would involve not only complex chemical reactions in the plumes, but also the interaction of the regional mix of sources and background levels. Such an assessment is limited to regional scale emissions and modeling and is outside the scope of the modeling analysis undertaken for this section. However, the potential consequences of regional emissions of VOCs and  $NO_x$  are presented in Section 6.5.3.

Thus, the AERMOD model was used with a set of emission rates and source parameters, in conjunction with other model input data discussed in the following subsections, to estimate maximum ambient impacts, which were then compared to established Federal and New York State ambient air quality standards (AAQS) and other ambient thresholds. The latter are essentially levels established by the Department's Division of Air Resources (DAR) program policy document DAR-1. These levels are the 1-hour SGCs and annual AGCs (short-term and annual guideline concentration, respectively). Where certain data on the chemicals modeled and the corresponding ambient thresholds were missing, New York State Department of Health (NYSDOH) staff provided the requested information. For the thresholds, the Department's Toxics Assessment section then calculated the applicable SGCs and AGCs. The modeling procedures also invoke a number of "default" settings recommended in the AERMOD user's guide and EPA's AERMOD Implementation Guide. For example, the settings of potential wells are not expected to be in "urban" locations, as defined for modeling purposes and, thus, the rural option was used. Other model input data are described next.

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<sup>351</sup> http://www.dec.ny.gov/chemical/30560.html.

### Meteorological Data

The AERMOD model requires the use of representative hourly meteorological data, which includes parameters such as wind speed, wind direction, temperature and cloud cover for the calculation of transport and dispersion of the plumes. A complete set of all the parameters needed for modeling is generally only available from National Weather Service (NWS) sites. The "raw" data from NWS sites are first pre-processed by the AERMET program and the AERSURFACE software using land use data at the NWS sites, which then create the necessary parameters to be input to AERMOD. There is a discrete set of NWS sites in New York which serves as a source of representative meteorological data sites for a given project. However, for this analysis, the large spatial extent of the Marcellus Shale necessitated the use of a number of the NWS site data in order to cover the meteorological conditions associated with possible well drilling sites throughout the State.

Figure 6.10 presents the spatial extent of the Marcellus Shale and the six NWS sites chosen within this area and deemed adequate for representing meteorological conditions for the purpose of dispersion modeling of potential well sites. It was judged that these sites would adequately envelope the set of conditions which would result in the maximum impacts from the relatively low-elevation or ground-level sources identified as sources of air pollutants. In addition, EPA and Department modeling guidance recommends the use of five years of meteorological data from a site in order to account for year to year variability. For the current analysis, however, the Department has chosen two years of data per site to gage the sensitivity of the maxima to these data and to limit the number of model calculations to a manageable set. It was determined that impacts from the relatively low-elevation sources would be well represented by the total of 12 years of data used in the analysis.

This analysis is conservative from the standpoint of the number of data years used. Certain public comments <sup>352</sup> recommended that the Department should use the EPA-recommended five years of data for its analysis. However, these comments do not fully recognize the conservative nature of using 12 years of meteorological data to determine the worst case impact for any potential site in the Marcellus Shale play. While the EPA and the Department guidance to use

<sup>&</sup>lt;sup>352</sup> AKRF Consultants, memo dated 12/3/2009, p. 2.

five years of data applies to individual meteorological site analysis to account for possible climatological variability at the particular site, the use of 12 years of data from six different sites has a similar conservatism built into it by the end use of the overall maxima for any well pads or compressor stations. That is, the overall maxima for any specific pollutant and averaging time could be controlled by meteorological data from different NWS sites, but these maxima are being used for all potential sites in the Marcellus Shale play regardless of whether they might experience these meteorological conditions. A review of the results discussed in the next section and in Table 6.16 confirms this conclusion. Thus, it is deemed that the use of two years of data from six NWS sites to assess the maximum potential impacts is conservative.

The NWS sites and the two years of surface meteorological data which were readily available from each site are presented in Table 6.13, along with latitude and longitude coordinates. In addition to these surface sites, upper air data is required as input to the AERMOD model in order to estimate certain meteorological parameters. Upper air data is only available at Buffalo and Albany for the sites chosen for this analysis, and were included in the data base. It should be noted that upper air data is not the driving force relative to the surface data in modeling low-elevation source impacts within close proximity of the sources, as analyzed in this exercise. The meteorological data for each year was used to calculate the maximum impacts per year of data and then the overall maxima were identified from these per the regulatory definitions of the specific AAQS and SGCs/AGCs, as detailed in the subsequent subsection.

#### Receptor and Terrain Input Data

Ground level impacts are calculated by AERMOD at user defined receptor locations in the area surrounding the source. These receptors are confined to "ambient air" locations to which the public has access. Current DMN regulations define a set of "set back" distances from the well sites to roadways and residences. However, these set back distances (e.g., 25m) are defined from the wellhead for smaller "footprint" vertical wells relative to the size of the multi-pad horizontal wells. Furthermore, EPA's strict definition of ambient air only excludes areas to which the public is explicitly excluded by enforceable measures such as fences, which might not be normally used by the industry. Thus, in order to determine the potential closest location of receptors to the well site, the modeling has considered receptors at distances as close as the boundary of a 150m by 150m well pad. On the other hand, it is clear from diagrams and pictures

of sample sites that the public would have no access to within the well pad area. However, the closest receptor to any of the sources was limited to 10m to allow for a minimum practical "buffer" zone between the equipment on the pad and its edge.

The location of the set of modeled receptors is an iterative process for each application in that an initial set is used to identify the distance to the maximum and other relatively high impacts, and then the grid spacing may need to be refined to assure that the overall maxima are properly identified. For the type of low-elevation and ground level sources which dominate the modeled set in this analysis, it is clear that maximum impacts would occur in close proximity to the sources. Thus, a dense grid of 10m spacing was placed along the "fencelines", and extended on a Cartesian grid at 10 m grid spacing out to 100 m from the sources in all directions. In a few cases, the modeling grid was extended to a distance of 1000 m at a grid spacing of 25 m from the 100 m grid's edge in order to determine the concentration gradients. For the combustion and venting sources, an initial grid at 10m increment was placed from the edge of the 150 m by 150 m pad area out to 1000 m, but this grid was reduced to a Cartesian grid of 20 m from spacing the "fenceline" to 500 m in order to reduce computation time. The revised receptor grid resolution was found to adequately resolve the maxima as well for the purpose of demonstrating the anticipated drop off of concentrations beyond these maxima.

The AERMOD model is also capable of accounting for ground level terrain variations in the area of the source by using U.S. Geological Survey Digital Elevation Model (DEM) or more recent National Elevation Data (NED) sets. However, for sources with low emission release heights, the current modeling exercise was performed assuming a horizontally invariant plane (flat terrain) as a better representation of the impacts for two reasons. First, given the large variety of terrain configurations where wells may be drilled, it was impractical to include a "worst case" or "typical" configuration. More importantly, the maximum impacts from the low-elevation sources are expected to occur close-in to the facility site, and any variations in topography in that area was determined to be best simulated by AERMOD using the concept of "terrain following" plumes.

It should be clarified that this discussion of terrain data use in AERMOD is distinct from the issue of whether a site might be located in a complex terrain setting which might create distinct

flow patterns due to terrain channeling or similar conditions. These latter mainly influence the location and magnitude of the longer term impacts and are addressed in this analysis to the extent that the set of meteorological data from six sites included these effects to a large extent. In addition, the air emission scenarios addressed in the modeling for the three operational phases and associated activities are deemed to be more constrained by short-term impacts due to the nature and duration of these operations, as discussed further below. For example, the emissions from any venting or well fracturing are intermittent and are limited to a few hours and days before gas production is initiated.

#### Emissions Input Data

EPA and Department guidance require that modeling of short-term and annual impacts be based on corresponding maximum potential and, when available, annual emissions, respectively. However, guidance also requires that certain conservative assumptions be made to assure the identification of maximum expected impacts. For example, the short-term emission rates have to represent the maximum allowable or potential emissions which could be associated with the operations during any given set of hours of the meteorological data set and the corresponding averaging times of the standards. This is to assure that conditions conducive to maximum impacts are properly accounted for in the varying meteorological conditions and complex dependence of the source's plume dispersion on the latter. Thus, for modeling of all short-term impacts (up to 24 hours); the maximum hourly emission rate is used to assure that the meteorological data hours which determine the maximum impacts over a given period of averaging time were properly assessed.

Based on the information and determinations presented in Section 6.5.1.2 on the set of sources and pollutants which need to be modeled, the necessary model input data was generated. This data includes the maximum and annual emission rates for the associated stack parameters for all of the pollutants for each of the activities. In response to the Department's request, industry provided the necessary model input data for all of the activities at the multi-well pad site, as well as at a potential offsite compressor. These data were independently checked and verified by DAR staff and the final set of source data information was supplied in the Industry Information Report noted previously. Although limited source data were also contained in the ICF report, the

data provided by industry were deemed more complete and could be substantiated for use in the modeling.

The sources of emissions specific to Marcellus Shale operations are treated by AERMOD as either point or area sources. Point sources are those with distinct stacks which can also have a plume rise, simulated by the model using the stack temperatures and velocities. An example of a point source is the flare used for short term periods. Area sources are generally low or ground level sources of distinct spatial dimensions which emit pollutants relatively uniformly over the whole of the area. The previously proposed flowback water impoundments are a good example of area sources. In addition to the emission rates and parameters supplied by industry, available photographs and diagrams indicated that many of the stacks could experience building downwash effects due to the low stack heights relative to the adjacent structure heights. In these instances, downwash effects were included in a simplified scheme in the AERMOD modeling by using the height and "projected width" of the structure. These effects were modeled to assure that worst case impacts for the compressors and engines were properly identified. The specific model input data used is described next, with criteria and non-criteria source configurations presented separately for convenience.

Criteria Pollutant Sources - The emission parameters and rates for the combustion source category at a multi-horizontal well pad were taken from data tables provided in the Industry Information Report. In some instances, additional information was gathered and assumptions made for the modeling. The report provides "average" and maximum hourly emission rates, respectively, of the criteria pollutants in Tables 7 and 8 for the drilling operations, Tables 14, 15, 20 and 21 for the completion phase operations, Table 18 for the production phase sources, and Table 24 for the offsite compressor. It should be noted that the criteria pollutant source emissions in these tables are not affected by the dry versus wet gas discussions, with the exception of SO<sub>2</sub> emissions from flaring of H<sub>2</sub>S in wet gas. For this particular pollutant, the flare emission rate from Table 21 was used. Furthermore, the modeling has included the off-site compressor in lieu of the smaller onsite compressor at the wellhead and an onsite line heater instead of an offsite one in order to determine expected worst case operations impacts.

As discussed previously, initial modeling of both short-term and annual impacts were based on the maximum hourly emissions rates, with further analysis of annual impacts performed using more representative long term emissions only when necessary to demonstrate compliance with corresponding annual ambient thresholds. For the short-term impacts (less than 24-hour), it was assumed that there could be simultaneous operations of the set of equipment at an on-site pad area for one well drilling, one well completion, and one well flaring, along with operations of the onsite line heater and off site compressor for the gas production phase for previously-completed wells. For the modeling of the 24-hour PM2.5 impacts for the Supplemental Modeling section, the simultaneous operation scenario was not used based on the Industry Information Report. It should be clarified that although AERMOD currently does not include the flare source option in the SCREEN3 model, the heat release rate provided in Table 15 of the Industry Information Report was used to calculate the minimum flare "flame height" as the stack height for input to AERMOD.

The placement of the various pieces of equipment in <u>Table 6.12</u> on a well pad site was chosen such as not to underestimate maximum offsite as well as combined impacts. For example, the schematic diagram in the ICF report represents a typical set up of the various equipment, but for the modeling of the sources which could be configured in a variety of ways on a given pad, the locations of the specific equipment were configured on a well pad without limiting their potential location being close to the property edge. That is, receptors were placed at distances from the sources as if these were near the edge of the property, with the "buffer zone" restriction noted previously. This was necessary since many of these low level sources could have maximum impacts within the potential 150m distance to the facility property and receptors could not be eliminated in this area.

At the same time, however, it would be unrealistic to locate all of the equipment or a set of the same multi-set equipment at an identical location. That is, certain sources such as the flare are not expected to be located next to the rig and the associated engines due to safety reasons. In addition, there are limits to the size of the "portable" engines which are truck-mounted, thus requiring a set of up to 15 engines placed adjacent to each other rather than treating these as a single emission point. Since there were some variations in the number and type of the multi-source engines and compressors specifically used for drilling and completion, a balance was

reached between using a single representative source, with the corresponding stack parameters and total emissions, versus using distinct individual source in the multi-source set. This determination was also dictated by the relative emissions of each source.

The modeling used a single source representation for the drilling engines and compressors from Table 8, while for the fracturing pump engines, five sources were placed next to each other to represent three-each of the potential fifteen noted in Table 15 of the Industry Information Report. The total emission rates for the latter sources were divided over the five representative sources in proper quantities. This scenario was revised for the Supplemental Modeling section by modeling each of the 15 completion equipment engines as individual point sources. The rest of the sources are expected to either be a single equipment or are in sets such that representation as a single source was deemed adequate. The one exception was the modeling of the NO<sub>2</sub> 1-hour standard as describe in the next section. Using sample photographs from existing operations in other states, estimates of both the location as well as the separation between sources were determined. For example, the size of the trucks with mounted fracturing engines was used to determine the separation between a row of the five representative sources. These photographs were also used to estimate the dimension of the "structures" which could influence the stack plumes by building downwash effects. All of the sources were deemed to have a potential for downwash effects, except for the flare/vent stack. The height and "effective" horizontal width of the structure associated with each piece of equipment were used in the modeling for downwash calculations.

It was also noted from the photographs that distinct types of rig engines and air compressors are used for the drilling operations, with one of the types having "rain-capped" stacks. This configuration could further retard the momentum plume rise out of the stack. Thus, for conservatism, this particular source was modeled using the "capped" stack option in AERMOD with the recommended low value for exit velocity. Revised industry information indicates that these "rain caps" open during engine operations and the supplemental modeling has incorporated this information. Furthermore, since the off-site "centralized" compressor could conceivably be located adjacent to one of the multi-well pads, this source was located adjacent to, but on the other side of the edge of the 150m by 150m pad site.

The placement of the various sources of criteria pollutants in the modeling is represented in Figure 6.11. The figure shows individual completion equipment engines as modeled in the supplemental analysis. This configuration was deemed adequate for the determination of expected worst-case impacts from a 'typical' multi-well pad site. Although the figure outlines the boundary of the 150m by 150m typical well pad area, it is again clarified that receptors were placed such that each source would have close-in receptors beyond the 10m "buffer" distance determined necessary from a practical standpoint. That is, receptors were placed in the pad area to assure simulation of any configuration of these sources on the pad at a given site.

Annual impacts were initially calculated using the maximum hourly emission rates, and the results reviewed to determine if any thresholds were exceeded. If impacts exceeded the annual threshold for a given pollutant, the "average" emission rates specifically for the drilling engines and air compressors in Table 7 and for the hydraulic fracturing and flaring operations from Table 20 of the Industry Information Report were used. For the other sources, such as the line-heater and offsite compressor, the average and maximum rates are the same as presented in Tables 18 and 24, respectively, and were not modified for the refined annual impacts. As these average rates account only for the variability of "source demand" for the specific duration of the individual operations, an additional adjustment needed to be made for the number of days in a year during which up to 10 such well operations would occur. Thus, from Tables 7 and 14, it is seen that there would be a maximum of 250 days of operations for the drilling engines, maximum of 20 days for hydraulic fracturing engines, and maximum of 30 days of flaring in a given year. Thus, for these sources, the annual average rate was adjusted accordingly. Although initial modeling included 10 wells per pad per year as an assumption, the resultant impacts were reviewed and relevant conclusions adjusted in the sections to follow where it was deemed of consequence to NAAQS or threshold compliance. That is, if the standards compliance was already demonstrated with the worst-case assumption of 10 wells, no revisions were necessary. On the other hand, the modeling has not included any operational limits on the use of the line heater and off-site compressor for the production phase and the annual emissions were represented by the maximum rates. Some of these considerations are further discussed in Section 6.5.2.4.

Lastly, in order to account for the possibility of well operations at nearby pads at the same time as operations at the modeled well pad configuration, a sensitivity analysis was performed to determine the potential contribution of an adjacent pad to the modeled impacts. This assessment addressed, in a simplified manner, the issue of the potential for cumulative effects from a nearby pad on the total concentrations of the modeled pad such that larger "background levels" for the determination of compliance with ambient threshold needed to be determined. The nearby pad with identical equipment and emissions as the pad modeled was located at a distance of one kilometer (km) from the 150m by 150m area of the modeled pad. This separation distance is the minimum expected for horizontal wells drilled from a single pad, which extends out to a rectangular area of 2500m by 1000m (one square mile).

Non-Criteria Pollutant Sources - There are a set of pollutants from two "distinct" sources in the Marcellus Shale operations for which there are no national ambient standards, but for which New York State has established either a state standard (H<sub>2</sub>S) or toxic guideline concentrations. These are VOC species and HAPs which are emitted from: a) sources associated with venting of gas prior to the production phase; or b) as by-products of combustion of gas or fuel oil. A review of the data on these pollutants and their sources indicated that the two distinct source types can be modeled independently, as described below.

First, of the sources which vent the constituents of the "wet" gas (if it is encountered), the flowback venting has by far the most dominant emissions of the toxic constituents. The other two sources of gas venting are the mud-gas separator and the dehydrator, and a comparison of the relative emissions of the five pollutants identified in the Industry Information Report (benzene, hexane, toluene, xylene, and H<sub>2</sub>S) from these three sources in Tables 8, 21 and 22 shows that the flowback venting has about two orders of magnitude higher emissions than the other two sources. As noted in the Industry Information Report, this venting is limited to a few hours before the flare is used, which reduces these emissions by over 90%. Thus, modeling was used to determine the short-term impacts of the venting emissions. Annual impacts were not modeled, due to the very limited time frame for gas venting, even if ten wells are to be drilled at a pad.

It was determined that during these venting events, essentially no other emissions of the same five toxics would occur from other sources. That is, even though a subset of these pollutants are also tabulated in the Industry Information Report at relatively low emissions for the engines, compressors and the flares, it is either not possible or highly unlikely that the latter sources would be operating simultaneously with the venting sources (e.g. gas is either vented or flared from the same stack). Thus, for the short-term venting scenario, only the impacts from the three sources need to be considered. It was also determined that rather than modeling each of the five pollutant for the set of the venting sources for each of the 12 meteorological years, the flowback venting source parameters of Table 15 were used with a unitized emission rate of 1 g/s as representative of all three sources. The actual pollutant specific impacts were then scaled with the total emissions from all three sources. This is an appropriate approximation, not only due to the dominance of the flowback vent emissions, but also since the stack height and the calculated plume heights for these sources are very similar. This simplification significantly reduced the number of model runs which would otherwise be necessary, without any real consequence to the identification of the maximum short-term impacts.

The next set of non-criteria pollutants modeled included those resulting from the combustion sources. It should be clarified that pollutants emitted from the glycol dehydrator (e.g. benzene), which are associated with combustion sources were also included in these model calculations for both the short-term and annual impacts. A review of the emissions in Tables 8, 18, 21, and 24 indicates seven toxic pollutants with no clear dominance of a particular source category. Furthermore, the sources associated with these pollutants have much more variability in the source heights than for the venting scenario. For example, the flare emissions of the three pollutants in Table 21 are higher than for the corresponding hydraulic fracturing pump engines, but the plume from the flame is calculated to be at a much higher elevation than those for the engines or compressors such that a "representative" source could not be simply determined in order to be able to model a unitized emission rate and limit the number of model runs.

However, it was still possible to reduce the number of model calculations from another standpoint. The seven pollutants associated with these sources were ranked according to the ratios of their emissions to the corresponding 1-hour SGCs and AGCs (SGCs for hexane and propylene were determined by Toxics Assessment section since these are not in DAR-1 tables).

These ratios allowed the use of any clearly dominant pollutants which could be used as surrogates to identify either a potential issue or compliance for the whole set of toxics. These calculations indicated that benzene and formaldehyde are clearly the two pollutants which would provide the desired level of scrutiny of all of the rest of the pollutants in the set. To demonstrate the appropriateness of this step, limited additional modeling for the annual impacts for acetaldehyde was also performed due to the relatively low AGC for this pollutant. These steps further reduced the number of model runs by a significant number.

The emission parameters, downwash structure dimension and the location of the sources were the same as for the criteria pollutant modeling. Similar to the case of the criteria pollutants, any necessary adjustments to the annual emission rates to provide more realistic annual impacts were made after the results of the initial modeling were reviewed to determine the potential for adverse impacts. These considerations are further discussed in the resultant impact section.

Pollutant Averaging Times, Ambient Thresholds and Background Levels

The AERMOD model calculates impacts for each of the hours in the meteorological data base at each receptor and then averages these values for each averaging time associated with the ambient standards and thresholds for the pollutants. For example, particulate matter (PM10 and PM2.5) has both 24-hour and annual standards, so the model would present the maximum impact at each receptor for these averaging times. As the form of the standards cannot be exceeded at any receptor around the source, the model also calculates and identifies the overall maximum impacts over the whole set of receptors.

For the set of pollutants initially modeled, the averaging times of the standards are: for SO<sub>2</sub>- 3-hour, 24-hour, and annual; for PM10/PM2.5 - 24-hour and annual; for NO<sub>2</sub> - annual; for CO - 1-hour and 8-hour; and for the set of toxic pollutants – 1-hour SGCs and annual AGCs. For most criteria pollutants, the annual standards are defined as the maxima not to be exceeded at any receptor, while the short-term standards are defined at the highest-second-highest (HSH) level wherein one exceedance is allowed per receptor. The exception is PM2.5 where the standards are defined as the 3 year averages, with the 24-hour calculated at the 98<sup>th</sup> percentile level. The toxic pollutant SGCs and AGCs are defined at a level not be exceeded. In the Department's assessments, the maximum impacts for all averaging times were used for all pollutants, except

for PM2.5, in keeping with modeling guidance for cases where less than five years of meteorological data per site is used.

In addition to the standards, EPA has defined levels which new sources or modifications after a certain time frame cannot exceed and cause significant deterioration in air quality in areas where the observations indicate that the standards are being met (known as attainment areas). The area depicted in Figure 6.4 for the Marcellus Shale has been classified as attainment for all of the pollutants modeled in the Department's analysis. Details on area designations and the state's obligation to bring a nonattainment area into compliance are available at the Department's public webpage as well as from EPA's webpage. 353 For the attainment areas, EPA's Prevention of Significant Deterioration (PSD) regulations define increments for SO<sub>2</sub>, NO<sub>2</sub> and PM10. More recently, EPA finalized the PSD increments for PM2.5; these are discussed below. Although, in the main, the PSD regulations apply only to major sources, the increments are consumed by both major and minor sources and would be modeled to assure compliance. However, the PSD regulations also exempt "temporary" sources from having to analyze for these increments. It is judged that essentially all of the emissions at the well pad can be qualified as temporary sources since the expectation is that the maximum number of wells at a pad can be drilled and completed well within a year. Even if a partial set of the wells is drilled in a year and these operations cease, the increment would be "expanded" as allowed by the regulations.

The only exception to the temporary designation would be the offsite compressor and the line heater which can operate for years. Thus, only these two sources were considered in the increment consumption analysis. The applicable standards and PSD increments are presented in Table 6.14 for the various averaging times. Table 6.14 reflects incorporation of the 1-hour SO<sub>2</sub> and NO<sub>2</sub> NAAQS which are addressed in the supplemental modeling section. Furthermore, the final PSD increments for PM2.5, which become effective on December 20, 2011, are added to the Table. In addition to these standards and increments, the table provides EPA's defined set of Significant Impact Levels (SILs) which exist for most of the criteria pollutants. These SILs are at about 2 to 4% of the corresponding standards and are used to determine if a project would

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<sup>353</sup> http://www.dec.ny.gov/chemical/8403.html and http://www.epa.gov/ttn/naaqs/.

<sup>&</sup>lt;sup>354</sup> Prevention of Significant Deterioration for PM2.5, final rule, Federal Register, Vol. 75, No. 202, October 20, 2010.

have a "significant contribution" to either an existing adverse condition or would cause a standards violation. Table 6.14 -also reflects the SILs for PM2.5 as contained in EPA's final PSD rule.

These SILs are also used to determine whether the consideration of background levels, which include the contribution of regional levels and local sources, need to be explicitly addressed or modeled. When the SILs are exceeded, it is necessary to explicitly model nearby major sources in order to establish potential "hot spots" of exceedances to which the project might contribute significantly. For the present analysis, if the SILs are exceeded for the single multi-well pad, the Department has considered the potential for the contribution of nearby pads to the impacts of the former on a simplified level. The approach used was noted previously and involves the modeling of a nearby pad placed at 1000m distance from the pad for which detailed impacts were calculated, in order to determine the relative contribution of the nearby pad sources. If these results indicate the potential for significant cumulative effects, then further analysis would need to be performed.

On the other hand, in order to determine existing criteria pollutant regional background levels, which would be explicitly included in the calculation of total concentrations for comparison to the standards, the Department has conservatively used the maximum observations from a set of Department monitoring sites in the Marcellus Shale region depicted in Figure 6.4. The location of these sites and the corresponding data is available in the Department's public webpage. The Department has reviewed the data from these sites to determine representative, but worst case background levels for each pollutant. The Department has used maximum values over a three year period from the latest readily available tabulated information from 2005 through 2007 from at least two sites per pollutant within the Marcellus Shale area, with two exceptions. First, in choosing these sites, the Department did not use "urban" locations, which could be overly conservative of the general areas of well drilling. This meant that for NO<sub>2</sub> and CO, data from Amherst and Loudonville, respectively, were used as representative of rural areas since the rest of the Department's monitor sites were all in urban areas for these two pollutants. Second, data for PM10 for the period chosen was not available from any of the appropriate sites due to

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<sup>355</sup> http://www.dec.nv.gov/chemical/8406.html.

switching of these sites to PM2.5 monitoring per EPA requirements. Thus, the Department relied on data from 2002-04 from Newburgh and Belleayre monitors. The final set of data used for background purposes are presented in Table 6.14. These data represent worst case estimates of existing conditions to which the multi-well pad impacts would be added in order to determine total concentrations for comparison to the AAQS. In instances where the use of these maxima causes an exceedance of the AAQS, EPA and Department guidance identify procedures to define more case specific background levels. Per the Department's Air Guide-1, since there are no monitoredbackground levels for the non-criteria pollutants modeled, the impacts of H<sub>2</sub>S and rest of the toxic chemicals are treated as incremental source impacts relative to the corresponding standard and SGCs/AGCs, respectively. Determinations on the acceptability of these incremental impacts are then made in accord with the procedures in Air Guide-1.

The background levels for criteria pollutants relied upon in the initial modeling analysis are still deemed conservative based on a review of observed monitoring levels in more recent years for pollutants such as PM2.5. Thus, most do not need to be updated. On the other hand, for PM2.5 24-hour averages and the new 1-hour NO<sub>2</sub> and SO<sub>2</sub> standards, more refined background levels were determined as discussed in the supplemental modeling section.

#### 6.5.2.4 Results of the Modeling Analysis

Using the various model input data described previously, a number of model calculations were performed for the criteria and toxic pollutants resulting from the distinct operations of the onsite and offsite sources. Each of the meteorological data years were used in these assessments and the receptors grids were defined such as to identify the maxima from the different sources. In some instances, it was possible to limit the number of years of data used in the modeling, as results from a subset indicated impacts well below any thresholds. In other cases, it was necessary to expand the receptor grid such that the decrease in concentration with downwind distance could be determined. These two aspects are described below in the specific cases in which they were used.

As described in the previous section, initial modeling of annual impacts was performed in the same model runs as for the short-term impacts, using the maximum emission rates. However, in a number of cases, this approach lead to exceedances of annual thresholds and, thus, more

appropriate annual emissions were determined in accord with the procedures described in Section 6.5.2.3, and the annual impacts were remodeled for all of the data years. These instances are also described below in the specific cases in which the annual emissions were used. The results from these model runs were then summarized in terms of maxima and compared to the corresponding SILs, PSD increments, ambient standards, and Air Guide-1 AGCs/SGCs.

This comparison indicated that, using the emissions and stack parameter information provided in the Industry Information Report, a few of the ambient thresholds could be exceeded. Certain of these exceedances were associated with conditions (such as very low stacks and downwash effects) which could be rectified relatively easily. Thus, some additional model runs were performed to determine conditions under which the ambient thresholds would be met. These results are presented below with the understanding that industry could implement these or propose their own measures in order to mitigate the exceedances. Results for the criteria pollutants are discussed first, followed by the results for the toxic/non-criteria pollutants.

### Criteria Pollutant Impacts

The set of sources identified in Table 6.12 for short-term simultaneous operations of the various combustion sources with criteria pollutant emissions were initially modeled with the maximum hourly emission rate and one year of meteorological data. It was clear from these results that the annual impacts for PM and NO<sub>2</sub> had to be recalculated using the more appropriate annual emissions procedures discussed in Section 6.5.2.3. That is, for these pollutants, the "average" rates in the Industry Information Report were scaled by the number of days/hours of operations per year for the drilling engine/compressor, the hydraulic fracturing engines and the flare, and then these results were multiplied by ten to account for the potential of ten wells being drilled at a pad for a year. The rest of the sources were modeled assuming full year operations at the maximum rates. In addition, based in part on the initial modeling, two further adjustments were made to the annual NO<sub>2</sub> impacts. First, the model resultant impacts were multiplied by the 0.75 default factor of the Tier 2 screening approach in EPA's modeling guidelines. This factor accounts for the fact that a large part of emissions of NO<sub>x</sub> from combustion sources are not in the NO<sub>2</sub> form of the standard. The second adjustment related to the stack height of the off-site compressor, which was raised to 7.6m (25ft) based on the results for the non-criteria pollutants

discussed below; that is, this height was deemed necessary in order to meet the formaldehyde AGC.

Each of the meteorological data years was used to determine the maximum impacts for all of the criteria pollutants and the corresponding averaging times of the standards. However, in the case of 24-hour particulate impacts, modeling was limited to the initial year (Albany, 2007) for reasons discussed below. The results for each year modeled are presented in Table 6.15. It should be noted that the SO<sub>2</sub> annual impacts in this table are based on the maximum hourly rates and are very conservative. In addition, the tabulated values for the 24-hour PM2.5 impacts are the eight highest in a year, which is used as a surrogate for the three year average of the eight highest value (i.e., 99<sup>th</sup> percentile form of the standard). It is seen that the short-term impacts do not show any significant variability over the twelve years modeled.

The overall maxima for each pollutant and averaging time from Table 6.15 are then transferred to Table 6.17 for comparison to the set of ambient thresholds. These maximum impacts are to be added to the worst case background levels from Table 6.14 (repeated in Table 6.16), with the sum presented in the total concentration column. The impacts of only the compressor and the line heater are also presented separately in Table 6.16 for comparison to the corresponding PSD increments. It should be noted that, due to the low impacts for many of the pollutants from all of the sources relative to the increments, only the 24-hour PM10 and PM2.5 and the annual NO<sub>2</sub> were re-calculated for the compressor and line heater, as noted in Table 6.16. In addition, due to the promulgated PSD increments for PM2.5 in the 10/20/10 final rule, the increments are reflected in Table 6.16, along with the corresponding PM2.5 impacts (conservatively assuming to equal PM10 impacts). The rest of the impacts are the same as those in the maximum overall impact column.

The results indicate that all of the ambient standards and PSD increments would be met by the multiple well drilling activities at a single pad, with the exception of the 24-hour PM10 and PM2.5 impacts. In fact, the 3-hour (and very likely the annual) SO<sub>2</sub> impacts are below the corresponding significant impact levels. This is a direct result of the use of the ultra low sulfur fuel assumed for the engines, which would have to be implemented in these operations. In addition, the level of compliance with standards for the maximum annual impacts for NO<sub>2</sub> and

PM2.5 are such as to require the implementation of the minimum 7.6 m (30feet) stack height for the compressor and general adherence to the annual operational restrictions identified in the Industry Information Report.

Table 6.16 results for 24-hour PM10 and PM2.5 impacts were limited to one year of meteorological data since these were found to be significantly above the corresponding standards, as indicated in Table 6.16. Unlike other cases, a simple adjustment to the stack height did not resolve these exceedances and it was determined that specific mitigation measures would need to be identified by industry. However, the Department determined one simple set of modeling conditions under which impacts can be resolved. It was noted that the relatively large PM10/PM2.5 impacts occurred very close to the hydraulic fracturing engines (and at lower levels near the rig engines) at a distance of 20 m, but there was also a very sharp drop-off of these concentration with distance away from these sources. Specifically, to meet the standards minus the background levels in Table 6.16, it was determined that the receptor distance had to be beyond 80 m for PM10, and 500 m for PM2.5. In an attempt to determine if a stack height adjustment in combination with a distance limitation for public access approach can also alleviate the exceedances, the rig engine and fracturing engine stacks heights were both extended by 3.1m (10ft). From the photographs of the truck-mounted engines, it was not clear if any extensions would be practical and, thus, only this minimal increase was considered. This scenario was modeled again with the Albany 2007 meteorological data. The resultant maximum impacts were reduced to 171 and 104 µg/m3 for PM10 and PM2.5, respectively. For this case, in order to achieve the standards using Table 6.16 background levels, the receptors would be beyond 40 m and 500 m for PM10 and PM2.5, respectively. Thus, the stack height extension did not significantly affect the concentrations at the farther distances, as would be expected from the fact that building downwash effects are largest near the source. However, the background level for PM2.5 can be adjusted from the standpoint that the expected averages associated with these operations at relatively remote areas are better represented by the regional component due to transport. If the contribution of the latter to the observed maxima is conservatively assumed to be half of the value in Table 6.17 (i.e., 15 µg/m<sup>3</sup>), then the receptor distance at which a demonstration of compliance can be made is approximately 150 m.

Thus, one practical measure to alleviate the PM10 and PM2.5 standard exceedances is to raise the stacks on the rig and hydraulic fracturing engines and/or erect a fence at a distance surrounding the pad area in order to preclude public access. Without further modifications to the industry stack heights, a fence out to 500 m would be required, but this distance could be reduced to 150 m with the taller stacks and a redefinition of the background levels. Alternately, there is likely control equipment which could significantly reduce particulate emissions. The set of specific control or mitigation measures would need to be addressed by industry.

Based on recent industry and public information, supplemental modeling analysis and detailed review of potential control measures and their practical use was undertaken. The preliminary results clearly indicate that certain levels of emission reductions are likely necessary for at least the completion equipment engines. The results of the supplemental modeling and the consequent recommended mitigation measure are presented in the two sections which follow.

An additional issue addressed in a simplified manner was the possibility of simultaneous operations at a nearby pad, which could be located at a minimum distance of one km from the one modeled, as described previously. It is highly unlikely than more than one additional pad would be operating as modeled simultaneously with other pads within this distance; it is more likely that drill rigs and other heavy equipment would be moved from one pad to another within a given vicinity, with sequenced operations. Regardless, the impacts of all the pollutants and averaging times were determined at a distance of 500 m from the modeled well pad for the years corresponding to the maximum impacts. This is half the distance to the nearest possible pad and allows the determination of potential "overlap" in impacts from the two pads. The concentrations at 500m drop off sharply from the maxima to below significance levels for almost all cases such that nearby pad emissions would not significantly contribute to the impacts from the modeled source. These impacts at 500m are presented in the last row of Table 6.16 and their comparisons to the corresponding SILs in Table 6.16 show only the 24-hour PM2.5 and annual NO<sub>2</sub> impacts are still significant at this distance.

Thus, there is a potential that for these two cases the nearby pad operations could contribute to another well operation's impacts. This scenario was assessed by placing an identical set of sources at another pad at a distance of 1km from the one modeled in the general upwind

direction from the latter. Impacts were then recalculated on the same receptor grid using the years of modeled worst case impacts for these two pollutants and averaging times. The results indicated that the maximum impacts presented in Table 6.16 for annual  $NO_2$  and 24-hour PM2.5 were essentially the same; in fact the 24-hour PM2.5 impacts are identical to the previous maxima while the  $NO_2$  annual impact of 63.2 increased by only  $1.2 \,\mu\text{g/m}^3$ . Annual impacts from any other pad not in the predominant wind direction would be lower. These results are judged not to effect the compliance demonstrations discussed above. Thus, it is concluded that minimal interactions from nearby pad well drilling operations would result, even if there were to be such simultaneous operations.

In addition to these results, the modeled impacts discussed in the supplemental modeling section and the remediation measures recommended to resolve modeled exceedances of both the 24-hour PM2.5 and 1-hour NO<sub>2</sub> NAAQS would substantially reduce both the PM2.5 and NO<sub>2</sub> impacts from the levels in Table 6.15 at the 500 m distance. Therefore, compliance with standards and increments can be said to be adequately demonstrated on the basis of individual pad results.

## Non-Criteria Pollutant Impacts

As discussed in Section 6.5.2.3, three "distinct" source types were independently modeled for a corresponding set of toxic pollutants: i) short-term venting of gas constituents, ii) combustion byproducts, plus the emissions of the same pollutants from the glycol dehydrator, and iii) a set of representative chemicals from the flowback impoundments. These impacts were determined for comparison to both the short-term 1-hour SGC and annual AGC, with the exception of the venting scenario which was limited to the short-term impacts due to the very short time frame of the practice. The gas venting emissions out of three sources (mud-gas separator, flowback venting, and the dehydrator) are essentially determined by the flowback phase. It was thus possible to model only this source with a unitized emission rate (1g/s) and then actual 1-hour impacts were scaled using the total maximum emission rates.

Each year of meteorological data was modeled with the flowback vent parameters to determine the maximum 1-hour impacts for 1 g/s emission rate. These results were then reviewed and the maximum overall normalized impact of 641  $\mu$ g/m³ (for Albany, 2008 data) was calculated as the worst case hourly impact. Using the total emissions from all three sources for

each of the vented toxic pollutants, as presented in Table 6.17, along with this maximum normalized impact, results in the maximum 1-hour pollutant specific values in the third column of Table 6.17. The pollutants "shaded out" in the table are not vented from these sources. All of the worst case 1-hour impacts are well below the corresponding SGCs, but the maximum 1-hour impact of  $61.5 \,\mu\text{g/m}^3$  for  $H_2S$  (underlined top entry in the box) is above the New York standard of  $14 \,\mu\text{g/m}^3$ .

Thus, if any "sour" gas is encountered in the Marcellus Shale, there would be a potential of exceedance of the  $H_2S$  standard. The maximum 1-hour impact occurred relatively close to the stack, and, in order to alleviate the exceedance, ambient air receptors would be excluded in all areas within at least 100 m of the stack. Alternately, it is possible to also reduce this impact by using a stack height which is higher than the conservative 3.7 m (12 ft) height provided in the Industry Information Report. Iterative calculations for the year with the maximum normalized impact indicated that a minimum stack height of 9.1 m (3 0 ft) would be necessary to reduce the impact to the  $12.1 \,\mu\text{g/m}^3$  value for  $H_2S$  reported in the "Max 1-hour" column of Table 6.18. With this requirement, all venting source impacts would be below the corresponding SGCs and standard.

For the set of seven pollutants resulting from the combustion sources and the dehydrator, it was previously discussed that it was only necessary to explicitly model benzene and formaldehyde, along with the annual acetaldehyde impacts, in order to demonstrate compliance with all SGCs and AGCs for the rest of the pollutants. The relative levels of the SGCs and AGCs presented in Table 6.18 for these pollutants and the corresponding emissions in the Industry Information Report tables clearly show the adequacy of this assertion. For the modeling of these pollutants, the maximum short-term emissions were used for the 1-hour impacts, but the annual emissions were used for the AGCs comparisons. The annual emissions were determined using the same procedures as discussed above for the criteria pollutants.

An initial year of meteorological data which corresponded to the worst case conditions for the criteria pollutants was used to determine the level of these impacts relative to the SGCs and AGCs before additional calculations were made. The results of this initial model run are presented in right-hand set of columns of <u>Table 6.18</u>. These indicate that, while the 1-hour

impacts are an order of magnitude below the benzene and formaldehyde SGCs and the acetaldehyde AGC, there were exceedances of the AGCs for the former two pollutants (the top underlined entries for each pollutant in the maximum annual column). It was determined that these exceedances were each associated with a particular source: the glycol dehydrator for benzene and the offsite compressor for formaldehyde. It should be noted that these exceedances occur even when the emissions from dehydrator are controlled to be below the National Emissions Standard for Hazardous Air Pollutants (NESHAP) imposed emission rate provided in Table 22 of the Industry Information Report and with 90% reduction in formaldehyde emissions accounted for by the installation of an oxidation catalyst, by NESHAP Subpart JJJJ requirement for the compressor. To assure the large margin of safety in meeting the benzene and formaldehyde SGCs and the acetaldehyde AGC, another meteorological data base was used to calculate these impacts. The results in Table 6.17 did not change from these calculations. Thus, it was determined that no further modeling was necessary for these. On the other hand, for the benzene and formaldehyde AGC exceedances, a few additional model runs were performed to test potential mitigating measures. It is clear that, similar to the criteria pollutant impacts, these high annual impacts are partially due to the low stacks and the associated downwash effects for both the dehydrator and the compressor sources. Given that these two sources already need to include NESHAP control measures, the necessary additional reduction in impacts can be practically achieved by either limiting public access to about 150m from these sources, or by raising their stacks.

An iterative modeling of increased stack heights for both the dehydrator and the compressor demonstrated that in order to achieve the corresponding AGCs, the stack of the dehydrator should be a minimum of 9.1m (30ft), in which case it would also avoid building downwash effects, while the compressor stack would be raised to 7.6m (25ft). These higher stacks were then modeled using each of the 12 years of meteorological data and the resultant overall maxima, tabulated in the bottom half of the "Max annual" column in Table 6.18. It should be noted that these modifications to stack height would also reduce the corresponding 1-hour maxima leading to a larger margin of compliance with SGCs. With these stack modifications and the required NESHAP control measures, all of the SGCs and AGCs are projected to be met by the various combustion operations and the dehydrator. It should be noted that appropriate stack height for

both the compressors and any associated dehydrators can be better determined by case-specific modeling during the compressor station permitting process if the dehydrator is to be located at the compressor station.

6.5.2.5 Supplemental Modeling Assessment for Short Term PM2.5, SO<sub>2</sub> and NO<sub>2</sub> Impacts and Mitigation Measures Necessary to Meet NAAQS.

As a supplement to the initial modeling, a number of additional model runs had to be made in order to address certain outstanding issues with PM10 and PM2.5 short term impacts from the original analysis, as well as to incorporate new information provided by industry. In addition, the re-assessment also addresses EPA's promulgated 1-hour NAAQS for SO<sub>2</sub> and NO<sub>2</sub> which became effective since September 2009. The modeling performed previously for PM10/PM2.5 was limited to a simplified set-up of the drilling and completion equipment engines and conservative set of assumptions which lead to substantial exceedances of the 24-hour NAAQS for both PM10 and PM2.5. Based on this preliminary result, it was deemed that further modeling would not resolve the exceedances without some level of emission mitigation.

Thus, industry was asked to provide a set of potential mitigation measures to alleviate these exceedances. In addition, the 2009 draft SGEIS identified a simple stack height and/or "fencing-in" of impacts option to be considered. This latter was not meant as the Department's suggested preferred mitigation option. Instead, the purpose behind the modeling with increased stack height was to provide a quantification of the level of simple physical adjustments to the operations in order for industry to incorporate the results in their assessment of mitigation and control measures. Based on both industry and public input, additional modeling analysis has been undertaken to address the PM10 and PM2.5 exceedances and the associated mitigation measures necessary to assume NAAQS compliance.

In addition to the PM10/PM2.5 issue, EPA promulgated new 1-hour standards for  $SO_2$  and  $NO_2$ . These standards are 100 ppb (or 188  $\mu g/m^3$ ) for  $NO_2$ , as the 3 year average of the 98<sup>th</sup> percentile of the daily maximum 1-hour values and 75 ppb (or 196  $\mu g/m^3$ ) for  $SO_2$ , as the 3 year average of the 99<sup>th</sup> percentile of the daily maximum 1-hour values, which became effective on April 12, 2010 and August 23, 2010, respectively<sup>356</sup>. These standards would be considered within the

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<sup>&</sup>lt;sup>356</sup> Federal Register: Vol 75, No. 26, pp 6474+ (2/9/10) and Vol. 75, No. 119, pp35520+ (6/22/10).

context of this SGEIS and in accordance with Subpart 200.6 requirement defined in Section 6.5.1 to assure all potential adverse impacts are identified and rectified. The additional assessments performed for these short term impacts are addressed separately to distinguish certain information for PM10/PM2.5 gathered from industry since the initial modeling analysis in the SGEIS.

## A) PM 10 and PM2.5 24-hour Impact Modeling and Potential Mitigation Measures.

As part of the Industry's Responses (dated September 16, 2009) to Information Requests, IOGA referenced a modeling assessment performed by consultants for Chesapeake Energy which incorporated a number of revisions to and recommendations on the Department's modeling analysis <sup>357</sup>. The analysis was based on one year of Binghamton meteorological data which indicated compliance with the PM10 NAAQS and much lower PM2.5 impacts than the Department's results, but still exceedances of the PM2.5 NAAQS. Mitigation measures were listed for resolving the latter exceedances. The analysis incorporated a set of assumptions which are summarized below with the Department's position on each of these:

The PM emissions provided by ALL consultants in the Industry Information Report were not speciated with respect to PM10 and PM2.5. Based on factors in EPA's AP-42 for large uncontrolled diesel engines, the PM10 and PM2.5 emissions represent 82% and 69%, respectively, of the total PM emissions. The Department has reviewed the information and agrees that the corresponding emissions should be adjusted accordingly;

The set of 15 completion equipment engines were represented in the Department's modeling as three sets of 5 units stationed next to each other. Industry noted that since these units contributed significantly to the modeled exceedances, each of the engines should be model as a separate point source. The Department had noted this conservative step and has remodeled the units are 15 separate sources. However, unlike Chesapeake's approach of separating the 15 units in two sets at the extreme ends of the pads, the Department has no reason to believe the engines would not be placed next to each other. Thus, the engines are re-modeled as depicted in revised <u>Figure 6.11</u>;

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<sup>&</sup>lt;sup>357</sup> June 21, 2010 letter from Brad Gill of IOGA-NY to Kathleen Sanford and associated modeling files.

It is claimed that the use of ULSF would result in an additional 10% reduction in PM emissions. The Department could not readily verify the level of reduction specifically for all diesel fuel sulfur contents, but it has been considered in our discussion of resultant impacts;

It was notes that the maximum emissions provided for the completion equipment engines are only representative of two hours in the operation cycle of these units. Thus, the hourly emission rate in the modeling was "prorated" to better characterize the likely 24-hour emission rate. The Department does not agree with this approach. As noted in our previous analysis, the ALL report noted a typical hydraulic fracturing operation can require up to 10 stages of total 5 hour periods. Thus, it is likely that a relevant portion of a day could experience the maximum hourly emission rate associated with worst case impacts, as we had previously assumed. Since there is no justified or simplified approach to account for this possibility, we believe it prudent to use the maximum hourly emission rate for the revised analysis; and

It was noted that for drilling engines, the use of the EPA "capping" stack option is not appropriate since the cap is "open" when the engines are in operation. This assumption has been revised in the reassessment by using the actual stack velocities and temperatures.

Finally, the Chesapeake modeling report noted that the background levels used were the maxima observed at representative monitors and are unreasonably high. The SGEIS recognizes the conservative nature of the background levels chosen as worst case observations, but notes that more representative values can be determined in instances where such refinement is necessary. For PM2.5, the reassessment has taken a less conservative approach in accord with the Department's and EPA's modeling guidance by reviewing the monitoring data and the expected associated average values in the Marcellus Shale area. In its March 23, 2010 guidance memo<sup>358</sup> on PM2.5, EPA provided a screening first Tier conservative approach to addressing NAAQS compliance which was to be followed by further guidance with more refined methods.

Lacking the follow-up guidance, most states, including New York, have allowed methods more in line with Section 8.2 of EPA's Modeling Guidelines. One such approach recognized by the March 23, 2010 memo is to allow for seasonal average observed concentrations. In reviewing

<sup>&</sup>lt;sup>358</sup> Modeling Procedures for Demonstrating Compliance with PM2.5 NAAQS, Stephen Page, 3/23/10.

the data at monitors in the Marcellus Shale area, especially for the latest three years, we have identified a value of  $15 \,\mu\text{g/m}^3$  as appropriate for the purpose of determining representative 24-hour "regional" background level. The data also indicates that more recent observations than the 2005-7 levels in the SGEIS have in general shown a downward trend. It is also noted that the modeled impacts would dominate the total impacts which are to be compared to the NAAQS. For this reason, it is deemed appropriate to use the  $8^{th}$  highest concentration, as the form of the NAAQS, instead of the maximum 24-hour value recommended as a first screening Tier. A conservative step was to use the  $8^{th}$  highest maximum from each year of meteorological data modeled since these were limited to only two years per site.

In addition to these modifications to the original PM10 and PM2.5 modeling in the SGEIS, we have incorporated industry's assertion that there would not be simultaneous drilling and hydraulic fracturing operations at a single well pad. In order to better characterize the contribution of the completion equipment engines, the drilling rig engine and the air compressors, in addition to calculating the maximum overall impacts, the modeling results were also separated for each operation to determine the need for mitigation associated with each engine type. The modeling approach was otherwise identical to the previous analysis, except the version of AERMOD was updated to the version (09292) available at the time of the analysis.

The first step in the modeling exercise was to determine the maximum 24-hour PM10 and PM2.5 impact for each of the modeled years. These results are presented in Table 6.19. It is seen that the refined impacts which incorporate the above considerations are much lower than the values in Table 6.15. This reduction is due mainly to the speciated emission rates and the modeling of completion equipment engines as individual point sources. However, the impacts are still projected to be above the PM10 and PM2.5 NAAQS, except for the PM10 impacts associated with the drilling engines. As was noted previously, these maximum impacts occur next to the well pad and concentrations drop-off relatively sharply with downwind distance. The modeled impacts were reviewed and indicate that impacts above the NAAQS-minus-background levels value occurred at distances up to a maximum of 60m for completion equipment engines and PM10, while for PM2.5 the corresponding maximum distances were 120 and 150m for the drilling and completion equipment engines, respectively. The levels of the maximum impacts

also indicate that the different sets of engines could be dealt with using different mitigation measures.

As required by Part 617.11(5) (see next section for more details), the Department would pursue mitigation measures which eliminate potential adverse impacts to the maximum extent practicable. The August 26, 2009 industry report, the Industry Information Report and technical information from the public 359 identified a set of such potential measures which have been reviewed with this SEQRA requirement in mind. Certain of these suggestions would unlikely be practically implemented to any extent; for example, the use of electric engines could be very limited due to the remote nature of the drilling sites, while cleaner fuel engines are currently being investigated by engine manufacturers for future use. To the extent these alternative cleaner engines are available, the Department recommends their use. On the other hand, PM control equipment or the use of newer and cleaner engines are two measures recognized by both industry and the public as viable and the Department's review has concluded that these measures are practical. Appendix 18A provides the Department's review of the emission factors for various tiers of engines and potential after-treatment methods. Its conclusions are incorporated in the following discussions.

The discussions are limited to PM2.5 since these are the controlling impacts; that is, any measures to eliminate the PM2.5 exceedances would also assure compliance with the PM10 NAAQS. For the drilling rig and air compressor engines, the results in Table 6.19 were further analyzed to determine the impacts from each. The contribution to the overall maximum impact (Buffalo, 2007) for drilling operations was associated with the rig engines. Furthermore, industry has suggested and operational diagrams confirm that these engines are used close to the center of the well pad where the drilling actually occurs. The modeling results in Table 6.19 indicate that at a distance of 75m (from the center to the edge of the well pad) the drilling engine impacts are 30  $\mu$ g/m³, essentially due to the rig engine, which would still require mitigation when a background level of 15  $\mu$ g/m³ is used. Even if the 10% reduction in PM emissions due to the use of ULSF is achieved, as argued by industry, the resultant impact would still exceed the NAAQS. The rig engine impacts, however, are associated with ALL report's assumed Tier 1

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For example, comments by AKRF consultants on behalf of NRDC, Memorandum from Hillel Hammer, dated December 3, 2009, page 5.

engine emission factor. If the rig engines class was restricted to the use of Tier 2 and higher, then the PM2.5 impacts would be reduced by at least a factor of 2.7 (see Table Two of Appendix 18A, 0.4/0.15) which would result in compliance with the NAAQS regardless of where these engines are located on the well pad.

Industry data in the IOGA-NY information responses indicate that a majority (71%) of engines currently in use are Tier 2 and Tier 3 engines. In addition, a small fraction (3.5%) are uncertified (Tier 0), with "unknown" emissions. It is the Department's conclusion that these latter engines cannot be used for drilling in New York's Marcellus Shale since it has not been demonstrated that these would result in NAAQS compliance. Furthermore, since 25% of the current drilling engines are Tier 1, their use in New York should only take place with certain control measures. The discussions in Appendix 18A conclude that of the two exhaust after-treatment measures, Diesel Oxidation Catalyst (DOC) and Continuously Regenerating Diesel Particulate Filter (CRDPF) or particulate "traps", the latter is by far the more effective method in that it achieves almost three times the emission reduction (i.e., 85% vs 30%). The level of control achieved by the traps is necessary to alleviate all PM2.5 NAAQS exceedances from any Tier 1 drilling engines. Thus, the CRDPF traps should be the after-treatment for Tier 1 drilling engines if these are to be used in New York. This conclusion also applies to the air compressors for which the maximum PM2.5 impact is calculated to be 65 ug/m<sup>3</sup> for Tier 1 emissions. On the other hand, Tier 2 and above drilling rig engines and air compressors demonstrate NAAQS compliance without these controls.

The Department also considered the "mitigation" of the NAAQS exceedances by stack height and distance restriction measures identified previously in the SGEIS. Although the IOGA-NY response also lists the stack height increase on the drilling engines as a potential measure, there is no indication from industry if such measures are practical given the stack configuration of these engines and the height to which these would be extended. In addition, this measure is not in strict accord with the need to mitigate the adverse impacts to the maximum extent practicable. The combination of operating these engines closer to the drilling rig, but more importantly the use of CRDPF traps on Tier 1 engines are deemed the necessary mitigation measures.

Turning next to the completion equipment engines, it seems even less practical to apply the distance and stack height increase restrictions to this class of engines. In fact, industry has previously indicated that stack height increase on these mobile units cannot be practically accomplished. A modeling run indicates that in order to meet the PM2.5 standard under the revised set of assumptions, the stack height would need to be at least doubled. Furthermore, the distance at which impacts are projected to be below the NAAQS-minus-background level was noted previously to be 150m. This is based on the Tier 2 emission factor modeled for these engines as provided by the ALL report. Consequently, the required practical approach to these engines would also require the use of the CRDPF traps as after-treatment on Tier 2 engines. For the maximum 24-hour PM2.5 case of Table 6.19 (Buffalo, 2006), the 202 µg/m<sup>3</sup> impact reduces to 44 µg/m<sup>3</sup> at a distance of 75m from the engines. Again, a 10% reduction in PM emissions due the use of ULSF does not alleviate these exceedances. Furthermore, unlike the smaller drilling engines, the ability of placing the 15 completion equipment engines (typically 14 used in Pennsylvania) near the center of the well pad is questionable. Based on industry's depiction, it is possible to separate these into two sets at either side of the hydraulic fracturing operations to further reduce impacts. In sum, however, the number of Tier 2 completion equipment engines which would require the installation of the particulate traps ranges from at least two thirds to all of the 15 engines per hydraulic fracturing job. For practical purposes, it is recommended that all Tier 2 engines be equipped with the CRDPF traps. Otherwise, each well operation might need to undergo more site specific analysis to demonstrate that a certain configuration or PM trap installation alternative would assure compliance with the 24-hour PM2.5 and PM10 NAAQS. Further details on the practicality of requiring these traps and other after-treatment control measures are discussed in the section following the SO<sub>2</sub> and NO<sub>2</sub> modeling results.

With respect to the Tier 0 and Tier 1 completion equipment engines, these emissions have not been analyzed or modeled, but for the same reasons as for the drilling engines, Tier 0 completion equipment engines should not be used in New York. In addition, based on the scaling of the maximum impact in <u>Table 6.19</u> by the ratio of Tier 1 to Tier 2 emission factors (2.7), it is determined that Tier 1 engines have the potential to cause a modeled exceedance even if equipped with a particulate trap (maximum impact of  $82 \,\mu\text{g/m}^3$  with 85% control). Industry can suggest impact mitigation in addition to the use of PM traps in order to show compliance with

the NAAQS, but lacking such a demonstration, it is the Department's interim conclusion that Tier 1 completion equipment engines should not be used in New York. On the other hand, and as also suggested by industry and the public, newer Tier 4 engines, which would likely be equipped with traps in order to achieve the required emission factors for those engines, can be used as an alternative to the Tier 2 engines with a PM trap.

## **B**) SO<sub>2</sub> and NO<sub>2</sub> 1-hour Impacts and Potential Mitigation Measures.

The 1-hour SO<sub>2</sub> and NO<sub>2</sub> NAAQS were promulgated since September 2009. Permitting and SEQRA actions after the effective date of an NAAQS are addressed by the Department to assure compliance with the NAAQS in accord with standard Department and EPA policy and requirements. EPA Region 2 recommended that the Department consider the new NAAQS in the SGEIS. In accord with the SEQRA process and the Department's Subpart 200.6 requirement, the Department has modeled the 1-hour SO<sub>2</sub> and NO<sub>2</sub> impacts to assure that all NAAQS are met.

With respect to the 1-hour  $SO_2$  standard of  $196 \, \mu g/m^3$ , no detailed modeling was determined necessary. Instead, the results of the previous  $SO_2$  3-hour modeling in Table 6.15 indicated that the use of the ULSF would likely result in 1-hour impacts being below the NAAQS. Thus, the 1-hour maximum CO impact in Table 6.15 was used to scale the corresponding 1-hour maximum  $SO_2$  impacts using the ratio of the fracturing engine  $SO_2$  and CO emissions since these engines were responsible for the overall maxima. The resultant maximum impact is calculated to be 24  $\mu g/m^3$ . Using a representative, yet conservative, maximum 1-hour  $SO_2$  level of  $126 \, \mu g/m^3$  from the Elmira monitor for 2009 gives a total impact of  $150 \, \mu g/m^3$  which is below the corresponding NAAQS of  $196 \, \mu g/m^3$ . Thus, no further modeling was necessary to demonstrate compliance with the 1-hour  $SO_2$  standard.

Simple scaling to demonstrate compliance was not possible for the NO<sub>2</sub> 1-hour impacts due to the very large concentrations projected using the same method. Instead, it was necessary to account for a number of refinements in the modeling based on EPA and Department guidelines. There are at least two main aspects to the NO<sub>2</sub> modeling which need to be addressed in such refinements. These issues have been raised by EPA, industry and regulatory agencies as needing

further guidance. Similar to the PM2.5 guidance, EPA released a memorandum<sup>360</sup> on June 29, 2010 which provides guidance on how to perform a first Tier assessment for the NO<sub>2</sub> NAAQS. More recently, EPA has provided further guidance <sup>361</sup> on particulars in the modeling approach for NO<sub>2</sub> 1-hour NAAQS compliance determinations.

The two main issues which have been raised deal with: 1) the form of the standard, as the 3 year average of the 98% of the daily maximum 1-hour value, which the AERMOD model used for the original modeling and the revised PM2.5 modeling are not set to calculate, and 2) the ratio of NO<sub>2</sub> to NO<sub>x</sub> emissions assumed for stacks from various source types. Of these, the latter is more critical since NO<sub>2</sub> is a small fraction of the NO<sub>x</sub> emissions in essentially all source types and assuming all of the NO<sub>x</sub> emissions are NO<sub>2</sub> is unrealistic. These issues, however, are not insurmountable. For example, there are model post processors offered by consultants which can readily resolve the first issue. At the time of our re-analysis, EPA provided the Department with a "beta" version of AERMOD which performs the correct averages for NO<sub>2</sub>. Some limited preliminary supplemental modeling used that model version, but the Department has recalculated these impacts using the final version of AERMOD (11059) released on 4/8/11 to assure proper calculation of the 8<sup>th</sup> highest 1-hour maximum per day of meteorological data. The results discussed below reflect the use of this version of AERMOD. It should be noted that the revised version of AERMOD does not contain any changes significant enough to affect the PM2.5 analysis.

With respect to the second issue, a number of entities, including EPA and the Department, have gathered information on the NO<sub>2</sub> to NO<sub>x</sub> ratios from various source types which can be incorporated in the modeling. For the specific drilling and completion equipment engines, Department staff has undertaken a review of available information and has made recommendations on this issue. The details of the recommendations are provided in Appendix 18A which are used in the analysis to be discussed shortly. In addition to this ratio, EPA and Department guidance allows the use of two methods to refine NO<sub>2</sub> modeled impacts; the Ozone

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<sup>&</sup>lt;sup>360</sup> Guidance Concerning the Implementation of the 1-hour NO<sub>2</sub> NAAQS for the Prevention of Significant Deterioration Program. Memo from Stephen Page, EPA OAQPS, dated June 29, 2010.

<sup>&</sup>lt;sup>361</sup> Additional Clarifications Regarding Application of Appendix W Modeling Guidance for the 1-hour NO<sub>2</sub> NAAQS. Memo from Tyler Fox, EPA OAQPS, dated March 1, 2011.

Limiting Method (OLM) and the Plume Volume Molar Ratio Method (PVMRM). There is no preference indicated in EPA guidance as to which method might provide more refinement. However, based on limited model evaluation results presented in the March 1, 2011 EPA guidance memorandum, the current analysis has relied upon the OLM method with the appropriate "source group" option (OLMGROUP ALL) noted in the EPA memo.

In addition to the  $NO_2/NO_x$  ratio, hourly  $O_3$  data is necessary for the use of the method. These were taken from available Department observations at monitor sites representative of the meteorological data bases discussed in the original analysis section. Furthermore, for the determination of background 1-hour  $NO_2$  values, we have refined EPA's first Tier screening approach of using the highest observed levels by calculating the average of the readily available  $3^{rd}$ -highest observations from the Department's Amherst and Pinnacle State Park monitors for the year 2009. This calculated value is  $50~\mu\text{g/m}^3$  and is still conservative relative to the form of the  $NO_2$  standard, as well as relative to further refinements allowed by EPA and Department guidance.

Appendix 18A recommends that, for engines for which emissions were calculated by the Industry Information Report and used in the Department's modeling, the  $NO_2$  fraction of  $NO_x$  is 11% without after-treatment. Thus, an initial set of model runs were performed for the completion equipment engines using the two years of Albany data and this ratio of 0.11 in AERMOD. The results indicate that the maximum impacts from the hydraulic fracturing operations with the 0.11 factor (without the OLM approach) were approximately 3500  $\mu$ g/m³ which, although lower than those from the simple scaling of the CO impacts, are still an order of magnitude above the 1-hour standard of 188  $\mu$ g/m³ for the hydraulic fracturing operations. The impact was noted to be above the NAAQS out to a distance of 300 m from the pad. Thus, further refinements were necessary by the AERMOD-OLM approach.

First to consider, however, is that a confounding issue which this initial modeling did not include was the discovery that the NO<sub>2</sub> to NO<sub>x</sub> ratio is increased by the particulate trap from 0.11 to 0.35 due to the generation of NO<sub>2</sub> in order to oxidize and remove the particulates (see Appendix 18A). This would lead to even higher NO<sub>2</sub> impacts. These results clearly indicate that some form of after-treatment exhaust control method is necessary for the completion equipment

engines. The after-treatment methods to reduce  $NO_x$  emissions are discussed in Appendix 18A which indicates that at present the recommended exhaust treatment method in practical use for on-road engines or engines in general is the SCR system. As noted in Appendix 18A, this preferred after-treatment method for  $NO_x$  control would reduce the  $NO_2$  to  $NO_x$  ratio (with the CRDPF traps in place) down to essentially the same value as without the traps (i.e. 0.10). Of course, the SCR system would also substantially reduces the  $NO_x$  emissions by 90%. Therefore, the last step in the modeling of the completion equipment engines was to use the 90% reduction in emissions and the  $NO_2$ /  $NO_x$  ratio of 0.10 with the OLM option. The analysis relied on the Tier 2 emissions provided by the Industry Information Report as the base emissions which were then reduced by 90% by the SCR controls. This level of modeling was deemed the most refinement allowed currently by Department and EPA guidance.

For the drilling engines, an initial modeling was performed first without the SCR controls and the 0.11 NO<sub>2</sub>/NO<sub>x</sub> ratio and the drilling rig Tier 1 emissions provided in the Industry Information Report as representative of the maximum emission case. For the compressors, Tier 2 was provided as the worst case emissions for the modeling of short term impacts. Based on two years of Albany meteorological data, it was found that the rig engines would exceed the NO<sub>2</sub> 1-hour standard by about a factor of two and impacts would be above the NAAQS-minus-background level out to a distance of 150 m. From the modeling for PM2.5, it was found that the Tier 1 rig engines would need to be equipped with a PM trap in order to project compliance with the 24-hour PM2.5 standard. Since the traps were found to increase the NO2/ NO<sub>x</sub> ratio by three fold, it is clear that the Tier 1 rig engine impacts would be substantially above the 1-hour NO<sub>2</sub> NAAQS without reductions in the NO<sub>2</sub> emissions. Thus, it is concluded that any Tier 1 rig engines (and compressors by analogy) would need to be equipped with both a PM trap and SCR for use in New York drilling activities.

Thus, the final set of modeling analysis used the SCR controlled Tier 2 completion equipment engine emissions with a  $NO_2/NO_x$  ratio of 0.10 and Tier 2 drilling rig engines and air compressor engines (both of which do not require PM traps) with the  $NO_2/NO_x$  ratio set to 0.11 as noted previously. As for the completion equipment engines, the  $NO_2$  modeling for the rig engines and compressors was based on more realistic representation of the units as individual units of five separate, but contiguous point sources as a further refinement to represent their configuration.

The emissions for each were scaled from the totals in Table 8 of the 8/26/09 Industry Report and these were placed in a north-south orientation at the same location as in Figure 6.11.

The set of NO<sub>2</sub> modeling with all of the meteorological data sites considered all potential sources as in previous analysis, but also provided the maximum impact for each of the three types of engines in order to determine specific potential necessary mitigation measures. However, initial modeling of the combined "drilling" scenario using two years of Albany data indicated an inconsistence in the total projected impacts in comparison to the results from the rig engines and compressors separately. This raised a potential issue with the "combined" impacts from these two operations which was related to the specifics of the OLM Ozone "distribution" approach. The resolution of this issue for the purposes of determining impacts from the rig engines and compressors and the need for potential mitigation measure was to recommend to place these two types of engines near the rig in the center of the well pad (as in the case of the PM results) and, furthermore, to separate these on either side of the drill rig to minimize combined impacts. A single year model run indicated this minimized combined impacts. From information and diagrams available, it is clear that these engines are in fact placed near the center of the pad when in actual operation.

The results of the 1-hour  $NO_2$  impacts are presented in <u>Table 6.19</u>. As noted in the table, all engine are based on Tier 2 emissions, with the completion equipment engines assume to use SCR controls. The results for each of the meteorological data years, the overall maxima, the impacts at a 75-m distance (from center of pad to boundary), and the distance at which the impacts fall off to the NAAQS-background value of  $138 \mu g/m^3$  are presented for the completion equipment engines, the rig engines and the compressors. It is seen that the overall maxima are above the NAAQS. However, these need to be qualified relative to the other information tabulated in terms of potential mitigation measures necessary. It should be noted that a number of conservative assumptions are related to these impacts. First, it is noted that if the sources are placed in the center of the pad, as recommended, the impacts are much lower and essentially below the 1-hour NAAQS. Furthermore, these impacts should be adjusted downward by 10% since the tiered emission "limits" for Tier 2 and above are at most 90%  $NO_x$  as described in Appendix 18A. In addition, the background level used is conservative in that it represents the average of the third highest observations in the shale area and can be adjusted downwards.

Lastly, the distance to achieve the NAAQS minus background level is seen in the Table to be very close to the edge of the well pad. Using concentration maps for the three engine types indicate a sharp drop off of impacts such that the NAAQS minus background level is reached essentially at the well pad edge with only the 10% downward adjustment to impacts. In total, these considerations result in the NO<sub>2</sub> impacts being below the 1-hour NAAQS with the proper placement of the engines near the center of the well pad and the use of SCR control on the fracturing engines, coupled with Tier 2 or higher engines.

As discussed in Appendix 18A, SCR control is the only currently available NO<sub>x</sub> reduction system for these size engines which has demonstrated the ability to practically achieve the level of reduction necessary (i.e., minimum 90%) to meet the NAAQS. Since the results of the PM2.5 modeling concluded that Tier 0 (uncertified) and Tier 1 completion equipment engines are not recommended for use in New York if CRDPF (particulate traps) are retrofitted to these, the application of SCR to Tier 2 and newer engines were considered. It is the Department's understanding from the manufacturers of these engines that the Tier 4 engines would have to be equipped with PM traps and SCR in order to meet the more stringent emission limits. It should be recalled that without the SCR control, the particulate traps increase the NO<sub>2</sub> to NO<sub>x</sub> ratio by three fold and the corresponding impacts by a similar magnitude. Thus, the SCR system should be installed on all engines in which PM traps are being required for PM2.5 NAAQS compliance purposes. Any alternate system proposed by industry which has a demonstrated ability to achieve the same level of PM and NO<sub>x</sub> reduction and, concurrently, resolve the NO<sub>2</sub> increase by the particulate traps in order to meet the NAAQS would be considered by the Department. At the present time, the Department is not aware of such an alternative system which has a proven record. For the purposes of the SGEIS, the Department has determined that the SCR system is necessary and adequate for this purpose. The next section discusses the practicality of using both the particulate traps and SCRs on completion equipment engines.

A summary of the Department's determination on the EPA Tier engines and the necessary mitigations to achieve the 24-hour PM2.5 and 1-hour NO<sub>2</sub> NAAQS is presented in tabular form in <u>Table 6.20</u>. The first column provides the various EPA tiers for the drilling and completion equipment engines and their time lines as presented in Appendix 18A. The next column presents sample percent of each Tier engines currently in use as provided by industry in the Information

Report. Note that based on the previous discussions, the uncertified (Tier 0) engines would not be allowed to be used in NY for Marcellus Shale activities. The third column provides the ratio of the Tier 1 emission rates for PM and NO<sub>x</sub> to the other tiers, based on the information in Appendix 18A. The last column summarizes the determinations made by the Department on the control requirements necessary to meet the 24-hour PM2.5 (and PM10) and the 1-hour NO<sub>2</sub> ambient standards. As seen from the table, Tier 1 drilling engines and air compressors would require a PM trap and SCR controls, with the same controls being required on most of the completion equipment engine tiers.

Another purpose of this table is to provide an important demonstration that the Department's recommendations on control measure for these engines would result in substantial emission reduction over the current levels allowed in any other operations in other states. That is, in terms of air quality impacts, the emission reduction factor column of <u>Table 6.20</u> indicates at least a factor of 3 and 2 reductions in PM2.5 and NO<sub>2</sub> emissions, respectively, from the Tier 1 engines. Thus, although Tier 2 and 3 drilling engines make up a majority of the engines in current use (71%), their relative emissions are much lower than the Tier 1 engines, which are recommended not to be used in NY (or have PM traps and SCR controls with about 90% reductions in emissions). Therefore, in terms of emissions reductions, the Department's requirements on the drilling engines would reduce emissions by at least half. Furthermore, since the completion equipment engines are about four times larger than the drilling engines, the imposition of PM traps and SCR on most completion equipment engines means a substantial reduction in overall PM and NO<sub>x</sub> emissions from the set of engines to be used in New York. Any alternative emission reduction schemes which industry might further pursue would be judged against these reductions. It is clear however, that the Department would assure that any such control or mitigation measure would explicitly demonstrate compliance with the ambient air quality standards.

6.5.2.6 The Practicality of Mitigation Measures on the Completion Equipment and Drilling Engines.

The supplemental modeling assessment has concluded that in order to meet the ambient standards for the 24-hr PM2.5 and the 1-hour NO<sub>2</sub> NAAQS, it is necessary that the completion equipment engines tiers allowed to be used in New York to be equipped with particulate filter

traps (CRDPF) and SCR control for NO<sub>x</sub>. These are Tier 2 and newer completion equipment engines. Similarly, the Tier 1 rig engines and air compressors would be required to be equipped with both control devices if these are used in New York. The determination on the specific after-treatment controls was based on the review of available control methods used in practice (see Appendix 18A). Currently available alternative control measures considered were deemed inadequate for the purpose of achieving the level of PM2.5 and NO<sub>x</sub> emission reductions necessary to demonstrate NAAQS compliance and/or having a proven record of use in practice.

Although industry can attempt to perform an independent assessment of alternatives to the recommended exhaust after-treatment controls, it is highly likely that a certain level of control equipment recommended would be necessary on these engines. If industry identifies viable alternative control measure which can be demonstrated to achieve the same level of emission reduction for NAAQS standard compliance, these alternative schemes would need to be submitted for Department review and concurrence prior to their use in New York. Furthermore, in recommending the use of particulate traps and the SCR technology, Department staff has considered the requirements of subsection 617.11.5 and the practicality of the chosen measures.

Taking the diesel particulate traps and the SCR controls separately, it is fair to say that since the former have a longer established history of actual use than the latter on types of engines of size in the rig engine class, the demonstration of practicality for the traps might be less onerous. For example, industry itself has identified these diesel particulate traps on Tier 2 and 3 engines in their list of mitigation measure. In addition, public information (see footnote 17) also has identified the ongoing use of diesel traps as a required mitigation measure by Metropolitan Transportation Authority (MTA) for non-road engines in major construction projects in NYC. These latter engines, however, are in the size range of the smaller rig engines and not in the completion equipment engine range. Information on the ongoing practical use of particulate traps in these and similar activities have been further confirmed by Department staff through publically available information. Thus, while it can be concluded that the requirement to use particulate traps on certain EPA tiered engines is in accord with Subsection 200.6 and 617.11 of the Department's requirements, it is nonetheless necessary for industry to further assess the

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<sup>&</sup>lt;sup>362</sup> ALL Consulting 2010. page 43 of the ALL/IOGA September 16, 2010 Information Request Report.

practicality of their use for the completion equipment engine size range. Based on limited conversations with two of the engine manufacturers indicated that the main issue still to be resolved is the details of the engineering necessary to use PM traps as after-treatment equipment. The concern relates to the need for "stand alone" equipment for each of the completion equipment engines which differs from the built-in or add on components being currently used for the smaller on-road or off-road engines. To the Department's knowledge, currently neither PM and NO<sub>2</sub> control measures are being used by the gas drilling industry for other shale activities to any extent. However, it is the Department's assumption that the PM traps can be feasibly used on the Tier 1 drilling engines and compressors and the Tier 1 and 2 completion equipment engines.

For the use of SCR as the Department's preferred control measure to reduce NOx emissions from all of the completion equipment engines allowed to be used in New York, there is less information on similar size engines. As Appendix 18A notes, however, these units are widely used in a package with particulate traps on heavy duty vehicles and there is no operational reason that the same cannot be achieved with the larger completion equipment engines. One way to judge the practicality of using SCR control on these engines is to consider the costs involved. The Department has undertaken a simple approach to this issue by using the analogy to reducing exhaust stream NOx emission and its "cost effectiveness" as a means for major stationary sources to get a "waiver" from the emission control limits set forth in Subpart 227-2 (Reasonably Available Control Technology (RACT) for Oxides of Nitrogen (NO<sub>x</sub>)). That is, if a source can demonstrate that the costs associated with the imposed emission limits are unreasonable, the Department and EPA would consider granting a waiver from meeting these limits.

Details of an analysis of the "cost effectiveness" of the SCR controls for completion equipment engines and the comparable value currently used by the Department for stationary sources is provided in Appendix 18B. It is important to note that the "cost effectiveness" is based on acceptable "engine size scaling-up" method for the completion equipment engines with certain assumptions which might not be representative of the actual cost of installation of SCR after treatment. The calculations in Appendix 18B indicate that the cost of requiring SCR on the completion equipment engines is within the value used by the Department for stationary sources

and is deemed reasonable. The cost effectiveness for the smaller drilling engines should be lower. It is recognized that the applicability of 227.2 RACT requirements are meant for major individual stationary sources, but it is also to be noted that the potential annual NO<sub>x</sub> emissions from the sum total of engine use throughout the Marcellus Shale are rather large, as discussed in the next section. Based on the conversations with the engine manufacturers, the main concern with the installation of SCR as an after-treatment control relates again to the need for a "standalone" system on the completion equipment engines, with the added complexity that these systems would require "continuous" maintenance to achieve the level of reduction assumed in the Department's analysis. In addition, these discussions indicate that the cost associated with the installation of the PM traps and SCR are likely above those assumed by the Department. A calculation using the approach in Appendix 18C for PM after-treatment indicates that the "cost effectiveness" value is well above the value used for NO<sub>x</sub> RACT waiver determinations. Thus, it is recommended that industry undertake a detailed assessment of the PM traps and SCR controls in addressing the Department's recommendations of these controls as the required mitigation measures on certain Tier drilling and completion equipment engines in order to demonstrate compliance with the 24-hour PM2.5 and 1-hour NO<sub>2</sub> NAAQS.

Based on the above discussions, the Department believes that the use of particulate traps and SCR controls are reasonable and practical in achieving the mitigation of potential adverse 24-hour PM2.5 and 1-hour NO<sub>2</sub> impacts, respectively. As noted previously, industry can present equivalent control measures and background information for further Department considerations. Regardless of the specific measure, however, it should be made clear that the Department is required to assure compliance with ambient standards with respect to any other control measures which could put forth by industry or the public. One of the mitigation "measures" noted by industry in their Information Report, at least for NO<sub>x</sub> emissions, is to allow for the "natural" fleet turnover of the EPA tiers as these requirements would "kick-in" over time. This suggestion is not an acceptable scheme, given that none of the engines currently in use or contemplated are the interim Tier 4 engines, which become effective in 2011, based on the Department's knowledge and industry data. If industry is to advance such a mitigation scheme, it would submit an acceptable timeline which clearly sets out an aggressive schedule to implement the Tier 4 engines. Based on engine manufacturer's information, there is ongoing efforts to achieve the

Tier 4 emission standards before the 2014/15 timelines noted in <u>Table 6.20</u>. Such an implementation schedule can be tied to the specific tiered engine after-treatment controls required by the Department.

## 6.5.2.7 Conclusions from the Modeling Analysis

An air quality impact analysis was undertaken of various sources of air pollution emissions from a multi-horizontal well pad and an example compressor station located next to a typical site in the area underlain by the Marcellus Shale. The analysis relied on recommended EPA and Department modeling procedures and input data assumptions. Due to the extensive area underlain by the Marcellus Shale and other low-permeability gas reservoirs in New York, certain assumptions and simplifications had to be made in order to properly simulate the impacts from a "typical" site such that the results would be generally applicable. At the same time, an adequate meteorological data base from a number of locations was used to assure proper representation of the potential well sites in the area underlain by the Marcellus Shale in New York.

Information pertaining to onsite and offsite combustion and gas venting sources and the corresponding emissions and stack parameters were initially provided by industry and independently verified by Department staff. The emission information was provided for the gas drilling, completion and production phases of expected operations. On the other hand, emissions of potential additive chemicals from the flowback water impoundments, which were proposed by industry as one means for reuse of water, were not provided by industry or an ICF report to NYSERDA. Thus, worst-case emission rates were developed by the Department using an EPA emission model for a set of representative chemicals which were determined to likely control the potential worst case impacts, using information provided by the hydraulic fracturing completion operators. The information included the compounds used for various purposes in the hydraulic fracturing process and the relative content of the various chemicals by percent weight. The resultant calculated emission rates were shared with industry for their input and comment prior to the modeling.

The modeling analysis of all sources was carried out for the short-term and annual averages of the ambient air quality standards for criteria pollutants and for Department defined threshold levels for non-criteria pollutants. The initial modeling used limitations on simultaneous operations of the various equipment at both onsite and offsite operations for a multi-well pad in the analysis for the short-term averages, while the annual impacts accounted for the potential use of equipment at the well pad over one year period for the purpose of drilling up to a maximum of ten wells. For the modeling of chemicals in the flowback water, two impoundments of expected worst case size were used based on information from industry: a smaller on-site and a larger offsite (or centralized) impoundment.

Initial modeling results indicated compliance with the majority of ambient thresholds, but also identified certain pollutants which were projected to be exceeded due to specific sources emission rates and stack parameters provided in the Industry Information Report. It was noted that many of these exceedances related to the very short stacks and associated structure downwash effects for the engines and compressors used in the various phases of operations. Thus, limited additional modeling was undertaken to determine whether simple adjustments to the stack height might alleviate the exceedances as one mitigation measure which could be implemented. An estimate of the distances at which the impacts would reduce to below all applicable SGCs and SGCs were provided as part of the original analysis.

Based on recent information provided by industry on the operational restrictions at the well pad, the elimination of the flowback impoundments, and a limited modeling of 24-hour PM2.5 impacts, the initial Department assessment was revisited. In addition, due to the promulgation of new 1-hour SO<sub>2</sub> and NO<sub>2</sub> NAAQS after September 2009, further modeling was performed. The significant consequences of the revised restrictions on simultaneous operations of the drilling and completion equipment engines, the number of wells to be drilled per year, and the elimination of the impoundments are incorporated in the initial modeling assessment. Further modeling details for the short term PM2.5, NO<sub>2</sub> and SO<sub>2</sub> impacts are presented in a supplemental modeling section. These results indicate the need for the imposition of certain control measures to achieve the NO<sub>2</sub> and PM2.5 NAAQS. These measures, along with all other restrictions reflecting industry's proposals and based on the modeling results, are detailed in Section 6.5.5 as well permit operation conditions.

Table 6.12 - Sources and Pollutants Modeled for Short-Term Simultaneous Operations

| Pollutant                        | SO2         | NO2         | PM10 & PM2.5 | СО          | Non-criteria combustion | H <sub>2</sub> S and other gas constituents |
|----------------------------------|-------------|-------------|--------------|-------------|-------------------------|---|
| Source                           |             |             | 2 2/2        |             | emissions               | Sms community                               |
| Engines for drilling             | >           | <b>&gt;</b> | <b>✓</b>     | <b>&gt;</b> |                         |   |
| Compressors for drilling         | >           | >           | <b>/</b>     | >           | ~                       |   |
| Engines for hydraulic fracturing | <b>&gt;</b> | <b>/</b>    | <b>✓</b>     | <b>&gt;</b> | <b>~</b>                |   |
| Line heaters                     | <b>V</b>    | ~           | ~            | •           | ~                       |   |
| Off-site compressors             | <b>V</b>    | ~           | ~            | ~           | ~                       |   |
| Flowback gas flaring             | <b>V</b>    | <b>V</b>    | ~            | <b>/</b>    | ~                       |   |
| Gas venting                      |             |             |              |             |                         | <b>'</b>                                    |
| Mud-gas separator                |             |             |              |             |                         | ~   |
| Glycol dehydrator                |             |             |              |             | <b>V</b>                | ~   |

Table 6.13 - National Weather Service Data Sites Used in the Modeling

| NWS Data Site | Meteorology Data Years | Latitude/Longitude Coordinates |
|---------------|------------------------|--------------------------------|
| Albany        | 2007-08                | 42.747/73.799                  |
| Syracuse      | 2007-08                | 43.111/76.104                  |
| Binghamton    | 2007-08                | 42.207/75.980                  |
| Jamestown     | 2001-02                | 42.153/79.254                  |
| Buffalo       | 2006-07                | 42.940/78.736                  |
| Montgomery    | 2005-06                | 41.509/74.266                  |

Table 6.14 - National Ambient Air Quality Standards (NAAQS), PSD Increments & Significant Impact Levels (SILs) for Criteria Pollutants ( $\mu g/m^3$ )

| Pollutant             | 1-hour | 3-hour | 8-hour | 24-hour | Annual |
|-----------------------|--------|--------|--------|---------|--------|
| SO <sub>2</sub> NAAQS | 196    | 1300   |        | 365     | 80     |
| PSD Increment         |        | 512    |        | 91      | 20     |
| SILs                  |        | 25     |        | 5       | 1      |
| PM10 NAAQS            |        |        |        | 150     | 50     |
| PSD Increment         |        |        |        | 30      | 17     |
| SILs                  |        |        |        | 5       | 1      |
| PM2.5 NAAQS           |        |        |        | 35      | 15     |
| PSD Increment         |        |        |        | 9       | 4      |
| SILs <sup>363</sup>   |        |        |        | 1.2     | 0.3    |
| NO2 NAAQS             | 188    |        |        |         | 100    |
| PSD Increment         |        |        |        |         | 25     |
| SILs                  |        |        |        |         | 1.0    |
| CO NAAQS              | 40,000 |        | 10,000 |         |        |
| SILs                  | 2000   |        | 500    |         |        |

<sup>&</sup>lt;sup>363</sup> The PM2.5 standards reflect the 3 year averages with the 24 hour standard being calculated as the 98<sup>th</sup> percentile value.

Table 6.15 - Maximum Background Concentration from Department Monitor Sites

| Pollutant | Monitor Sites                     | Maximum Observed Values<br>for 2005-2007 (μg/m³)     |
|-----------|-----------------------------------|--|
| $SO_2$    | Elmira* and Belleayre             | 3 hour - 125 24-hour - 37<br>Annual - 8              |
| $NO_2$    | Amherst                           | Annual - 26  |
| PM10**    | Newburgh* and Belleayre           | 24-hour - 49 Annual - 13                             |
| PM2.5     | Newburgh* and Pinnacle State Park | 24-hour - 30 Annual - 11 (3 year averages per NAAQS) |
| СО        | Loudonville                       | 1-hour - 1714 8 hour - 1112                          |

<sup>Denotes the site with the higher numbers.
For PM10, data from years 2002-4 was used.</sup> 

Table 6.16 - Maximum Impacts of Criteria Pollutants for Each Meteorological Data Set

| Meteorological D | ata Year |        | $SO_2$  |               | P              | M10    | PM2     | 2.5*   | CO     | )      | $NO_2$        |
|------------------|----------|--------|---------|---------------|----------------|--------|---------|--------|--------|--------|---------------|
| & Location       | 1        | 3-hour | 24-hour | <u>Annual</u> | <u>24-hour</u> | Annual | 24-hour | Annual | 1-hour | 8-hour | <u>Annual</u> |
| Albany           | 2007     | 15.4   | 13.3    | 3.1           | 459            | 2.7    | 355     | 2.7    | 9270   | 8209   | 57.9          |
|                  | 2008     | 15.3   | 13.2    | 2.9           |                | 2.4    |         | 2.4    | 9262   | 8298   | 51.0          |
| Syracuse         | 2007     | 15.9   | 12.6    | 2.8           |                | 2.7    |         | 2.7    | 8631   | 7849   | 57.1          |
|                  | 2008     | 15.8   | 14.3    | 2.7           |                | 2.7    |         | 2.7    | 8626   | 7774   | 55.4          |
| Binghamton       | 2007     | 18.5   | 13.4    | 2.3           |                | 2.1    |         | 2.1    | 10122  | 8751   | 45.5          |
|                  | 2008     | 18.6   | 15.4    | 1.9           |                | 1.8    |         | 1.8    | 9970   | 8758   | 37.6          |
| Jamestown        | 2001     | 16.7   | 14.0    | 2.4           |                | 2.1    |         | 2.1    | 8874   | 8193   | 46.4          |
|                  | 2002     | 16.8   | 14.4    | 2.7           |                | 2.3    |         | 2.3    | 8765   | 8199   | 50.9          |
| Buffalo          | 2006     | 16.6   | 15.7    | 3.2           |                | 2.9    |         | 2.9    | 9023   | 8067   | 63.2          |
|                  | 2007     | 16.9   | 14.4    | 3.1           |                | 2.8    |         | 2.8    | 8910   | 8270   | 60.8          |
| Montgomery       | 2005     | 17.4   | 11.6    | 1.9           |                | 1.8    |         | 1.8    | 9362   | 8226   | 38.4          |
|                  | 2006     | 14.4   | 14.0    | 2.2           |                | 2.0    |         | 2.0    | 9529   | 8301   | 41.9          |
| Maximum          |          | 18.6   | 15.7    | 3.2           |                | 2.9    |         | 2.9    | 10122  | 8758   | 63.2          |
| Impact at 500m   |          | 0.3    | 0.3     | 0.05          | 7.1            | .11    | 5.0     | .11    | 480    | 253    | 2.5           |

Note: 24-hour PM2.5 values are the 8<sup>th</sup> highest impact per the standard.

Table 6.17 - Maximum Project Impacts of Criteria Pollutants and Comparison to SILs, PSD Increments and Ambient Standards

| Pollutant and Averaging Time | Maximum Impact (μg/m³) | SIL* | Worst Case Background Level (µg/m³) | Total<br>(μg/m³) | NAAQS<br>(μg/m³) | Increment Impact** (µg/m³) | PSD* Increment $(\mu g/m^3)$ |
|------------------------------|------------------------|------|-------------------------------------|------------------|------------------|----------------------------|------------------------------|
| SO2 - 3 hour                 | 18.6                   | 25   | 125                                 | 143.6            | 1300             | 18.6                       | 512                          |
| SO <sub>2</sub> - 24-hour    | 15.7                   | 5    | 37                                  | 52.7             | 365              | 15.7                       | 91                           |
| SO <sub>2</sub> - Annual     | 3.2                    | 1    | 8                                   | 11.2             | 80               | 3.2                        | 20                           |
| PM10 - 24-hour               | 459***                 | 5    | 49                                  | 508***           | 150              | 6.5**                      | 30                           |
| PM10 - Annual                | 2.9                    | 1    | 13                                  | 15.9             | 50               | 2.9                        | 17                           |
| PM2.5 - 24-hour              | 355***                 | 1.2  | 30***                               | 385***           | 35               | 6.5**                      | 9                            |
| PM2.5 - Annual               | 2.9                    | 0.3  | 11                                  | 13.9             | 15               | 2.9                        | 4                            |
| NO2 - Annual                 | 63.2                   | 1.0  | 26                                  | 89.2             | 100              | 5.6**                      | 25                           |
| CO - 1-hour                  | 10,122                 | 2000 | 1714                                | 11,836           | 40,000           | NA                         | None                         |
| CO - 8 hour                  | 8758                   | 500  | 1112                                | 9870             | 10,000           | NA                         | None                         |

<sup>\*</sup> SILs and increments for PM2.5 included in revised Table from EPA's final PSD rule for PM2.5

<sup>\*\*</sup> Impacts from the off-site compressor plus the line heater only for PSD increment comparisons were recalculated for annual NO<sub>2</sub> and PM10 and PM2.5 24-hour cases. NA means not applicable

<sup>\*\*\*</sup> See Supplemental Modeling Section for revised analysis

Table 6.18 - Maximum Impacts of Non-Criteria Pollutants and Comparisons to SGC/AGC and New York State AAQS

| Pollutant           | Total Venting Emission Rate (g/s) | Impacts from all Venting Sources (µg/m³)  Max 1-hr SGC |        |      | tor Impac | ources and ts (μg/m³)  GC A | <u>nnual</u> |
|---------------------|-----------------------------------|--|--------|------|-----------|-----------------------------|--------------|
| Benzene***          | 0.218                             | 140  | 1,300  | 13.2 | 1,300     | <u>0.90</u><br>0.10         | 0.13         |
| Xylene              | 0.60                              | 365  | 4,300  | NA** | 4,300     | NA                          | 100          |
| Toluene             | 0.78                              | 500  | 37,000 | NA   | 37,000    | NA                          | 5,000        |
| Hexane              | 9.18                              | 5,888  | 43,000 |      |           |                             |              |
| H <sub>2</sub> S*** | 0.096                             | 61.5<br>12.1   | 14*    |      |           |                             |              |
| Formaldehyde* *     |                                   |  |        | 4.4  | 30        | <u>0.20</u><br>0.04         | 0.06         |
| Acetaldehyde        |                                   |  |        | NA   | 4,500     | 0.06                        | 0.45         |
| Naphthalene         |                                   |  |        | NA   | 7,900     | NA                          | 3.0          |
| Propylene           |                                   |  |        | NA   | 21,000    | NA                          | 3,000        |

Denotes the New York State 1-hour standard for H<sub>2</sub>S

The standard exceedance for H<sub>2</sub>S is eliminated by using a minimum stack height of 9.1m for gas venting

The AGC exceedance for formaldehyde is eliminated by using a compressor stack height of 7.6m

<sup>\*\*</sup> Denotes not analyzed by modeling, but the SGCs and AGCs would be met (see text)

<sup>\*\*\*</sup> AGC exceedance for benzene is eliminated by raising the dehydrator stack to  $9.1 \mathrm{m}$ 

Table 6.19 - Modeling Results for Short Term PM10, PM2.5 and  $NO_2$  (New July 2011)

| Met Data Met             |               | PM10, 24-hr (μg/m <sup>3</sup> ) |          |                         | PM2.5, 24-hr<br>(μg/m <sup>3</sup> ) |                         | NO <sub>2</sub> , 1-hour impact (μg/m <sup>3</sup> ) (see NOTE) |            |  |
|--------------------------|---------------|----------------------------------|----------|-------------------------|--------------------------------------|-------------------------|---|------------|--|
| Location                 | Data<br>Year  | Hydraulic<br>Fracturing          | Drilling | Hydraulic<br>Fracturing | Drilling                             | Hydraulic<br>Fracturing | Rig Engine  | Compressor |  |
| Albony                   | 2007          | 313                              | 76       | 152                     | 36                                   | 198                     | 256   | 216        |  |
| Albany                   | 2008          | 268                              | 84       | 129                     | 40                                   | 198                     | 259   | 230        |  |
| Cymanica                 | 2007          | 224                              | 95       | 144                     | 34                                   | 156                     | 196   | 198        |  |
| Syracuse                 | 2008          | 327                              | 81       | 120                     | 27                                   | 161                     | 180   | 208        |  |
| Dinghomton               | 2007          | 281                              | 87       | 154                     | 34                                   | 194                     | 239   | 208        |  |
| Binghamton               | 2008          | 327                              | 89       | 121                     | 35                                   | 213                     | 231   | 220        |  |
| Lamastarum               | 2001          | 339                              | 74       | 151                     | 29                                   | 180                     | 237   | 221        |  |
| Jamestown                | 2002          | 229                              | 83       | 155                     | 33                                   | 181                     | 248   | 217        |  |
| Buffalo                  | 2006          | 338                              | 106      | 202                     | 55                                   | 147                     | 269   | 231        |  |
| Dullaio                  | 2007          | 318                              | 102      | 189                     | 59                                   | 148                     | 272   | 231        |  |
| Montgomowy               | 2005          | 255                              | 77       | 104                     | 28                                   | 169                     | 198   | 202        |  |
| Montgomery               | 2006          | 301                              | 66       | 108                     | 21                                   | 155                     | 211   | 200        |  |
| Maximum (µ               | ug/m³)        | 339                              | 106      | 202                     | 59                                   | 213                     | 272   | 231        |  |
| Max @ 75m                | $(\mu g/m^3)$ | 92                               | 75       | 44                      | 30                                   | 100-140                 | 140-170   | 120-150    |  |
| Max Dist to N Background | ~             | 60                               | 60       | 150                     | 120                                  | <90                     | <100  | <100       |  |

**NOTE**: NO<sub>2</sub> results reflect SCR controls on the completion equipment engines, with Tier 2 emissions used for all completion equipment, rig engines and compressors. Results are from the OLM option in AERMOD. See text for details.