

Permit ID: 3-3926-00729/00054 Renewal Number: 3 03/28/2022

Facility Identification Data

Name: STRIDES PHARMA INC Address: 1 RAM RIDGE RD CHESTNUT RIDGE, NY 10977

Owner/Firm

Name: STRIDES PHARMA INC

Address: 2 TOWER CTR BLVD STE 1102 EAST BRUNSWICK, NJ 08816, USA

Owner Classification: Corporation/Partnership

Permit Contacts

Division of Environmental Permits:

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Phone:

Permit Description Introduction

The Title V operating air permit is intended to be a document containing only enforceable terms and conditions as well as any additional information, such as the identification of emission units, emission points, emission sources and processes, that makes the terms meaningful. 40 CFR Part 70.7(a)(5) requires that each Title V permit have an accompanying "...statement that sets forth the legal and factual basis for the draft permit conditions". The purpose for this permit review report is to satisfy the above requirement by providing pertinent details regarding the permit/application data and permit conditions in a more easily understandable format. This report will also include background narrative and explanations of regulatory decisions made by the reviewer. It should be emphasized that this permit review report, while based on information contained in the permit, is a separate document and is not itself an enforceable term and condition of the permit.

Summary Description of Proposed Project

Application for renewal of Air Title V Facility. There are no changes to the current ATV permit as part of this renewal.



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Attainment Status

STRIDES PHARMA INC is located in the town of RAMAPO in the county of ROCKLAND. The attainment status for this location is provided below. (Areas classified as attainment are those that meet all ambient air quality standards for a designated criteria air pollutant.)

Criteria Pollutant

Attainment Status

Particulate Matter (PM)	ATTAINMENT
Particulate Matter< 10μ in diameter (PM10)	ATTAINMENT
Sulfur Dioxide (SO2)	ATTAINMENT
Ozone*	SEVERE NON-ATTAINMENT
Oxides of Nitrogen (NOx)**	ATTAINMENT
Carbon Monoxide (CO)	ATTAINMENT

Facility Description:

Strides Pharma Inc is a manufacturer as well as a research and development facility of pharmaceutical products. The facility is located in Chestnut Ridge, Rockland County, New York. Potential emissions of Volatile Organic Compounds exceed major source thresholds subjecting the facility to Title V permitting. The facility emissions of individual and total Hazardous Air Pollutants (HAPs) are limited below major source thresholds. The facility is subject to monitoring and record keeping requirement under 6 NYCRR Parts 201 and 212. The Standard Industrial Code (SIC) is 2834 - Pharmaceutical Preparations.

Permit Structure and Description of Operations

The Title V permit for STRIDES PHARMA INC

is structured in terms of the following hierarchy: facility, emission unit, emission point, emission source and process. A facility is defined as all emission sources located at one or more adjacent or contiguous properties owned or operated by the same person or persons under common control. The facility is subdivided into one or more emission units (EU). Emission units are defined as any part or activity of a stationary facility that emits or has the potential to emit any federal or state regulated air pollutant. An emission unit is represented as a grouping of processes (defined as any activity involving one or more emission sources (ES) that emits or has the potential to emit any federal or state regulated air pollutant). An emission source is defined as any apparatus, contrivance or machine capable of causing emissions of any air contaminant to the outdoor atmosphere, including any appurtenant exhaust system or air cleaning device. [NOTE: Indirect sources of air contamination as defined in 6 NYCRR Part 203 (i.e. parking lots) are excluded from this definition]. The applicant is required to identify the principal piece of equipment (i.e., emission source) that directly results in or controls the emission of federal or state regulated air pollutants from an activity (i.e., process). Emission sources are categorized by the following types:

combustion - devices which burn fuel to generate heat, steam or power

incinerator - devices which burn waste material for disposal

control - emission control devices

process - any device or contrivance which may emit air contaminants

^{*} Ozone is regulated in terms of the emissions of volatile organic compounds (VOC) and/or oxides of nitrogen (NOx) which are ozone precursors.

^{**} NOx has a separate ambient air quality standard in addition to being an ozone precursor.



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that is not included in the above categories.

STRIDES PHARMA INC is defined by the following emission unit(s):

Emission unit 00000T -

Drying ovens No. 6 (DRY06), No. 7 (DRY07), and No. 8 (DRY08) are used to remove isopropyl alcohol (IPA) and/or ethanol (EtOH) from solvent-based pharmaceutical products or water from aqueous based pharmaceutical products. Solvent and aqueous modes are identified as process D03. In solvent mode, the dryers are vented to a dedicated scrubber for the first six hours of the drying cycle as follows: DRY06 vents to Scrubber No. 2 (SCR02) then exhausts via EP00006; DRY07 vents to Scrubber No. 1 (SCR01) then exhausts via EP00005; DRY08 vents to Scrubber No. 3 (SCR03) then exhausts via EP00007. Then the scrubbers will be turned off and emissions from the remaining portion of the drying cycle exhausts through a bypass vent (DRY06 to EP00008, DRY07 to EP00009, DRY08 to EP00010). When operating in aqueous mode, the dryers will exhaust via dedicated bypass vents (DRY06 to EP00008, DRY07 to EP00009, DRY08 to EP00010).

This emission unit also consists of a 48" (COAT4) and 60" (COAT7) Accela-Cota Tablet Coaters, two Glatt GPCG-60 coater/granulation/dryer units (COAT5 and COAT6), and one ACG Fluid Bed Dryer (FBD) (COAT8). Each has a dedicated fume hood (HOOD1, HOOD2, HOOD3, HOOD4, and HOOD5), which is used to dispense raw materials used in the process. Solvent mode operation is identified as one process ID (P07). IPA and/or EtOH are used in the 48" Tablet Coater, two Glatts, FBD #3, and associated fume hoods. Acetone is used in the 48" and 60" Tablet Coaters and associated fume hoods. Methanol (MeOH) is also used in the two Glatts and associated fume hoods. This equipment can be used in aqueous mode (process IDs: P08, P09, P10, P13 & P14).

The 48" Tablet Coater (COAT4) is associated with HOOD1. Particulate emissions from COAT4 are controlled by dust collector (DC-11). COAT4/HOOD1 is controlled by the carbon absorption system (CA1RR) and exhausted via EP00040 when using IPA and EtOH. When acetone is used, emissions are controlled by Condenser (COND1) and exhausted via EP00045. When operating in aqueous mode, COND1 and CA1RR are bypassed and emissions exhaust via bypass vent EP00041.

The 60" Tablet Coater (COAT7) is associated with HOOD4. Particulate emissions from COAT7 are controlled by dust collector (DC-18). When acetone is used, COAT7/HOOD4 vents to Condenser (COND2) and exhaust via EP00046. When operating in aqueous mode, COND2 is bypassed, and emissions exhaust via bypass vent EP00047.

COAT4/HOOD1 and COAT7/HOOD4 will not operate at the same time when processing acetone batchers.

Glatt #1 (COAT5) is associated with HOOD2 and Glatt #2 (COAT6) is associated with HOOD3. Particulate emissions from COAT5 and COAT6 are controlled by internal dust collectors DC-12 and DC-13, respectively. When using solvent in the Glatts and their associated fume hoods, the emissions are either directed to SCR01 and exhausted via EP00005 if IPA and/or EtOH is used or CA1RR and exhausted via EP00040 if MeOH is used. CA1RR can also be used as a backup to SCR01 when using IPA and/or EtOH in COAT5/HOOD2 and COAT6/HOOD3. This equipment can also operate in aqueous mode. When doing so, SCR01 and CA1RR are bypassed and emissions from COAT5/HOOD2 exhaust via bypass vent EP00026; and COAT6/HOOD3 exhaust via bypass vent EP00042.

AGC (COAT8) is associated with HOOD5. Particulates from COAT8 will be controlled by an internal dust collector DC-19. SCR03 will be used to reduce the solvent emissions from COAT8 and HOOD5



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(EP00007). When in aqueous mode, SCR03 is bypassed and emissions from COAT8 and HOOD5 exhaust via bypass vent EP00048.

Emission unit 00000T is associated with the following emission points (EP): 00005, 00006, 00007, 00008, 00009, 00010, 00026, 00040, 00041, 00042, 00045, 00046, 00047, 00048 Process: D03 Drying ovens #283 (DRY06), #284 (DRY07), and #285 (DRY08) are used to remove isopropyl alcohol and/or ethanol from pharmaceutical products. Emissions will exhaust through wet scrubbers as follows: DRY07 to SCR01 (EP00005), DRY06 to SCR02 (EP00006), and DRY08 to SCR03 (EP00007) for the first six hours of the drying cycle. Then scrubbers will be turned off and emissions will exhaust through bypass emission points for the remainder of the drying cycle as follows: DRY06 to EP00008, DRY07 to EP00009, DRY08 to EP00010.

Process: D04 Drying ovens #283 (DRY06), #284 (DRY07), and #285 (DRY08) when operating in aqueous mode. Emissions will exhaust through the following emission points: DRY06 to EP00008, DRY07 to EP00009, DRY08 to EP00010.

Process: P07 Operation of the following equipment in solvent mode utilizing isopropyl alcohol and/or ethanol or methanol:

The 48" Accela-Cota Tablet Coater (COAT4), two Glatt GPCG-60 coater/granulation/dryer units (COAT5 & COAT6), and one ACG Fluid Bed Dryer (FBD) (COAT8), each with a dedicated fume hood used to dispense raw materials (HOOD 1/COAT4, HOOD2/COAT5, HOOD3/COAT6, and HOOD5/COAT8).

Methanol is also used in the two Glatts and associated fume hoods (COAT5/HOOD2 and COAT6/HOOD3).

All equipment is used in pharmaceutical production and/or research & development purposes. All equipment can be used in solvent mode (VOCs). COAT5/HOOD2 and COAT6/HOOD3 can use methanol (HAP) in addition to VOC.

Particulate emissions from COAT4, COAT5, COAT6, and COAT8 are controlled by dust collectors (DC-11, DC-12 [internal], DC-13 [internal], and DC-19 [internal], respectively) before being manifolded to its dedicated fume hood and directed to VOC control.

VOC emissions are controlled by a wet scrubber or the carbon absorption system as follows: COAT4/HOOD1 to CAIRR (EP00040); COAT5/HOOD2 to SCR01 (EP00005) or CA1RR (EP00040) as a backup; COAT6/HOOD3 to SCR01 (EP00005) or CA1RR (EP00040) as a backup, and COAT8/HOOD5 to SCR03 (EP00007).

Methanol emissions are controlled as follows: COAT5/HOOD2 to CA1RR (EP00040) and COAT6/HOOD3 to CA1RR (EP00040).

Process: P08 A Glatt GPCG-60 coater/granulation/dryer units (COAT5) with a dedicated fume hood (HOOD2) used to dispense raw materials operates in aqueous mode.



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Particulate emissions from COAT5 are controlled by an internal dust collector (DC-12) before being manifolded to HOOD2 and vented to atmosphere (EP00026).

Process: P09 is located at Building 01 - A 48" Accela-Cota Tablet Coater (COAT4) with a dedicated fume hood (HOOD1) used to dispense raw materials operates in aqueous mode.

Particulate emissions from COAT4 are controlled by an external dust collector (DC-11) before being manifolded to HOOD1 and vented to atmosphere (EP00041).

Process: P10 is located at Building 01 - A Glatt GPCG-60 coater/granulation/dryer units (COAT6) with a dedicated fume hood (HOOD3) used to dispense raw materials operates in aqueous mode.

Particulate emissions from COAT6 are controlled by an internal dust collector (DC-13) before being manifolded to HOOD3 and vented to atmosphere (EP00042).

Process: P13 A 60" Accela-Cota Tablet Coater (COAT7) with a dedicated fume hood (HOOD4) used to dispense raw materials operates in aqueous mode.

Particulate emissions from COAT7 are controlled by an external dust collector (DC-18) before being manifolded to HOOD4 and vented to atmosphere (EP00047).

Process: P14 is located at Building 01 - An ACG Fluid Bed Dryer (FBD) (COAT8) with a dedicated fume hood (HOOD5) used to dispense raw materials operates in aqueous mode.

Particulate emissions from COAT8 are controlled by an internal dust collector (DC-19) before being manifolded to HOOD5 and vented to atmosphere (EP00048).

Process: P16 Operation of the following equipment when utilizing acetone:

The 48" (COAT4) and 60" (COAT7) Accela-Cota Tablet Coaters, each with a dedicated fume hood used to dispense raw materials (HOOD1/COAT4, and HOOD4/COAT7).

COAT4/HOOD1 & COAT7/HOOD4 will not operate at the same time when processing acetone batches.

Particulate emissions from COAT4 & COAT7 are controlled by dust collectors (DC-11, & DC-18, respectively) before being manifolded to its dedicated fume hood & directed to acetone control.

Acetone emissions are controlled by condensers as follows: COAT4/HOOD1 to COND1(EP00045) & COAT7/HOOD4 to COND2 (EP00046).

Emission unit 00000A - Drying pharmaceutical products using oven No. 4 and oven No. 5. Products are aqueous based only.



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Emission unit 00000A is associated with the following emission points (EP):

00001, 00002 Process: D01 is located at GROUND - Drying of aqueous based pharmaceutical products.

Emission unit 00000B - Drying aqueous based pharmaceutical products using an aeromatic fluidized bed dryer. Emissions are exhausted through emission point 00003.

Emission unit 00000B is associated with the following emission points (EP):

00003

Process: D02 is located at GROUND - Drying aqueous based pharmaceutical products.

Emission unit 00000C - Chemical storage room.

Emission unit 00000C is associated with the following emission points (EP):

00004

Process: S01 Chemical Storage.

Emission unit 00000E - Aqueous coating of pharmaceutical tablets. Particulate emissions are controlled by a dust collector exhausted to emission point 00020.

Emission unit 00000E is associated with the following emission points (EP):

00020

Process: P01 is located at GROUND - Aqueous coating of pharmaceutical tablets.

Emission unit 00000F - Aqueous coating of pharmaceutical tablets. Particulate emissions are controlled by a dust collector exhausted to emission point 00021.

Emission unit 00000F is associated with the following emission points (EP):

00021

Process: P02 is located at GROUND - Aqueous coating of pharmaceutical tablets.

Emission unit 00000G - Aqueous based compu-lab tablet coater. Particulate emissions are controlled by a dust collector exhausted to emission point 00022.

Emission unit 00000G is associated with the following emission points (EP): 00022

Process: P03 is located at GROUND - Aqueous based compu-lab tablet coater.

Emission unit 00000H - This emission unit defines overall facility fugitive volatile organic compound (VOC) emissions associated with sanitizing solvents used and batch production of pharmaceutical



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products. Fugitive VOC emissions are assumed exhausting through emission point 00023.

Emission unit 00000H is associated with the following emission points (EP): 00023

Process: F01 is located at GROUND - Fugitive volatile organic compound (VOC) emissions associated with sanitizing solvents used and batch production of pharmaceutical products.

Emission unit 00000J - Operations associated with the granulation area. Particulate emissions are controlled by a dust collector (0DC05) exhausted to emission point 00025.

Emission unit 00000J is associated with the following emission points (EP): 00025

Process: P05 is located at GROUND - Operations associated with the granulation area.

Emission unit 00000K - Chemical storage building.

Emission unit 00000K is associated with the following emission points (EP): 00031

Process: S02 Chemical storage.

Emission unit 00000L - Operations associated with the storage of hazardous waste.

Emission unit 00000L is associated with the following emission points (EP): 00032

Process: S03 Storage of hazardous wastes.

Emission unit 00000M - This emission unit is comprised of the Compression Area (Rooms 23, 27 and 28) and relocated Pharmacy Dispensing Area (Rooms 1, 2 and 3). Negative pressure is maintained on emission sources consisting of the Pharmacy Dispensing Area room and three Compression Area tablet presses.

Particulate emissions are controlled by a DFT 2-4 PulseJet Cartridge Dust Collector (0DC16) and exhausted through Emission Point 00034.

Emission unit 00000M is associated with the following emission points (EP): 00034

Process: P06 is located at Ground - Combined process consisting of the pharmacy dispensing area and compression area activities.

Emission unit 00000I - Operations associated with the compression area. Particulate emissions are controlled by a dust collector (0DC04) exhausted to emission point 00024.



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Emission unit 00000I is associated with the following emission points (EP): 00024

Process: P04 is located at GROUND - Operations associated with the compression area.

Emission unit 00000N - Packaging lines A, D, E and F. Particulate emissions are controlled by dust collector DC-07 vented to EP00035.

Emission unit 00000N is associated with the following emission points (EP): 00035

Process: PLA is located at Ground, Building 01 - Packaging of pharmaceutical products Lines A, D, E and F in support of pharmaceutical production and or R & D. Dust collector DC-07 is utilized to control particulate emissions exhausted through emission point 35.

Emission unit 000000 - Two Packaging lines B and C and one blister packaging line. Particulate emissions are controlled by dust collector DC-08 vented to EP00036.

Emission unit 00000O is associated with the following emission points (EP): 00036

Process: PLB is located at Building 01 - Packaging of pharmaceutical products Lines B, C and Blister.

Emission unit 00000R - Activities associated with granulation rooms No. 2, No. 4 and No. 6A. Emission sources are a portable Fitzmill, a granulator, a blender and a sifter. Air is collected from each room, routed to separate in line HEPA filters, combined, and vented to a common dust collector to control particulate emissions prior to exiting through emission point 00043.

Emission unit 00000R is associated with the following emission points (EP): 00043

Process: P11 is located at Building 01 - Process associated with granulation.

Emission unit 00000S - Creams and Gels Area consisting of one - Fette 1200i tablet press, one - Woowon mixer and dust pickups in six pharmaceutical manufacturing rooms. Particulate emissions are controlled by a DFT 2-4 PulseJet Cartridge Dust Collector (DC-17).

Emission unit 00000S is associated with the following emission points (EP):

Process: P12 Pharmaceutical production associated with Creams and Gel areas.



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Emission unit 00000U - A dust collector system will pick up particulate matter (PM) from spot areas and/or room ventilation pick up points in five new manufacturing clean rooms (MCR) (rooms 37, 38, 39, 40, & 41) (MCR37, MCR38, MCR39, MCR40, MCR41) plus one new packaging line (0PL-J). The PM collected from this system will be manifolded and directed to a Torit dust collector (DC-20) and exhausted via Emission Point EP00049. Only aqueous products will be produced in the manufacturing clean rooms and pre made pharmaceutical products will be packaged on the packaging line 0PL-J.

Emission unit 00000U is associated with the following emission points (EP): 00049

Process: P15 is located at Ground, Building 01 - A dust collection system will pick up particulate matter (PM) from spot areas and/or room ventilation pick up points in five new manufacturing clean rooms (MCR) (rooms 37, 38, 39, 40 & 41) (MCR37, MCR38, MCR39, MCR40, MCR41) plus one new packaging line (0PL-J). The PM collected from this system will be manifolded and directed to a Torit dust collector (DC-20) and exhausted via Emission Point EP00049. Only aqueous products will be produced in the manufacturing clean rooms and pre made pharmaceutical products will be packaged on the packaging line 0PL-J.

Title V/Major Source Status

STRIDES PHARMA INC is subject to Title V requirements. This determination is based on the following information:

Strides Pharma Inc is subject to Title V requirements. This determination is based on the following information: Source is major since potential to emit (PTE) for volitile organic carbons (VOC) is greater than 25 tons per year.

Program Applicability

The following chart summarizes the applicability of STRIDES PHARMA INC with regards to the principal air pollution regulatory programs:

Regulatory Program

Applicability

PSD	NO
NSR (non-attainment)	NO
NESHAP (40 CFR Part 61)	NO
NESHAP (MACT - 40 CFR Part 63)	NO
NSPS	NO
TITLE IV	NO
TITLE V	YES
TITLE VI	NO
RACT	NO
SIP	YES



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NOTES:

PSD Prevention of Significant Deterioration (40 CFR 52, 6 NYCRR 231-7, 231-8) - requirements which pertain to major stationary sources located in areas which are in attainment of National Ambient Air Quality Standards (NAAQS) for specified pollutants.

NSR New Source Review (6 NYCRR 231-5, 231-6) - requirements which pertain to major stationary sources located in areas which are in non-attainment of National Ambient Air Quality Standards (NAAQS) for specified pollutants.

NESHAP National Emission Standards for Hazardous Air Pollutants (40 CFR 61, 6 NYCRR 200.10) - contaminant and source specific emission standards established prior to the Clean Air Act Amendments of 1990 (CAAA) which were developed for 9 air contaminants (inorganic arsenic, radon, benzene, vinyl chloride, asbestos, mercury, beryllium, radionuclides, and volatile HAP's).

MACT Maximum Achievable Control Technology (40 CFR 63, 6 NYCRR 200.10) - contaminant and source specific emission standards established by the 1990 CAAA. Under Section 112 of the CAAA, the US EPA is required to develop and promulgate emissions standards for new and existing sources. The standards are to be based on the best demonstrated control technology and practices in the regulated industry, otherwise known as MACT. The corresponding regulations apply to specific source types and contaminants.

NSPS New Source Performance Standards (40 CFR 60, 6 NYCRR 200.10) - standards of performance for specific stationary source categories developed by the US EPA under Section 111 of the CAAA. The standards apply only to those stationary sources which have been constructed or modified after the regulations have been proposed by publication in the Federal Register and only to the specific contaminant(s) listed in the regulation.

Title IV Acid Rain Control Program (40 CFR 72 thru 78, 6 NYCRR 201-6) - regulations which mandate the implementation of the acid rain control program for large stationary combustion facilities.

Title VI Stratospheric Ozone Protection (40 CFR 82, Subpart A thru G, 6 NYCRR 200.10) - federal requirements that apply to sources which use a minimum quantity of CFC's (chlorofluorocarbons), HCFC's (hydrofluorocarbons) or other ozone depleting substances or regulated substitute substances in equipment such as air conditioners, refrigeration equipment or motor vehicle air conditioners or appliances.

RACT Reasonably Available Control Technology (6 NYCRR Parts 212-3, 220-1.6, 220-1.7, 220-2.3, 220-2.4, 226, 227-2, 228, 229, 230, 233, 234, 235, 236) - the lowest emission limit that a specific source is capable of meeting by application of control technology that is reasonably available, considering technological and economic feasibility. RACT is a control strategy used to limit emissions of VOC's and NOx for the purpose of attaining the air quality standard for ozone. The term as it is used in the above table refers to those state air pollution control regulations which specifically regulate VOC and NOx emissions.

SIP State Implementation Plan (40 CFR 52, Subpart HH, 6 NYCRR 200.10) - as per the CAAA, all states are empowered and required to devise the specific combination of controls that, when implemented, will bring about attainment of ambient air quality standards established by the federal government and the individual state. This specific combination of measures is referred to as the SIP. The term here refers to those state regulations that are approved to be included in the SIP and thus are considered federally enforceable.



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Compliance Status

Facility is in compliance with all requirements.

SIC Codes

SIC or Standard Industrial Classification code is an industrial code developed by the federal Office of Management and Budget for use, among other things, in the classification of establishments by the type of activity in which they are engaged. Each operating establishment is assigned an industry code on the basis of its primary activity, which is determined by its principal product or group of products produced or distributed, or services rendered. Larger facilities typically have more than one SIC code.

SIC Code Description

2834 PHARMACEUTICAL PREPARATIONS

SCC Codes

SCC or Source Classification Code is a code developed and used" by the USEPA to categorize processes which result in air emissions for the purpose of assessing emission factor information. Each SCC represents a unique process or function within a source category logically associated with a point of air pollution emissions. Any operation that causes air pollution can be represented by one or more SCC's.

SCC Code	Description
3-01-060-09	CHEMICAL MANUFACTURING CHEMICAL MANUFACTURING - PHARMACEUTICAL PREPARATIONS Air Dryers
3-01-060-10	CHEMICAL MANUFACTURING CHEMICAL MANUFACTURING - PHARMACEUTICAL PREPARATIONS
3-01-060-11	Storage/Transfer CHEMICAL MANUFACTURING CHEMICAL MANUFACTURING - PHARMACEUTICAL PREPARATIONS
3-01-060-12	Coating Process CHEMICAL MANUFACTURING CHEMICAL MANUFACTURING - PHARMACEUTICAL PREPARATIONS
3-01-060-22	Granulation Process CHEMICAL MANUFACTURING CHEMICAL MANUFACTURING - PHARMACEUTICAL PREPARATIONS
3-01-060-99	CHEMICAL MFG:PHARMACEUTICAL PRODUCTION:MISCELLANEOUS FUGITIVES CHEMICAL MANUFACTURING
	CHEMICAL MANUFACTURING - PHARMACEUTICAL PREPARATIONS Other Not Classified

Facility Emissions Summary

In the following table, the CAS No. or Chemical Abstract Service code is an identifier assigned to every chemical compound. [NOTE: Certain CAS No.'s contain a 'NY' designation within them. These are not true CAS No.'s but rather an identification which has been developed by the department to identify groups



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of contaminants which ordinary CAS No.'s do not do. As an example, volatile organic compounds or VOC's are identified collectively by the NY CAS No. 0NY998-00-0.] The PTE refers to the Potential to Emit. This is defined as the maximum capacity of a facility or air contaminant source to emit any air contaminant under its physical and operational design. Any physical or operational limitation on the capacity of the facility or air contamination source to emit any air contaminant, including air pollution control equipment and/or restrictions on the hours of operation, or on the type or amount or material combusted, stored, or processed, shall be treated as part of the design only if the limitation is contained in federally enforceable permit conditions. The PTE for each contaminant that is displayed represents the facility-wide PTE in tons per year (tpy) or pounds per year (lbs/yr). In some instances the PTE represents a federally enforceable emissions cap or limitation for that contaminant. The term 'HAP' refers to any of the hazardous air pollutants listed in section 112(b) of the Clean Air Act Amendments of 1990. Total emissions of all hazardous air pollutants are listed under the special NY CAS No. 0NY100-00-0. In addition, each individual hazardous air pollutant is also listed under its own specific CAS No. and is identified in the list below by the (HAP) designation.

Cas No.	Contaminant ACETONITRILE	PTE lbs/yr	PTE tons/yr	Actual lbs/yr	Actual tons/yr
000630-08-0	CARBON MONOXIDE	21526		3908	
000067-66-3	CHLOROFORM			0.31	
000067-64-1	DIMETHYL KETONE	27966		1209	
000067-56-1	METHYL ALCOHOL	19000		16	
011104-93-1	NITROGEN OXIDE- (USE 0NY210-00-0)	41524		4710	
0NY075-00-0	PARTICULATES	4442		496	
007446-09-5	SULFUR DIOXIDE	949		32	
0NY100-00-0	TOTAL HAP	49000		30	
0NY998-00-0	VOC	73948		20429	

NOTIFICATION OF GENERAL PERMITTEE OBLIGATIONS

Item A: Public Access to Recordkeeping for Title V Facilities - 6 NYCRR 201-1.10(b) The Department will make available to the public any permit application, compliance plan, permit, and monitoring and compliance certification report pursuant to Section 503(e) of the Act, except for information entitled to confidential treatment pursuant to 6 NYCRR Part 616 - Public Access to records and Section 114(c) of the Act.

Item B: Timely Application for the Renewal of Title V Permits -6 NYCRR Part 201-6.2(a)(4)

Owners and/or operators of facilities having an issued Title V permit shall submit a complete application at least 180 days, but not more than eighteen months, prior to the date of permit expiration for permit renewal purposes.

Item C: Certification by a Responsible Official - 6 NYCRR Part 201-6.2(d)(12)

Any application, form, report or compliance certification required to be submitted pursuant



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to the federally enforceable portions of this permit shall contain a certification of truth, accuracy and completeness by a responsible official. This certification shall state that based on information and belief formed after reasonable inquiry, the statements and information in the document are true, accurate, and complete.

Item D: Requirement to Comply With All Conditions - 6 NYCRR Part 201-6.4(a)(2)

The permittee must comply with all conditions of the Title V facility permit. Any permit non-compliance constitutes a violation of the Act and is grounds for enforcement action; for permit termination, revocation and reissuance, or modification; or for denial of a permit renewal application.

Item E: Permit Revocation, Modification, Reopening, Reissuance or Termination, and Associated Information Submission Requirements - 6 NYCRR Part 201-6.4(a)(3)

This permit may be modified, revoked, reopened and reissued, or terminated for cause. The filing of a request by the permittee for a permit modification, revocation and reissuance, or termination, or of a notification of planned changes or anticipated noncompliance does not stay any permit condition.

Item F: Cessation or Reduction of Permitted Activity Not a Defense - 6 NYCRR 201-6.4(a)(5)

It shall not be a defense for a permittee in an enforcement action to claim that a cessation or reduction in the permitted activity would have been necessary in order to maintain compliance with the conditions of this permit.

Item G: Property Rights - 6 NYCRR 201-6.4(a)(6)

This permit does not convey any property rights of any sort or any exclusive privilege.

Item H: Severability - 6 NYCRR Part 201-6.4(a)(9)

If any provisions, parts or conditions of this permit are found to be invalid or are the subject of a challenge, the remainder of this permit shall continue to be valid.

Item I: Permit Shield - 6 NYCRR Part 201-6.4(g)

All permittees granted a Title V facility permit shall be covered under the protection of a permit shield, except as provided under 6 NYCRR Subpart 201-6. Compliance with the conditions of the permit shall be deemed compliance with any applicable requirements as of the date of permit issuance, provided that such applicable requirements are included and are specifically identified in the permit, or the Department, in acting on the permit application or revision, determines in writing that other requirements specifically identified are not applicable to the major stationary source, and the permit includes the determination or a concise summary thereof. Nothing herein shall preclude the Department from revising or revoking the permit pursuant to 6 NYCRR Part 621 or from exercising its summary abatement authority. Nothing in this permit shall alter or affect the following:

i. The ability of the Department to seek to bring suit on behalf of the State of New York, or the Administrator to seek to bring suit on behalf of the United States, to immediately restrain any person causing or contributing to pollution presenting an imminent and substantial endangerment to public health, welfare or the environment to stop the emission of air pollutants causing or contributing to



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such pollution;

- ii. The liability of a permittee of the Title V facility for any violation of applicable requirements prior to or at the time of permit issuance;
- iii. The applicable requirements of Title IV of the Act;
- iv. The ability of the Department or the Administrator to obtain information from the permittee concerning the ability to enter, inspect and monitor the facility.

Item J: Reopening for Cause - 6 NYCRR Part 201-6.4(i)

This Title V permit shall be reopened and revised under any of the following circumstances:

- i. If additional applicable requirements under the Act become applicable where this permit's remaining term is three or more years, a reopening shall be completed not later than 18 months after promulgation of the applicable requirement. No such reopening is required if the effective date of the requirement is later than the date on which this permit is due to expire, unless the original permit or any of its terms and conditions has been extended by the Department pursuant to the provisions of Part 2 01-6.7 and Part 621.
- ii. The Department or the Administrator determines that the permit contains a material mistake or that inaccurate statements were made in establishing the emissions standards or other terms or conditions of the permit.
- iii. The Department or the Administrator determines that the Title V permit must be revised or reopened to assure compliance with applicable requirements.
- iv. If the permitted facility is an "affected source" subject to the requirements of Title IV of the Act, and additional requirements (including excess emissions requirements) become applicable. Upon approval by the Administrator, excess emissions offset plans shall be deemed to be incorporated into the permit.

Proceedings to reopen and issue Title V facility permits shall follow the same procedures as apply to initial permit issuance but shall affect only those parts of the permit for which cause to reopen exists.

Reopenings shall not be initiated before a notice of such intent is provided to the facility by the Department at least thirty days in advance of the date that the permit is to be reopened, except that the Department may provide a shorter time period in the case of an emergency.

Item K: Permit Exclusion - ECL 19-0305

The issuance of this permit by the Department and the receipt thereof by the Applicant does not and shall not be construed as barring, diminishing, adjudicating or in any way affecting any legal, administrative or equitable rights or claims, actions, suits, causes of action or demands whatsoever that the Department may have against the Applicant for violations based on facts and circumstances alleged to have occurred or existed prior to the effective date of this permit, including, but not limited to, any enforcement action authorized pursuant to the provisions of applicable federal law, the Environmental Conservation Law of the State of New York (ECL) and Chapter III of the Official Compilation of the Codes, Rules and Regulations of the State of New York (NYCRR). The



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issuance of this permit also shall not in any way affect pending or future enforcement actions under the Clean Air Act brought by the United States or any person.

Item L: Federally Enforceable Requirements - 40 CFR 70.6(b)

All terms and conditions in this permit required by the Act or any applicable requirement, including any provisions designed to limit a facility's potential to emit, are enforceable by the Administrator and citizens under the Act. The Department has, in this permit, specifically designated any terms and conditions that are not required under the Act or under any of its applicable requirements as being enforceable under only state regulations.

NOTIFICATION OF GENERAL PERMITTEE OBLIGATIONS

Item A: Emergency Defense - 6 NYCRR 201-1.5

- An emergency, as defined by subpart 201-2, constitutes an affirmative defense to penalties sought in an enforcement action brought by the Department for noncompliance with emissions limitations or permit conditions for all facilities in New York State.
- (a) The affirmative defense of emergency shall be demonstrated through properly signed, contemporaneous operating logs, or other relevant evidence that:
 - (1) An emergency occurred and that the facility owner or operator can identify the cause(s) of the emergency;
 - (2) The equipment at the permitted facility causing the emergency was at the time being properly operated and maintained;
 - (3) During the period of the emergency the facility owner or operator took all reasonable steps to minimize levels of emissions that exceeded the emission standards, or other requirements in the permit; and
 - (4) The facility owner or operator notified the Department within two working days after the event occurred. This notice must contain a description of the emergency, any steps taken to mitigate emissions, and corrective actions taken.
- (b) In any enforcement proceeding, the facility owner or operator seeking to establish the occurrence of an emergency has the burden of proof.
- (c) This provision is in addition to any emergency or upset provision contained in any applicable requirement. item 02

Item B: General Provisions for State Enforceable Permit Terms and Condition - 6 NYCRR Part 201-5

Any person who owns and/or operates stationary sources shall operate and maintain all emission units and any required emission control devices in compliance with all applicable Parts of this Chapter and existing laws, and shall operate the facility in accordance with all criteria, emission limits, terms, conditions, and standards in this permit. Failure of such



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person to properly operate and maintain the effectiveness of such emission units and emission control devices may be sufficient reason for the Department to revoke or deny a permit.

The owner or operator of the permitted facility must maintain all required records on-site for a period of five years and make them available to representatives of the Department upon request. Department representatives must be granted access to any facility regulated by this Subpart, during normal operating hours, for the purpose of determining compliance with this and any other state and federal air pollution control requirements, regulations or law.

Regulatory Analysis

Location Facility/EU/EP/Proce	Regulation ess/ES	Condition	Short Description
 FACILITY	ECL 19-0301	41	Powers and Duties of the Department with respect to air pollution control
FACILITY	40CFR 68	18	Chemical accident
FACILITY	40CFR 82-F	19	prevention provisions Protection of Stratospheric Ozone - recycling and emissions reduction
FACILITY	6NYCRR 200.6	1	Acceptable ambient air quality.
FACILITY	6NYCRR 200.7	10	Maintenance of
FACILITY	6NYCRR 201-1.4	42	equipment. Unavoidable noncompliance and violations
FACILITY	6NYCRR 201-1.7	11	Recycling and Salvage
FACILITY	6NYCRR 201-1.8	12	Prohibition of reintroduction of collected contaminants to the air
FACILITY	6NYCRR 201-3.2(a)	13, 14	Exempt Activities - Proof of eligibility
FACILITY	6NYCRR 201-3.3(a)	15	Trivial Activities - proof of eligibility
FACILITY	6NYCRR 201-6	20, 21, 22, 23, 38, 39	Title V Permits and the Associated Permit Conditions
FACILITY	6NYCRR 201-6.4(a)(4)	16	General Conditions - Requirement to Provide Information
FACILITY	6NYCRR 201-6.4(a)(7)	2	General Conditions - Fees
FACILITY	6NYCRR 201-6.4(a)(8)	17	General Conditions -
FACILITY	6NYCRR 201-6.4(c)	3	Right to Inspect Recordkeeping and Reporting of Compliance Monitoring
FACILITY	6NYCRR 201-6.4(c)(2)	4	Records of Monitoring, Sampling and Measurement
FACILITY	6NYCRR 201-	5	Reporting



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	6.4(c)(3)(ii		Requirements -
			Deviations and
			Noncompliance
FACILITY	6NYCRR 201-6.4(d)(4)	24	Compliance Schedules
			- Progress Reports
FACILITY	6NYCRR 201-6.4(e)	6	Compliance
			Certification
FACILITY	6NYCRR 201-6.4(f)	25	Operational
			Flexibility
FACILITY	6NYCRR 201-6.5(a)	43	State Enforceable
			Requirements
FACILITY	6NYCRR 201-7	26, 40	Federally Enforceable
			Emissions Caps
FACILITY	6NYCRR 202-1.1	29	Required emissions
			tests.
FACILITY	6NYCRR 202-2.1	7	Emission Statements -
			Applicability
FACILITY	6NYCRR 202-2.5	8	Emission Statements -
			record keeping
			requirements.
FACILITY	6NYCRR 211.1	44	General Prohibitions
			- air pollution
			prohibited
FACILITY	6NYCRR 211.2	30	General Prohibitions
			- visible emissions
			limited.
FACILITY	6NYCRR 212-1.6(a)	31	Limiting of Opacity
FACILITY	6NYCRR 212-1.7(b)(3)	32	The VOC outlet
			concentration from
			fixed-bed carbon
			adsorption units
FACILITY	6NYCRR 212-2.3(b)	45, 46, 47, 48	State Air Program
			Non-Criteria air
			contaminants subject
			Table 4
FACILITY	6NYCRR 212-2.4(b)	33, 34	Control of
			Particulate from New
			and Modified Process
			Emission Sources
FACILITY	6NYCRR 212-	35 , 36	RACT compliance plan
	3.1(c)(4)(i)		control limits for
			Capture and Control
FACILITY	6NYCRR 215.2	9	Open Fires -
			Prohibitions
FACILITY	6NYCRR 225-1.2(d)	37	Sulfur-in-Fuel
			Limitation -
			Distillate Oil

Applicability Discussion:

Mandatory Requirements: The following facility-wide regulations are included in all Title V permits:

ECL 19-0301

This section of the Environmental Conservation Law establishes the powers and duties assigned to the Department with regard to administering the air pollution control program for New York State.

6 NYCRR 200.6

Acceptable ambient air quality - prohibits contravention of ambient air quality standards without mitigating measures

6 NYCRR 200.7

Anyone owning or operating an air contamination source which is equipped with an emission control



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device must operate the control consistent with ordinary and necessary practices, standards and procedures, as per manufacturer's specifications and keep it in a satisfactory state of maintenance and repair so that it operates effectively

6 NYCRR 201-1.4

This regulation specifies the actions and recordkeeping and reporting requirements for any violation of an applicable state enforceable emission standard that results from a necessary scheduled equipment maintenance, start-up, shutdown, malfunction or upset in the event that these are unavoidable.

6 NYCRR 201-1.7

Requires the recycle and salvage of collected air contaminants where practical

6 NYCRR 201-1.8

Prohibits the reintroduction of collected air contaminants to the outside air

6 NYCRR 201-3.2 (a)

An owner and/or operator of an exempt emission source or unit may be required to certify that it operates within the specific criteria described in this Subpart. All required records must be maintained on-site for a period of 5 years and made available to department representatives upon request. In addition, department representatives must be granted access to any facility which contains exempt emission sources or units, during normal operating hours, for the purpose of determining compliance with this and any other state and federal air pollution control requirements, regulations, or law.

6 NYCRR 201-3.3 (a)

The owner and/or operator of a trivial emission source or unit may be required to certify that it operates within the specific criteria described in this Subpart. All required records must be maintained on-site for a period of 5 years and made available to department representatives upon request. In addition, department representatives must be granted access to any facility which contains trivial emission sources or units subject to this Subpart, during normal operating hours, for the purpose of determining compliance with this and any other state and federal air pollution control requirements, regulations, or law.

6 NYCRR Subpart 201-6

This regulation applies to those terms and conditions which are subject to Title V permitting. It establishes the applicability criteria for Title V permits, the information to be included in all Title V permit applications as well as the permit content and terms of permit issuance. This rule also specifies the compliance, monitoring, recordkeeping, reporting, fee, and procedural requirements that need to be met to obtain a Title V permit, modify the permit and demonstrate conformity with applicable requirements as listed in the Title V permit. For permitting purposes, this rule specifies the need to identify and describe all emission units, processes and products in the permit application as well as providing the Department the authority to include this and any other information that it deems necessary to determine the compliance status of the facility.

6 NYCRR 201-6.4 (a) (4)

This mandatory requirement applies to all Title V facilities. It requires the permittee to provide information that the Department may request in writing, within a reasonable time, in order to determine whether cause exists for modifying, revoking and reissuing, or terminating the permit or to determine compliance with the permit. The request may include copies of records required to be kept by the permit.

6 NYCRR 201-6.4 (a) (7)

This is a mandatory condition that requires the owner or operator of a facility subject to Title V requirements to pay all applicable fees associated with the emissions from their facility.



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6 NYCRR 201-6.4 (a) (8)

This is a mandatory condition for all facilities subject to Title V requirements. It allows the Department to inspect the facility to determine compliance with this permit, including copying records, sampling and monitoring, as necessary.

6 NYCRR 201-6.4 (c)

This requirement specifies, in general terms, what information must be contained in any required compliance monitoring records and reports. This includes the date, time and place of any sampling, measurements and analyses; who performed the analyses; analytical techniques and methods used as well as any required QA/QC procedures; results of the analyses; the operating conditions at the time of sampling or measurement and the identification of any permit deviations. All such reports must also be certified by the designated responsible official of the facility.

6 NYCRR 201-6.4 (c) (2)

This requirement specifies that all compliance monitoring and recordkeeping is to be conducted according to the terms and conditions of the permit and follow all QA requirements found in applicable regulations. It also requires monitoring records and supporting information to be retained for at least 5 years from the time of sampling, measurement, report or application. Support information is defined as including all calibration and maintenance records and all original strip-chart recordings for continuous monitoring instrumentation, and copies of all reports required by the permit.

6 NYCRR 201-6.4 (c) (3) (ii)

This regulation specifies any reporting requirements incorporated into the permit must include provisions regarding the notification and reporting of permit deviations and incidences of noncompliance stating the probable cause of such deviations, and any corrective actions or preventive measures taken.

6 NYCRR 201-6.4 (d) (4)

This condition applies to every Title V facility subject to a compliance schedule. It requires that reports, detailing the status of progress on achieving compliance with emission standards, be submitted semiannually.

6 NYCRR 201-6.4 (e)

Sets forth the general requirements for compliance certification content; specifies an annual submittal frequency; and identifies the EPA and appropriate regional office address where the reports are to be sent.

6 NYCRR 202-1.1

This regulation allows the department the discretion to require an emission test for the purpose of determining compliance. Furthermore, the cost of the test, including the preparation of the report are to be borne by the owner/operator of the source.

6 NYCRR 202-2.1

Requires that emission statements shall be submitted on or before April 15th each year for emissions of the previous calENDar year.

6 NYCRR 202-2.5

This rule specifies that each facility required to submit an emission statement must retain a copy of the statement and supporting documentation for at least 5 years and must make the information available to department representatives.

6 NYCRR 211.2



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This regulation limits opacity from sources to less than or equal to 20 percent (six minute average) except for one continuous six-minute period per hour of not more than 57 percent opacity.

6 NYCRR 215.2

Except as allowed by section 215.3 of 6 NYCRR Part 215, no person shall burn, cause, suffer, allow or permit the burning of any materials in an open fire.

40 CFR Part 68

This Part lists the regulated substances and there applicability thresholds and sets the requirements for stationary sources concerning the prevention of accidental releases of these substances.

40 CFR Part 82, Subpart F

Subpart F requires the reduction of emissions of class I and class II refrigerants to the lowest achievable level during the service, maintenance, repair, and disposal of appliances in accordance with section 608 of the Clean Air Act AmENDments of 1990. This subpart applies to any person servicing, maintaining, or repairing appliances except for motor vehicle air conditioners. It also applies to persons disposing of appliances, including motor vehicle air conditioners, refrigerant reclaimers, appliance owners, and manufacturers of appliances and recycling and recovery equipment. Those individuals, operations, or activities affected by this rule, may be required to comply with specified disposal, recycling, or recovery practices, leak repair practices, recordkeeping and/or technician certification requirements.

Facility Specific Requirements

In addition to Title V, STRIDES PHARMA INC has been determined to be subject to the following regulations:

6 NYCRR 201-6.4 (f)

This section describes the potential for certain operational changes to be made by the facility owner or operator without first obtaining a permit modification. Changes made pursuant to this provision must meet all of the criteria described in this section to qualify for consideration as operational flexibility. The Department reserves the right to require the facility owner or operator to obtain a permit modification prior to making any changes at the facility pursuant to this section.

6 NYCRR 201-6.5 (a)

This subdivision states that the Department shall include state enforceable conditions in Title V permits. State enforceable conditions related to regulations developed pursuant to the Climate Leadership and Community Protection Act (CLCPA) and Article 75 of New York State Environmental Conservation Law may be included in future versions of this permit, as applicable.

6 NYCRR 211.1

This regulation requires that no person shall cause or allow emissions of air contaminants to the outdoor atmosphere of such quantity, characteristic or duration which are injurious to human, plant or animal life or to property, or which unreasonably interfere with the comfortable enjoyment of life or property.

6 NYCRR 212-1.6 (a)



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This provisions requires that the facility owner or operator not cause or allow emissions having an average opacity during any six consecutive minutes of 20 percent or greater from any process emission source or emission point, except for the emission of uncombined water.

6 NYCRR 212-1.7 (b) (3)

This paragraph requires the source owner to monitor the VOC outlet concentration of the fixed-bed carbon adsorption unit to demonstrate on-going compliance.

6 NYCRR 212-2.3 (b)

Table 4 of 212-2.3 describes the reduction in emissions required for a non-criteria air contaminant based on its uncontrolled emission rate. The uncontrolled emission rate in conjunction with the assigned environmental rating determines the degree of controlled applied.

6 NYCRR 212-2.4 (b)

Particulate emissions from any process emission source, which received a B or C Environmental Rating, and for which an application was received by the department after July 1, 1973 are restricted to 0.050 grains per cubic foot of exhaust gas, expressed at standard conditions on a dry gas basis.

6 NYCRR 212-3.1 (c) (4) (i)

This provision states that owners and/or operators of emission points subject to Part 212-3 operating prior to October 20, 1994 must submit a compliance plan to the department. The compliance plan must demonstrate that the VOC emission points are equipped with a capture system and a control device with an overall removal efficiency of at least 81 percent.

6 NYCRR 225-1.2 (d)

This subdivision sets the sulfur-in-fuel limitation for distillate oil fired emission sources throughout the State.

6 NYCRR Subpart 201-7

This regulation sets forth an emission cap that cannot be exceeded by the facility. In this permit that cap



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is

Compliance Certification Summary of monitoring activities at STRIDES PHARMA INC:

Location Facility/EU/EP/Process/ES	Cond	No. Type of Monitoring
FACILITY	14	work practice involving specific operations
FACILITY	21	monitoring of process or control device parameters as surrogate
FACILITY	22	monitoring of process or control device parameters as surrogate
FACILITY	23	work practice involving specific operations
FACILITY	5	record keeping/maintenance procedures
FACILITY	6	record keeping/maintenance procedures
FACILITY	27	work practice involving specific operations
FACILITY	28	work practice involving specific operations
FACILITY	7	record keeping/maintenance procedures
FACILITY	31	monitoring of process or control device parameters as surrogate
FACILITY	32	continuous emission monitoring (cem)
FACILITY	45	monitoring of process or control device parameters as surrogate
FACILITY	46	monitoring of process or control device parameters as surrogate
FACILITY	47	intermittent emission testing
FACILITY	48	intermittent emission testing
FACILITY	33	monitoring of process or control device parameters as surrogate
FACILITY	34	intermittent emission testing
FACILITY	35	monitoring of process or control device parameters as surrogate
FACILITY	36	intermittent emission testing
FACILITY	37	work practice involving specific operations

Basis for Monitoring 6 NYCRR Part 201-3

For the emergency generators on site to be considered emergency, the facility must maintain monthly records which demonstrate that each engine is operated less than 500 hours per year, on a 12-month rolling total basis.

6 NYCRR Part 201-6

Strides Pharma shall limit the acetone-based production associated with Process P07, process equipment COAT4 and COAT7, fume hoods HOOD1 and HOOD4, and control equipment COND1 and COND2 to 65 commercial batches per year.



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The Glatt 1 and Glatt 2 solvent based commercial batches limit of 100 per year is based on a historical permit action to avoid New Source Review nonattainment implications.

The facility is required to comply with the maximum number of total batches for COAT8/HOOD5 of 20 commercial batches per 12-month period.

6 NYCRR Part 201-7

Strides Pharma shall demonstrate that its emissions are below the 24.5 tons per year of total HAPs based on a 12-month rolling average by maintaining records of all required measurements and emission calculations of individual HAP and total HAPs from coating, drying, clearing, and miscellaneous activities.

Strides Pharma shall demonstrate that its emissions are below the 9.5 tons per year of any individual HAP, including Methyl Alcohol, based on a 12-month rolling average by maintaining records of all required measurements and emission calculations of individual HAP and total HAPs from coating, drying, cleaning, and miscellaneous activities.

6 NYCRR Part 212-1

Opacity is limited to 20 percent. Formal method 9 observations to verify compliance shall be conducted annually. However, the facility must conduct a monthly visual survey and perform corrective action as needed.

The VOC monitoring requirement that applies to the carbon adsorption system is based on reasonable engineering practice to identify cycles of needed maintenance or repair. The concentration of 150 ppm VOC is an indicator established as part of previous permit actions.

6 NYCRR Part 212-2

Particulate emissions are limited to 0.05 grains per dscf. The facility will monitor, and record pressure drop across the dust collectors manually on a weekly basis, when in operation, to verify compliance. The dust collectors must not exceed a pressure drop of 6 inches of water. If the pressure drop exceeds 6 inches of water, the filters must be changed. The Department reserves the right to require additional testing, if deemed necessary.

The facility is required to maintain a capture system and a control device with an overall removal efficiency of at least 77 percent for the removal on acetone associated with Process 07, process equipment COAT4, HOOD1, Control equipment COND1 and Dust Collector DC-11. The facility will monitor and record the coil temperature manually on a weekly basis, when in operation, to verify compliance. Condenser 1 must not exceed a coil temperature of -18 degrees Fahrenheit. The Department reserves the right to require additional testing, if deemed necessary.

The facility is required to maintain a capture system and a control device with an overall removal efficiency of at least 75 percent for the removal on acetone associated with Process 07, process equipment COAT7, HOOD4, control equipment COND2, and dust collector DC-18. The facility will monitor and record the coil temperature manually on a weekly basis, when in operation, to verify compliance.



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Condenser 2 must not exceed a coil temperature of -15 degrees Fahrenheit. The Department reserves the right to require additional testing, if deemed necessary.

6 NYCRR Part 212-3

VOC RACT of 81 percent capture and control applies. This is based on the function of process equipment, exhaust and control devices employed. The facility has a 5-year historic emissions values averaging less than 8 tons VOC per year. The owner or operators will monitor and record the scrubber water flow rate manually on a weekly basis when in operation to verify compliance. The water flow rate range must be within a range of 14-20 gallons per minute based off of vendor specifications. The Department reserves the right to require additional testing, if deemed necessary.

6 NYCRR Part 225-1

Operators of a stationary combustion installations that fire distillate oil are limited to the firing of distillate oil with 0.0015 percent sulfur by weight or less on or after July 1, 2016. Compliance with this limit will be based on vendor certifications.

6 NYCRR Part 233

The facility does not engage in chemical synthesis and is therefore no subject to Part 233. Packaging and formulation are not considered to be synthesized pharmaceutical manufacturing as defined in 233.2(13) of the regulation.