



The State of Florida's Section 804 Importation Program (SIP) Proposal for the Importation of Prescription Drugs from Canada

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History of Updates

Version #	Update	Change/Update Details
1	11/23/2020	Submission of SIP document
1.1	04/19/2021	Updated responsible individual for SIP sponsor
1.2	09/15/2021	<p>The following updates were made:</p> <ul style="list-style-type: none"> • Added History of Updates • Updated name and contact information for the Responsible Individual of the SIP co-sponsor • Updated address of importer and FDA-registered relabeler • Updated the name and contact information for the State's point of contact • Provided updated signatures for the sponsor and co-sponsor • Updated key personnel for importer • Updated Attachment C with evidence of relabeler's FDA registration • Updated Attachment D with information for FDA-approved drug, HPFB-approved drug, proposed labeling, and comparison information between FDA-approved label and proposed label • Updated Compliance Plan <ul style="list-style-type: none"> ○ Lines of Communication and Processes ○ Procedures for Noncompliance, Misconduct, and Conflicts of Interest
1.3	11/11/2021	<ul style="list-style-type: none"> • Updated status on foreign seller • Matched the proposed list of drugs to match attachment

Introduction

The State of Florida is submitting this proposal for its Section 804 Importation Program (SIP) as a component of the effort to reduce spending on prescribed drugs. With the state's Agency for Health Care Administration serving as the sponsor and Department of Business and Professional Regulation as co-sponsor, Florida is seeking to begin importing medications for consumers receiving services through the following state agencies/government programs:

- Department of Health (patients served through county health departments)
- Department of Corrections (inmates in the custody of the Department of Corrections)
- Department of Children and Families (patients in a public state mental hospital/treatment facility)
- Agency for Persons with Disabilities (clients residing in a public Institution for Individuals with Development Disabilities)
- Agency for Health Care Administration (recipients served in the Medicaid program)

By implementing an importation program across government agencies, the State of Florida will be able to reap significant savings.

This proposal will describe how the Canadian Prescription Drug Importation program will operate alongside our partners to yield savings to Floridians.

The Agency for Health Care Administration (Agency) is responsible for licensing and regulating over 40,000 health care facilities in the State and is responsible for the administration of the Medicaid program. As such, the Agency is best poised to implement and administer Florida's importation program. It currently oversees the Statewide Medicaid Managed Care program and provides health care for over four million recipients. Additionally, the Agency has experience monitoring large-scale programs for quality and compliance with federal regulations. Acting as the co-sponsor, the Department of Business and Professional Regulation (DBPR) enforces regulations and provides oversight of Florida's prescription drug wholesalers. Its expertise in this area will allow it to support compliance with requirements such as supply chain standards, relabeling, repackaging, and recalling suspect products.

Because of the intricacies involved in operating an importation program, the State will enter into contractual relationships with entities to meet all requirements of the program. This will enable the State to establish business relationships with an importer, foreign seller, manufacturer, and relabeler. In addition, the State will establish relationships with qualifying laboratories to ensure prescription drug authenticity and compliance with U.S. Food and Drug Administration (FDA) requirements. Given the integral role that DBPR plays as the state's regulating authority for drug distribution, it will play a strategic part in implementing Florida's compliance plan, especially related to the handling of recalls and returns.

In regard to prescription drugs, Florida has chosen a more limited set of classes for which to initiate its importation program, focusing on medications that are used to treat certain conditions (e.g., HIV/AIDS, asthma, chronic obstructive pulmonary disease, diabetes, etc.) and that will yield the highest potential savings. After the program has proven successful, the State intends to amend its SIP to expand the list of medications that will be imported.

Ensuring the safe handling of these prescription drugs and having a secure supply chain is paramount to the SIP's success. To maintain safety, the Agency and DBPR will work with its

importer or their designee and contracted third parties to prevent shipments and batches from becoming lost or contaminated through the process. This begins when the prescription drugs marked for sale in Canada are sold to the foreign seller and imported into the U.S. and continues through laboratory testing, relabeling, repackaging, and distribution in Florida. Strict adherence to safety standards is not only necessary for protecting Floridians but also for instilling public trust and confidence in prescription drug importation.

Contained in the following proposal is key information pertaining to Florida's importation program. In addition, the proposal outlines how the Agency and DBPR will maintain a secure supply chain, test sample batches, and label accordingly, all while bringing substantial savings to the state.

Florida's Canadian Prescription Drug Importation Program

The federal rule requires the SIP proposal to include the name of the program, identify the sponsor and co-sponsors, list prescription drugs to be imported, provide addresses of participating parties and companies, and give a summary of how the importation program will function securely. The chart below provides identifying information for the sponsor/co-sponsor and entities involved in the administration/operation of the program.

Name of the Program:	Florida's Canadian Prescription Drug Importation Program
Importation Program Sponsor:	The Florida Agency for Health Care Administration Address: 2727 Mahan Drive, Mail Stop #20 Tallahassee, FL 32308
Importation Program Co-Sponsor:	The Florida Department of Business and Professional Regulation Address: Division of Drugs, Cosmetics, and Devices 2601 Blair Stone Road Tallahassee, FL 32399-1047
Responsible Individuals:	Secretary Simone Marstiller Simone.Marstiller@ahca.myflorida.com (850) 412- 4264 2727 Mahan Drive Bldg. 3 Mailstop 1 Tallahassee, FL 32308 Secretary Julie Brown Julie.Brown@myfloridalicense.com (850) 413-0755 2601 Blair Stone Road Tallahassee, FL 32399-1047
Name and Address of Foreign Seller (must include a copy of their license to operate in Canada):	Methapharm Inc. 81 Sinclair Boulevard Brantford, Ontario N3S 7X6 Canada See Attachment E for a copy of the foreign seller's attestations, licenses, and inspectional histories.
Name and Address of Importer:	Importer of Record Dr. Niaz Siddiqui (Licensed Pharmacist) Back-up: Dr. Danyelle Williams (Licensed Pharmacist) The Florida Department of Health (DOH) Central Pharmacy 104-2 Hamilton Drive Tallahassee, FL 32304 The DOH designee for certain functions: LifeScience Logistics, LLC (Licensed Wholesale Distributor)

	3100 Olympus Blvd, Suite 100 Dallas, TX 75019 See Attachment C for a copy of the importer and their designee's licenses.
Name and Address of the FDA-Registered Relabeler (must include inspection history)	Relabeling: LifeScience Logistics, LLC 3100 Olympus Blvd, Suite 100 Dallas, TX 75019 See Attachment C for inspection history.

Importation Program Summary:

As stated above, the DOH Central Pharmacy serves as the importer under the Florida program. However, as a state entity, they will need logistics support to fulfill all of the requirements of this program. As such, the State is contracting with a licensed wholesale distributor, LifeScience Logistics, LLC to assist the state and importer with the following:

- Work with Methapharm Inc. to import prescription drugs from Canada
- Negotiate drug prices from Methapharm Inc./manufacturer that will yield savings under the program
- Relabel and repackage the product
- Provide logistics support in transporting the eligible drugs into the U.S., including customs clearance, ensuring all laboratory testing is complete, and that the product is trackable and traceable throughout the supply chain.
- Distributing the imported eligible drugs to the end user (pharmacies dispensing on behalf of the state programs).

LifeScience Logistics, LLC is an experienced provider focused solely on the health care supply chain. They are a Verified-Accredited Wholesale Distributor (VAWD), ISO 13485 certified, licensed in all 50 States, and have an excellent state and federal audit/inspection history. They are also fully compliant with the Drug Supply Chain Security Act (DSCSA) requirements. The State is confident in their ability to meet all expectations related to safety and efficacy and will describe how it anticipates those requirements will be met throughout the SIP.

Florida already has robust statutes and rules in place to ensure the safe handling and distribution of prescription drugs, which are more stringent than those of the FDA. As the agency that oversees the regulation of the state's prescription drug market, DBPR will ensure that the SIP participants will adhere to federal, state, and Canadian requirements. This will result in a secure supply chain that verifies the authenticity and purity of imported prescription drugs as well as maintaining strict labeling and packaging standards.

In addition to the statutory requirements listed in Chapter 499, Florida Statutes (F.S.), DBPR's Division of Drugs, Cosmetics, and Devices is responsible for enforcing the rules listed in Chapter 61N of the Florida Administrative Code (F.A.C.). These rules provide requirements that include but are not limited to the following:

- Drug labeling (Rule 61N-1.006, F.A.C.)
- Product tracking and tracing:
 - Manufacturer requirements (Rule 61N-1.029, F.A.C.)

- Wholesale distributor requirements (Rule 61N-1.030, F.A.C.)
- Dispenser requirements (Rule 61N-1.031, F.A.C.)
- Inspections, investigations, and monitoring (Rule 61N-1.019, F.A.C.)

With a rigorous system already in place, DBPR will use its existing infrastructure to oversee the operation of a secure supply chain that safely distributes authentic prescription drugs that complies with and exceeds the FDA's requirements.

Florida intends for its imported prescription drug supply chain to function in the same manner as the domestic one. Once consumers receive their medication, the only difference they should be able to see is a label indicating that their medication was imported from Canada.

The total savings that Florida's importation program can realize is open-ended as continual analyses will be performed to optimize the impact. For the first year, the State is conservatively projecting that it can save between approximately \$80 to \$150 million. These savings will also benefit the federal government because less federal financial participation will be required for Medicaid. However, what Florida's population can save annually once the importation program's benefit fully matures should amount to the hundreds of millions.

Florida's List of Prescription Drugs to Import

The FDA rule requires the SIP to include the following information related to the prescription drugs that will be imported:

- Names and Drug Identification Numbers (DIN) of selected drugs to import
- Information of the applicant that holds the New Drug Applications or Abbreviated New Drug Applications
- Name and address of the manufacturer of the finished dosage form
- Names and addresses of manufacturers of the prescription drugs and active ingredients

As part of this proposal, the State is providing the list of prescription drugs it will initially attempt to import under the SIP. The final list of imported prescription drugs is subject to change and can be addressed in an amended SIP or through the pre-import request, based on the FDA's preference. The State will also update the list of prescription drugs with the manufacturer information once that is confirmed (known).

Florida has chosen a more limited set of eligible prescription drug classes for which to initiate its importation program, focusing on medications that are used to treat certain conditions (e.g., HIV/AIDS, asthma, chronic obstructive pulmonary disease, diabetes, etc.) and that will yield the highest potential savings. Also, these specific medications allow Florida the best opportunity to maximize the importation program's benefits while remaining compliant with federal law. After the program has proven to be a success, the State intends to amend its SIP to expand the list of medications that will be imported.

The list of proposed drugs for this SIP is given below. Full information regarding each drug, including the proposed labels, can be found in Attachment D.

Hepatitis C Prescription Drugs

Drug Name	Active Ingredient	NDC	U.S. Price	NDA/ ANDA	DIN	Canadian Name	Sold in U.S.
Harvoni 90-400mg	Ledipasvir, Sofosbuvir	61958-1801-01	\$1,098.94	NDA205834	02432226	Harvoni	Yes
Epclusa 400-100mg	Sofosbuvir, Velpatavir	61958-2201-01	\$869.05	NDA208341	02456370	Epclusa	Yes
Mavyret 100-40mg	Glecaprevir, Pibrentasvir	00074-2625-80	\$153.11	NDA209394	02467550	Maviret	Yes
Zepatier 50-100mg	Elbasvir, Grazoprevir	00006-3074-01	\$260.00	NDA208261	02451131	Zepatier	Yes

Psychiatric Prescription Drugs

Drug Name	Active Ingredient	NDC	U.S. Price	NDA/ ANDA	DIN	Canadian Name	Sold in U.S.
Latuda 20mg tab	Lurasidone HCL	63402-0302-30	\$40.99	NDA200603	02422050	Latuda	Yes

Drug Name	Active Ingredient	NDC	U.S. Price	NDA/ ANDA	DIN	Canadian Name	Sold in U.S.
Latuda 40mg	Lurasidone HCL	63402-0304-10	\$41.00	NDA200603	02387751	Latuda	Yes
Latuda 60mg	Lurasidone HCL	63402-0306-30	\$41.04	NDA200603	02413361	Latuda	Yes
Latuda 80mg	Lurasidone HCL	63402-0308-30	\$41.03	NDA200603	02387778	Latuda	Yes
Latuda 120mg	Lurasidone HCL	63402-0312-30	\$61.13	NDA200603	02387786	Latuda	Yes

Diabetes Prescription Drugs

Drug Name	Active Ingredient	NDC	U.S. Price	NDA/ ANDA	DIN	Canadian Name	Sold in U.S.
Farxiga 5mg tabs	Dapagliflozin Propanediol	00310-6205-30	\$16.55	NDA202293	02435462	Farxiga	Yes
Farxiga 10mg tabs	Dapagliflozin Propanediol	00310-6210-30	\$16.54	NDA202293	02435470	Farxiga	Yes
Tradjenta 5mg tabs	Linagliptin	00597-0140-30	\$14.79	NDA201280	02370921	Tradjenta	Yes
Januvia 50mg tabs	Sitagliptin Phosphate	00006-0112-31	\$15.13	NDA021995	02388847	Januvia	Yes
Januvia 25mg tabs	Sitagliptin Phosphate	00006-0221-31	\$15.13	NDA021995	02388839	Januvia	Yes
Januvia 100mg tabs	Sitagliptin Phosphate	00006-0277-31	\$15.14	NDA021995	02303922	Januvia	Yes

Asthma and Chronic Obstructive Pulmonary Disease Prescription Drugs

Drug Name	Active Ingredient	NDC	U.S. Price	NDA/ ANDA	DIN	Canadian Name	Sold in U.S.
Spiriva 30 caps, 18mcg	Tiotropium Bromide	00597-0075-41	\$436.80	NDA021395	02246793	Spiriva	Yes
Spiriva Respimat 2.5 mcg	Tiotropium Bromide	00597-0100-61	\$461.62	NDA021936	02435381	Spiriva	Yes
Combivent Respimat 20-100 mcg	Ipratropium Bromide, Albuterol Sulfate (aka salbutamol sulfate)	00597-0024-02	\$408.76	NDA021747	02419106	Combivent Respimat	Yes
Wixela Inhub 100-50 mcg	Fluticasone Propion, Salmeterol	00378-9320-32	\$123.00	ANDA208891	02495597	Wixela Inhub	Yes

Drug Name	Active Ingredient	NDC	U.S. Price	NDA/ ANDA	DIN	Canadian Name	Sold in U.S.
Wixela Inhub 250-50 mcg	Fluticasone Propion, Salmeterol	00378-9321-32	\$150.00	ANDA208891	02495600	Wixela Inhub	Yes
Wixela Inhub 500-50 mcg	Fluticasone Propion, Salmeterol	00378-9322-32	\$199.20	ANDA208891	02495619	Wixela Inhub	Yes
Incruse Ellipta 62.5 mcg	Umeclidinium	00173-0873-10	\$329.40	NDA205382	02423596	Incruse Ellipta	Yes

HIV/AIDS Prescription Drugs

Drug Name	Active Ingredient	NDC	U.S. Price	NDA/ ANDA	DIN	Canadian Name	Sold in U.S.
Kaletra 200-50mg	Lopinovir, Ritonovir	00074-2605-21	\$8.53	NDA021906	02285533	Kaletra	Yes
Kaletra 100-25mg	Lopinovir, Ritonovir	00074-0522-60	\$4.27	NDA021906	02312301	Kaletra	Yes
Genvoya 150-150-200-10 mg tabs	Elviteg, COB, Emtri, Tenof, Alafen	61958-1901-01	\$104.93	ANDA207561	02449498	Genvoya	Yes
Complera 200-25-300 mg tabs	Emtricit, Rilpivirine, Tenofovir DF	61958-1101-01	\$95.52	NDA202123	02374129	Complera	Yes
Stribild 150-150-200-300 mg	Elviteg, COB, Emtricit, Tenofovir Disop	61958-1201-01	\$109.51	NDA203100	02397137	Stribild	Yes
Emtricitabine, Tenofovir DF 200-300mg	Emtricitabine, Tenofovir DF	00093-7607-56	\$45.69	ANDA090894	02399059	TEVA-Emtricitabine, Tenofovir	Yes
Odefsey 200-25-25 mg	Emtricit, Rilpivirine, Tenofovir Ala	61958-2101-01	\$95.77	NDA208351	02461463	Odefsey	Yes
Edurant 25 mg	Rilpivirine HCL	59676-0278-01	\$37.58	NDA202022	02370603	Edurant	Yes
Intelence 100mg	Etravirine	59676-0570-01	\$10.92	NDA022187	02306778	Intelence	Yes
Intelence 200mg	Etravirine	59676-0571-01	\$22.41	NDA022187	02375931	Intelence	Yes
Triumeq 50-600-300 mg	Abacavir, Dolutegravir, Lamivudine	49702-0231-13	\$98.19	NDA205551	02430932	Triumeq	Yes
Biktarvy	Bictegravir, Emtricit, Tenofovir Ala	61958-2501-01	\$104.52	NDA210251	02478579	Biktarvy	Yes

Drug Name	Active Ingredient	NDC	U.S. Price	NDA/ ANDA	DIN	Canadian Name	Sold in U.S.
Isentress 400mg	Raltegravir	00006-0277-61	\$26.74	NDA022145	02301881	Isentress	Yes
Descovy	Emtricitabine, Tenofovir 200mg-25mg	61958-2002-01	\$59.67	NDA208215	02454424	Descovy	Yes
Juluca	Dolutegravir, Rilpivirine	49702-0242-13	\$91.75	NDA210192	02475774	Juluca	Yes

Attestations and Information Statement

The FDA final rule language requires the SIP to:

Include an attestation and information statement containing a complete disclosure of any past criminal convictions or violations of State, Federal, or Canadian laws regarding drugs or devices against or by the responsible individual(s), Foreign Seller, or Importer or an attestation that the responsible individual(s), Foreign Seller, or Importer has not been involved in, or convicted of, any such violations. Such attestation and information statement must include principals, any shareholder who owns 10 percent or more of outstanding stock in any non-publicly held corporation, directors, officers, and any facility manager or designated representative of such manager.

The State is including the necessary attestations in Attachment B except for the foreign seller, which is included in Attachment E.

Disciplinary Actions and Inspectional History

The FDA final rule requires the SIP proposal to include:

- A list of all disciplinary actions, to include the date of and parties to any action imposed against the responsible individual(s), Foreign Seller, or Importer by State, Federal, or Canadian regulatory bodies, including any such actions against the principals, owners, directors, officers, quality unit, or any facility manager or designated representative of such manager for the previous 7 years prior to submission of the SIP Proposal.
- The Health Canada inspectional history for the Foreign Seller for the previous 5 years or, if the Foreign Seller has been licensed for less than 5 years, for the duration of its period of licensure; and the State and Federal inspectional history for the Importer for the previous 5 years or, if the Importer has been licensed for less than 5 years, for the duration of its period of licensure.

The State's importers are employed by the Florida Department of Health (DOH) Central Pharmacy. The DOH Central Pharmacy has an exemplary track record and performance history. Evidence of the importer's performance is provided in Attachment A.

Florida is also including the inspectional history for LifeScience Logistics, LLC. This information is included in Attachment C.

In addition, the importer has entered into an agreement with a Canadian prescription drug wholesaler, Methapharm Inc., to serve as the foreign seller. Methapharm Inc.'s performance and inspectional history is included in Attachment E.

Evidence that Imported Drugs are Commercially Available

The FDA final rule requires that the SIP proposal provide adequate evidence that each Health Products and Food Branch (HPFB)-approved drug's FDA-approved counterpart drug is currently commercially marketed in the United States.

The State has verified that the prescription drugs it seeks to import have FDA-approved counterparts that are readily available in the U.S. market. Florida can provide evidence as listed in the chart beginning on **pages 10-13** that identifies drug names, active pharmaceutical ingredients, National Drug Codes (NDC), and shared manufacturers with locations in both the U.S. and Canada.

Description of Qualifying Laboratory Testing Techniques

The FDA final rule requires that the SIP proposal:

Describe, to the extent possible, the testing that will be done to establish that the HPFB-approved drug meets the conditions in the NDA or ANDA for the HPFB-approved drug's FDA-approved counterpart. The SIP Sponsor's importation plan must also identify the qualifying laboratory that will conduct the Statutory Testing for the Importer, if the Importer is responsible for conducting the Statutory Testing, and it must establish that the laboratory is qualified in accordance with § 251.15 to conduct the tests.

Overview

Ensuring prescription drug purity and authenticity is essential to the success of Florida's SIP. The State intends to make every effort to partner with manufacturers that will perform the FDA required testing on each imported drug. Some of the eligible prescription drugs Florida proposes to import are manufactured by the same companies for the U.S. and Canadian markets. In addition, these products are produced in the same facilities, on the same manufacturing lines, and contain identical specifications and standards. Because these Canadian products are fully compliant with FDA-approved New Drug Applications (NDA) (except for labeling), it will not be necessary to perform statutory testing on these products.

The State will provide to the FDA evidence to establish that these products are manufactured according to the specifications in the FDA-approved NDAs and will ensure that these products are relabeled appropriately. This will avoid duplicative testing efforts and reduce overall costs by avoiding unnecessary markups for added steps as drugs proceed through the supply chain. However, if that is not possible, through its contracted entities, the State will ensure that a qualifying laboratory will test statistically valid sample batches or shipments of imported prescription drugs in accordance with the FDA's current and good manufacturing practices (CGMPs) as specified in Title 21 Code of Federal Regulations Sections 211 and 251.16.

The qualifying lab will obtain a statistically valid sample sufficient for testing and retesting as necessary. In addition, it will retain these samples for at least one year following the batch's expiration date as stated in federal rule (or longer as necessary). Each selected sample will undergo testing to evaluate authenticity, stability, and contamination. If a sample fails in any of the three categories, it will be prohibited from distribution and dispositioned.

The following sections outline the specific laboratory tests Florida is considering utilization when screening imported prescription drugs.

Selecting Samples for Testing

For laboratory testing to accurately verify whether a prescription drug is authentic, the selected samples analyzed must be randomly chosen using a statistically valid sampling plan (i.e., ANSI/ASQ Z1.4-2008 or MIL-STD-105E). To ensure this, the Agency and DBPR will require the importer or its designee to take the following steps when selecting samples:

- The importer or its designee must pull samples directly from the shipment or batch and cannot require the foreign seller or manufacturer to submit samples separately.
- The selection process must not expose the prescription drugs to possible contamination or adulteration. The importer cannot unseal containers and reseal them.

- The importer or its designee must select samples from multiple points in a shipment and not a single area. This is to ensure that temperature and environmental conditions have not adversely affected certain parts of a shipment and not others.
- The importer or its designee must segregate selected samples from the shipment or batch and ensure they are kept in the same environmental and climate conditions prior to testing.
- The importer or its designee and qualifying laboratory must not disclose how they selected samples.

Evaluating for Authenticity

When assessing the authenticity of a prescription drug, Florida will require its qualifying lab to use visual inspections, spectrometry, and chromatography. Combined, these will examine visual characteristics (color, labeling, identifying marks) and physical properties (active pharmaceutical ingredients and excipients) to determine whether they are identical to their FDA-approved counterparts in the U.S. In addition to identifying counterfeits, the authenticity testing can also discern whether the prescription drug meets purity requirements by checking for the presence of foreign substances or chemical toxins. If a sample fails in any one of the categories, its originating batch will be removed from the supply chain and dispositioned.

Visual Inspections: Before testing, laboratories can start ascertaining whether a prescription drug is authentic by examining its visual properties such as labeling, pill color and shape, and pill markings. Counterfeits sometimes lack expiration dates or instructions and can use poor quality ink or have grammatical mistakes. Additionally, pills improperly colored or that have the wrong markings are direct indications of fakes. The qualifying laboratory will be required to have a process to visually inspect selected samples and document their authenticity. Any prescription drugs identified as being inconsistent with the actual product will be dispositioned immediately and not undergo further testing.

Laboratory Testing: Florida will require its qualifying laboratory to verify authenticity based on the results of multiple tests. This is due to no single technique delivering the universal results needed for verification. Additionally, the qualifying lab will have the necessary equipment to perform detailed testing on the samples using spectroscopic and high-performance liquid chromatography, as applicable, based upon the methods specified by the manufacturer

Assessing Stability

Unlike evaluating for authenticity, laboratory tests for stability do not assess whether a prescription drug is genuine but whether its active pharmaceutical ingredient and excipients will retain their medicinal properties to be of benefit to individuals taking them. This is necessary for not only measuring effectiveness but ensuring that certain ingredients will not become toxic before use, particularly those that are unstable such as nitroglycerine. In addition, analyzing stability provides the opportunity to assign expiration dates to batches. In regard to imported prescription drugs, stability testing is essential to determine those coming into the U.S. have not expired prior to entry and remain just as effective as newly manufactured ones in Canada. To test for stability, two methods are available, real time and accelerated. The qualifying lab will

have the necessary equipment to perform either stability test on the sample batch, based upon the method specified by the manufacturer.

In addition to testing prescription drugs following entry into the U.S., the qualifying laboratory will be required to retain samples for retesting at certain intervals (e.g., six months, one year) depending upon each product's FDA-approved counterpart's shelf life. Prescription drugs deemed to have expired or will expire before being able to be safely consumed will be designated for disposition.

Testing for Biological Contamination

To evaluate whether a batch poses biological hazards, the qualifying lab will test for harmful bacteria by using culture media swabs on the selected sample. This includes gathering swabs on pills in bottles or other containers and blister packs, although the risk of biological contamination in blister packs is reduced. Following an incubation period of 48 to 72 hours, the qualifying laboratory will identify any microorganisms present and assess whether they can potentially harm humans. Samples that present evidence of contamination will have their originating batches removed and dispositioned.

Names and Addresses of Qualified Laboratories

The State will use the following laboratories to meet the testing requirements:

Contract Laboratories		
Name of the Organization / Address	Registrations	FDA Audit History
Micro Quality Labs 3125 N. Damon Way Burbank, California 91505 818-845-0070 www.microqualitylabs.com	FDA Registered (DUNS 149592615) ISO 17025 (Certificate 3034.1)	Feb 2018 – FDA (In good standing)
Avomeen, LLC 4840 Venture Drive Ann Arbor, MI 48108 833-507-6831 www.avomeen.com	FDA Registered (FEI 3008808597) ISO 17025 (Certificate L19-183)	Jun 2016 – FDA Mar 2018 – FDA Oct 2018 - FDA (In good standing)
Avista Pharma Solutions, Inc dba Cambrex Corp 104 Gold St Agawam, MA 01001 843-312-6529 www.cambrex.com	FDA Registered (FEI 1220785) ISO 17025 (Certificate L2190)	2016 – FDA 2018 – FDA 2019 – FDA (In good standing)
BioChroma Analytical 1309 Record Crossing Rd Dallas, TX 75235 972-454-9166 www.biochromalabs.com	FDA Registered (FEI 3012816120) ISO 17025 (Certificate L20-206)	Feb 2017 - FDA (In good standing)

Prescription Drug Labeling Comparison

The FDA final rule requires the SIP to:

Include a copy of the FDA-approved drug labeling for the FDA-approved counterpart of the eligible prescription drug, a copy of the proposed labeling that will be used for the eligible prescription drug, and a side-by-side comparison of the FDA-approved labeling and the proposed labeling, including the Prescribing Information, carton and container labeling, and patient labeling (e.g., medication guide, instructions for use, patient package inserts), with all differences annotated and explained. The SIP Proposal must also include a copy of the HPFB-approved labeling.

For this submission, the State is providing in Attachment D the proposed labels that it will use for imported prescription drugs when they are distributed in Florida.

As for the remaining components, the State will include the following in a forthcoming submission:

- FDA-approved labels for prescription drugs that it seeks to import from Canada
- Proposed labeling for each imported prescription drug
- HPFB-approved labeling for each imported prescription drug
- Side-by-side comparison of the FDA-approved labeling and proposed labeling

Explanation of Cost Savings

The FDA final rule requires the SIP to:

Explain how the SIP Sponsor will ensure that the SIP will result in a significant reduction in the cost to the American consumer of the eligible prescription drugs that the SIP Sponsor seeks to import. The explanation must include any assumptions and uncertainty, and it must be sufficiently detailed to allow for a meaningful evaluation.

The Agency for Health Care Administration collected information from each participating state agency on the amount spent on all prescription drugs in the second quarter of calendar year 2018 and then compared the unit cost of each drug to its Canadian equivalent formulation. Because many of the state programs already benefit from deep discounts through other government programs (e.g., 340B covered entities, Medicaid supplemental rebate program requirements, etc.), the State removed drugs that were already deeply discounted and would not yield any greater savings or were ineligible for inclusion due to other restrictions. This analysis will be performed on a quarterly basis to continually assess opportunities to focus in on a narrow list of prescription drugs that yields the greatest amount of savings.

Florida has identified the initial list of prescription drugs that will be included in the initial launch of the importation program. As previously stated, the State will be evaluating the list on a quarterly schedule to determine if changes are needed and if it continues to be cost-effective to import those specific drugs.

Following full implementation, Florida is projecting over \$150 million dollars in annual savings. The below table presents an example of the analysis conducted to determine the potential cost savings under the SIP using a sample of drugs used to treat HIV/AIDS (the data represents utilization and costs for one quarter of 2018).

Brand Name	Utilization	Net Unit Cost	Total Spend/Actual Spend	Canadian Unit Cost *	Potential Spend	Estimated Difference
Atripla Tablet	48,513	\$ 87.90	\$ 4,264,305	\$ 42.25	\$ 2,049,674	\$ 2,214,618
Complera Tablet	15,477	\$ 85.66	\$ 1,325,757	\$ 42.65	\$ 660,094	\$ 665,666
Genvoya Tablet	225,815	\$ 94.98	\$ 21,448,515	\$ 47.63	\$ 10,755,568	\$ 10,692,340
Isentress 400 Mg Tablet	104,607	\$ 24.15	\$ 2,525,748	\$ 12.51	\$ 1,308,634	\$ 1,217,625
Odefsey Tablet	66,699	\$ 86.46	\$ 5,766,699	\$ 42.66	\$ 2,845,379	\$ 2,921,416
Prezista 800 Mg Tablet	71,933	\$ 51.09	\$ 3,675,303	\$ 21.26	\$ 1,529,296	\$ 2,145,761
						\$ 19,857,428

*Includes a markup

The State has included a markup on the Canadian price to account for additional costs potentially imposed by the foreign seller.

Since Florida's program exclusively focuses on serving consumers who are receiving their medications through government programs (some funded with both state and federal dollars), any savings derived will benefit all American taxpayers. The savings will not only reduce the overall state budget, but those dollars can be reinvested in other programs/services that support Floridians. To the extent consumers pay co-payments on the imported drugs, the savings can

also be used to offset copayment amounts, again resulting in savings to the American consumer.

Storage, Handling, Supply Chain, and Reporting Guidelines

The FDA final rule language requires the SIP proposal to:

- Explain how the SIP Sponsor will ensure that all the participants in the SIP comply with the requirements of section 804 of the Federal Food, Drug, and Cosmetic Act.
- Describe the procedures the SIP Sponsor will use to ensure that the requirements are met, including the steps that will be taken to ensure that the:
 - Storage, handling, and distribution practices of supply chain participants, including transportation providers, meet the requirements and do not affect the quality or impinge on the security of the eligible prescription drugs
 - Supply chain is secure
 - Importer screens the eligible prescription drugs it imports for evidence that they are not adulterated, counterfeit, damaged, tampered with, expired, suspect foreign product, or illegitimate foreign product
 - Importer fulfills its responsibilities to submit adverse event, field alert, and other reports required by the SIP, the Federal Food, Drug, and Cosmetic Act, or this part.

As the most essential element to ensuring that imported prescription drugs are identical to their FDA-approved counterparts, the State understands that maintaining a secure supply chain is integral to the process. By requiring safe storage, handling, and transportation practices along with robust screening regimens, Florida will prevent counterfeit, contaminated, or adulterated drugs from entering the market. In the event that the importer's screening process detects unfit prescription drugs, it will immediately take actions to maintain the health and safety of Floridians.

Having a robust and closed supply chain beginning in Canada and ending with the delivery of prescription drugs to individuals in Florida requires the foreign seller, importer, and manufacturer to follow multiple requirements. In addition to complying with the U.S. Drug Supply Chain Security Act (DSCSA), participating parties in Florida's program must also adhere to the minimum requirements for storage and handling as specified in Title 21 CFR Section 205.50 and Section 499.0121, F.S. (Note that Florida's requirements mirror those listed in the CFR). Because Florida is considering the importation of HIV/AIDS medications, all of which have specific temperature requirements, ensuring a secure supply chain and safe handling and storage practices is paramount to providing Floridians reliable imported prescription drugs.

The primary responsibility of ensuring the delivery of safe imported prescription drugs belongs to the importer or its designee. The importer (or designee) will maintain all transaction histories, information, and statements in addition to having adequate facilities that meet cleanliness and climate standards. The following describes the supply chain and the handling, storage, and transportation practices Florida's importer (or designee) will utilize to import prescription drugs.

Storage, Handling, and Distribution

All storage facilities and vehicles used to transport imported prescription drugs must meet specific state and federal guidelines. This includes not only those located within the U.S. but in Canada as well. To further ensure compliance and safeguard the integrity of the supply chain, LifeScience Logistics will provide dedicated and fully licensed distribution space within Florida. This will prevent the possibility of inadvertently comingling drug products from the program with other drug products and distribution channels. The importer or its designee will need to provide

documentary proof that Canadian facilities and vehicles meet the same requirements as their counterparts in the U.S. These requirements as listed in federal rule include but are not limited to the following:

- Facilities used for storing and/or marketing prescription drugs must have adequate size, storage conditions, quarantine areas, cleanliness, and security.
- Storage areas must have climate control and accurate instrumentation for measuring temperature and humidity.
- Having written policies and procedures that ensure the oldest approved stock is distributed first, handling recalls and withdrawals, ensuring the facility can function during a crisis, and removing outdated prescription drugs from those designated for distribution.

Additionally, the importer or its designee will need to ensure that the prescription drugs remain in temperature-controlled climates throughout importation and distribution. This is due to HIV/AIDS anti-retroviral medications requiring an environment that cannot exceed 20-25 degrees Centigrade to maintain potency and effectiveness. Due to Florida's tropical climate, controlling temperature becomes more necessary as the imported prescription drugs get distributed across the state.

The importer or its designee will be responsible for providing the Agency and DBPR with a list of vendor-approved Canadian and U.S. facilities that will store the prescription drugs in addition to the vendor-approved carriers that will transport and distribute. The list must include not only the facility names and addresses but proof that they meet FDA and Health Canada's licensing standards. Also, the importer or its designee will provide a flow chart that presents the route imported prescription drugs and their active pharmaceutical ingredients (APIs) will take beginning with the country of origin through the port of entry in the U.S. and where they will be stored during laboratory testing and relabeling before going to their final points of distribution in Florida.

Having a Secure Supply Chain

Signed into law in 2013, the DSCSA updated the requirements that pharmaceutical companies, wholesalers, and distributors must follow to prevent counterfeit, adulterated, or contaminated prescription drugs from reaching consumers. As the SIP sponsor, the State will require the importer or its designee to verify at receipt, maintain, and submit all transaction histories, information, and statements. When monitoring for compliance, the Agency and DBPR will review the transaction documents and verify their accuracy as well as confirm that all prescription drugs being imported meet Health Canada and FDA guidelines.

The State understands the purpose of the DSCSA and plans to hold the foreign seller, importer, and manufacturer accountable for documenting each change of ownership during the process. The State's selected vendor, LifeScience Logistics, LLC fully complies with the DSCSA (including the components that are not yet enforced by the FDA – i.e., serialization). Its system will receive inbound electronic data using industry standard formats, such as the Advance Ship Notice (ASN/856) and Electronic Product Code Information Services (EPICS) documents. Additionally, its system also captures and stores all transaction information, histories, and statements and supports full serialization of the imported product. LifeScience Logistics will also accommodate the transmission of the transaction information, histories, and statements via

paper, as allowed by the DSCSA regulations, and its system will capture the following transactional information:

- Product name
- Strength and dosage
- National Drug Code
- Number and size of containers
- Lot number
- Transaction and shipment dates
- Names and addresses of the businesses that complete transactions.

Because these prescription drugs will be imported from Canada, Florida will require transactional information to include Canadian information such as Drug Identification Numbers and Health Products and Food Branch (HPFB) proprietary names.

LifeScience Logistics will also physically inspect each drug shipment received from Methapharm Inc. against shipping paperwork and a set of specifications developed for each drug imported. These specifications include damage, tamper seal intact, lot number, DIN number, and determining whether expiration dating on packaged units aligns with shipping paperwork and if there is no presence of counterfeit or illegitimate products. All packaging inspections will be documented, reviewed by the quality assurance staff, and included in the import receipt files.

Data Availability and Documentation

Florida's importation program will use industry-leading software to provide the required features, functions, and capabilities of a warehouse management system and a transportation management system. When a product is received into the system, important transaction information is captured and stored. Within the warehouse, all products are tracked by their lot numbers. On outbound shipments, all required information is provided in print and electronically to comply with applicable federal regulations. The system also tracks when it receives, stores, and ships by individual serial numbers.

LifeScience Logistics has developed a CGMP compliant set of standard operating procedures (SOPs) that ensure each product is handled, stored, and distributed in accordance with applicable FDA, Drug Enforcement Agency, and State of Florida guidelines. In addition to the guidelines associated with facilities, training, document control, change control, equipment, temperature monitoring, vendor qualification, security, pest control, redundancy, deviation and corrective action/preventative action, Florida will maintain SOPs governing all processes associated with products inbound, inventory management, order management, returns, and preventive/corrective maintenance. The State will also require LifeScience Logistics to maintain policies and procedures for identifying, recording, and reporting losses or thefts and for correcting all errors and inaccuracies in inventories. They must include in their written policies and procedures:

1. A procedure whereby the oldest approved stock of a prescription drug product is distributed first. The procedure may permit deviation from this requirement if the deviation is temporary and appropriate.
2. A procedure for addressing any crisis that affects security or operation of any facility if a strike, fire, flood, or other natural disaster, or a local, state, or national emergency occurs.

3. A procedure to ensure that any outdated prescription drugs are segregated from other drugs and returned to the manufacturer or destroyed.
4. A procedure that prevents the diversion of prescription drugs.

LifeScience Logistics leverages a cloud data center provider to supply infrastructure for its technology systems. Specifically, their system provides real-time redundancy across two data centers in different geographic regions within the United States. The system is load-balanced across these two data centers. A failure of any or all components in a single data center would cause a real-time failover to the second data center with no user impact or loss of data.

Reporting Adverse Incidents and Filing Field Alerts

When an imported prescription drug fails testing, becomes compromised, has a recall issued, or results in patient injury, the Agency and DBPR will require the importer or its designee to conduct adverse incident reporting and issue field alerts to state and federal agencies. All adverse incidents must be reported to the FDA's Adverse Event Reporting System (FAERS) and the Agency and DBPR. Additionally, LifeScience Logistics is required to follow FDA guidelines when filing field alerts by doing so within 72 hours of becoming informed of one or more of the following issues:

- Patient injury or death
- Labeling problems that can cause the prescription drug to be identified as another product
- Biological contamination
- Changes in the chemical or physical composition of the prescription drug that leads to deterioration, degradation, or toxicity
- Any failure of a shipment or batch of prescription drugs to meet the specifications in its NDA or ANDA

In addition to submitting these reports to state and federal agencies, the State will also require the importer or its designee to inform Health Canada and the HPFB of any defect, contamination, or adulteration of a prescription drug. The importer will report these issues formally in accordance with Canadian standards and procedures.

Additional Reporting

The SIP Sponsor and co-sponsor will submit quarterly reports to the FDA consisting of the following information required by Title 21 CFR Section 251.19:

- The name, address, telephone number, and professional license number of the importer
- The name and quantity of the active ingredient of the imported eligible prescription drug(s)
- A description of the dosage form of the eligible prescription drug(s)
- The date(s) on which the eligible prescription drug(s) were shipped; the lot or control number assigned to the eligible prescription drug(s) by the manufacturer of the eligible prescription drug(s)
- The point of origin (i.e., manufacturer) and the destination (i.e., the wholesale, pharmacy, or patient to whom the importer sells the drug) of the eligible prescription drug(s)
- The per unit price paid by the importer for the prescription drug(s) in U.S. dollars

- Any other information the FDA determines is necessary for the protection of the public health.

The quarterly reports will also include the importer's confirmation that it purchased eligible prescription drug(s) directly from the foreign seller. In addition, the quarterly reports will include the following documentation:

- A listing of manufacturers of each eligible prescription drug
- The quantity of each lot of eligible prescription drug received by the foreign seller from the manufacturer
- Proof that the eligible prescription drug was received by the foreign seller from the manufacturer and subsequently shipped by the foreign seller to the importer
- Results of the statutorily required laboratory testing and descriptions of the sample selection methods used for each eligible prescription drug

The State will ensure that the report contains a certification from the importer that each shipment of each eligible prescription drug is approved for marketing in the United States and is not adulterated or misbranded and that it meets all labeling requirements under the Federal Food, Drug, and Cosmetic Act. The certifications will note the following:

- That there is an authorized SIP
- That the imported drug is covered by the authorized SIP
- That the drug is an eligible prescription drug as defined by this rule
- That the FDA-approved counterpart of the drug is currently commercially marketed in the United States
- That the drug is approved for marketing in Canada.

Lastly, the quarterly reports will also include data, information, and analyses on the SIP's cost savings to the American consumer.

Education and Outreach Plan

The FDA final rule language requires the SIP proposal to:

Explain how the SIP Sponsor will educate pharmacists, healthcare providers, pharmacy benefit managers, health insurance issuers and plans, as appropriate, and patients about the eligible prescription drugs imported under its SIP.

The SIP is a novel concept, and Florida is a trailblazer by working to influence the cost of prescription drugs. Because of how innovative this program is, the Agency in coordination with DBPR will be taking great steps when considering how to provide education and training resources to state-run facilities, other agencies, and Florida Medicaid providers and recipients.

These will include webpages, webinars, written guides available online, brochures, and infographics. In addition, the Agency will prepare materials to inform Medicaid beneficiaries of what prescription drug importation means and how it does not pose a risk to their health and safety. As the SIP sponsor and co-sponsor, the Agency and DBPR believe that everyone involved in prescription drug importation, beginning with organizations to the consumers should be aware of where their medications originate and how obtaining them from Canada benefits the entire state.

Webpages and Webinars

When seeking to understand new programs and regulations, the State understands that many individuals search for information independently. To accommodate these people, the Agency is contemplating the construction of a webpage that provides detailed information on the SIP and how it works. Available on this page will be resources such as Florida's original concept paper, the approved SIP proposal draft, links to the FDA Importation of Prescription Drugs final rule, the Florida Statutes, a list of all imported prescription drugs, and any guides or brochures created. The goal of the webpage is to serve as a continuous resource to answer any questions about the program. In addition, it will provide a link for consumers to make complaints and offer contact information for further questions. For ease of use, the webpage will be linked to the Agency's homepage, making it accessible with minimal "clicks" to access the page.

During the beginning phases of prescription drug importation, the Agency will also schedule multiple webinars to train relevant stakeholders (state agencies, and providers/facilities). These webinars will go over how the importation program functions and the stakeholder roles to ensure program success.

For interested parties unable to attend a webinar, the Agency will record them and make the recordings accessible on the prescription drug importation webpage. Each webinar will be accompanied by a PowerPoint presentation and provide opportunities for attendees to ask questions.

Written Guides

For detailed information, the Agency and DBPR will prepare written guides that will provide more specific information than what is covered in the webinars and narrative content on the webpage. Each guide will address one of the following importation program components for interested parties to research:

- Labeling and packaging, with detailed visual examples and comparisons

- Qualifying laboratory testing methods and standards
- Safe storage, handling, and transporting processes and procedures
- Recall processes, with specific information on procedures for each of the three tiers

During the drafting process, the Agency and DBPR will work with the importer or its designee to include thorough and accurate information that can address in-depth questions. As with the webinars and other materials, these written guides will be available on the Agency's prescription drug importation webpage.

Brochures and Infographics

To promote the SIP and generate stakeholder support, the Agency will design multiple brochures and infographics. Each of which will provide high-level information on imported prescription drugs and visuals showing the supply chain, labeling, and how savings will be achieved. Although these materials will not be intended to provide detailed training to providers, they can serve as a useful resource to assist with educating the public and communicating the general purpose and function of the SIP.

Other Education and Training Measures

The State understands that additional training may be needed to fully convey the scope of importing prescription drugs. To ensure that all interested parties are clear on the concept and operationalization of the importation program, both the Agency and DBPR can hold conference calls on an ad hoc basis to discuss specific issues, assign staff to focus solely on handling questions, and send representatives to meetings.

Recall and Return Plan

The FDA final rule language requires the SIP proposal to:

- Include the SIP's recall plan, including an explanation of how the SIP Sponsor will obtain recall or market withdrawal information and how it will ensure that recall or market withdrawal information is shared among the SIP Sponsor, the Foreign Seller, the Importer, and FDA and provided to the manufacturer.
- Include the SIP's return plan, including an explanation of how the SIP Sponsor will ensure that product that is returned after distribution in the United States is properly dispositioned in the United States, if it is a non-saleable return, in order to protect patients from expired or unsafe drugs, and an explanation of how the SIP Sponsor will prevent the non-saleable returned eligible prescription drugs from being exported from the United States. In the event that a returned eligible prescription drug may be considered saleable, include an explanation for how the returned product will be determined to be saleable and under what circumstances such eligible prescription drugs may be re-distributed in the United States.

Currently, U.S. prescription drug manufacturers have to follow FDA guidelines when recalling products. However, medications imported to the U.S. under a SIP will not only have to follow domestic policies but adhere to Canadian standards as well. Depending on the medication and where its active pharmaceutical ingredient is manufactured, the State will monitor FDA recall alerts as well as Canadian recalls.

If a recall is ordered on a shipment of imported prescription drugs, the Agency and DBPR will immediately halt the importation of affected prescription drugs under the SIP in accordance with the FDA's Importation of Prescription Drugs final rule. Additionally, they will take actions to work with the importer or its designee and those participating in the SIP to communicate the need to isolate those drugs and return them for disposition. To inform stakeholders and affected parties, the Agency and DBPR understand that messaging must go out to health plans, pharmacies, state run facilities (e.g., public health clinics, prisons, state mental hospitals), and other state agencies. The importer's (or their designee's) role will be to follow a process for safe handling and disposal of the recalled products in addition to ensuring all non-dispensed inventory is collected.

If at any time, the Agency or DBPR determines that an issue is present in the SIP, it can issue a recall and halt the importation of prescription drugs. Below is the recall and return plan that the State will follow when the SIP becomes operational.

Agency and DBPR Communication Plan

In addition to requiring the importer to monitor the FDA's MedWatch and Health Canada's Recalls and Safety Alerts, the Agency and DBPR will check daily for any notifications pertaining to imported prescription drugs. Both the FDA and Health Canada use the following three-tiered system for classifying recalls.

- **Tier 1:** Recalled prescription drug poses severe risks to individuals that can result in serious health complications or death.
- **Tier 2:** Recalled prescription drug may cause a temporary health problem or have a slight chance of posing a serious health complication.

- **Tier 3:** Recalled prescription drug is in violation of labeling or manufacturing laws and does not pose a significant risk to individuals' health.

Given the consequences of delayed or insufficient action, the Agency and DBPR will implement immediate measures beginning with communication to all stakeholders involved in the importation program, including: SMMC health plans, participating pharmacies, county health departments, state-run facilities, and state agencies. Communications will be made via email blasts and direct calls to administrators to inform them of the recall and potential risks as well as instructions for returning drugs to the importer. Those entities will be responsible for notifying patients that have received the recalled medications.

While the Agency and DBPR handle communications with stakeholders, the importer or its designee will administer collecting the recalled prescription drugs and their disposition. Additionally, the importer will submit a report to the Agency and DBPR explaining the quantity of prescription drugs recovered, the dates of recovery, the number of those unaccounted for, and where they are stored. The importer will also include in its report the number of recalled prescription drugs distributed to each provider/facility and the individual quantities recovered from them. For Tier 1 and Tier 2 recalls, the Agency and DBPR will contact those providers/facilities unable to collect recalled prescription drugs to determine what actions may be needed to get the products off the market.

The importer will also need to provide the Agency and DBPR with specific information on which recalled prescription drugs went through the supply chain and were distributed to providers/facilities. This information must consist of the following:

- Section 804 serial identifier (SSI), National Drug Code (NDC), Drug Identification Number (DIN), and manufacturer's assigned lot number
- Number and size of containers
- Dates of transactions and shipments between the foreign seller, manufacturer, and importer

Importer Recall Plan

If a recall is required, the importer or its designee will be responsible for collection, documentation, storage, and destruction of the suspect prescription drugs. In addition, it will also halt the importation of affected prescription drugs in accordance with federal rule. While the Agency and DBPR oversee the communications aspect, the importer or its designee will immediately begin working with distributors, providers, and facilities to collect the recalled products. Once retrieved, the importer or its designee will gather all the returned prescription drugs and store them in a single facility under quarantine. Depending on the reasons for the recall, the importer or designee will also oversee their secure destruction. Unless otherwise specified, the importer will follow the same process for all three tiers of recalls.

At the recall's outset, the importer or its designee must use its track and trace procedures as established under the Drug Supply Chain Security Act (DSCSA) to identify which batches or shipments it received require collecting. It will verify this information with the foreign seller and manufacturer. The importer or its designee must confirm with the manufacturer that it has identified all suspect shipments by comparing lot numbers, DINs, and dates of transactions and shipments. Additionally, the importer will locate where all recalled prescription drugs are at in

the supply chain (e.g., in storage, at the laboratory, distributed to providers, etc.) and submit a report consisting of the following information to the Agency and DBPR.

- Location of each batch and shipment
- Quantity of prescription drugs at each location
- Identification information (SSI, DIN, NDC, lot number, dates of transactions and shipments)
- Dates of distribution to providers/facilities, if applicable

During the recall process, the importer (or designee) will provide daily updates to the Agency and DBPR on the quantities collected for Tier 1 and 2 recalls. For Tier 3, the importer can provide one update per week. The importer will provide updates until the recall process is complete and then on an ad hoc basis as required.

When disposing of recalled prescription drugs, the FDA and Drug Enforcement Agency (DEA) do not require a specific method for destruction. However, the importer or its designee needs to ensure that disposition occurs in the U.S. and does not involve discarding prescription drugs as trash or possibly contaminating a water supply. In addition, the destruction process needs to ensure the recalled products are physically destroyed or rendered as non-retrievable. Following disposition, the importer must submit a report to the Agency and DBPR specifying that each batch or shipment was destroyed and provide identification information.

In accordance with FDA and Health Canada requirements, manufacturers can voluntarily engage in recalls. To further ensure that suspect imported prescription drugs do not enter the market, the Agency and DBPR are granting the importer this same ability. If at any time, the importer or its designee determines that a recall is necessary, it can issue one.

Return Plan

For imported prescription drugs that must go through the return process, the Agency and DBPR will require the importer to ensure that all collected products remain in the original supply chain (e.g., its own storage facilities). The importer or its designee will return the prescription drugs to its storage facility and keep them isolated from non-recalled ones. At no time, can the importer send returned prescription drugs to an outside facility or transport them via a different means without direct approval from the Agency and DBPR and only with justification (e.g., facility is beyond capacity or has been compromised). By mandating that these prescription drugs remain within the supply chain, the State can ensure that they do not enter the black market or are exported to another country.

At each point during the return process, the Agency will review the importer's reports and assess whether any prescription drugs are missing from the list of batches and shipments. Additionally, the Agency will immediately work with the importer to resolve any discrepancies. In the event that a discrepancy cannot be resolved, the Agency and importer will ascertain at which point the prescription drugs became misplaced and issue communications to affected parties or contact law enforcement if theft is suspected.

In the event a recalled prescription drug can be returned to market, the Agency will require its importer to use the following procedures to ascertain whether the product is saleable. This can only apply to Tier 3 recalls that occurred due to a labeling mishap or other issue that poses no risk to individuals taking the medication.

- Assess whether the prescription drugs have expired, and if not, determine if a reasonable timeframe exists to verify purity and potency and return to the market.
- For non-expired prescription drugs, the importer must randomly select new samples for testing to evaluate purity, potency, and the presence of contamination.
- For prescription drugs recalled due to labeling issues, the importer will have all batches and shipments relabeled after they have passed laboratory testing.

Prior to returning to market, the importer or its designee will have verified that the prescription drug is saleable and that the issue prompting the recall was resolved. It will report this information to the Agency and DBPR. In addition, the importer will not begin redistributing the prescription drugs until the Agency and DBPR have given approval.

Compliance Plan

The FDA final rule language requires the SIP's compliance plan to include:

- A description of the division of responsibilities among co-sponsors, if any, which includes a plan for timely communication of any compliance issues to the SIP Sponsor
- Identification of responsible individual(s) and a description of the respective area(s) of the SIP, the Federal Food, Drug, and Cosmetic Act, or this part that will be under each responsible individual's oversight
- The creation of written compliance policies, procedures, and protocols
- The provision of education and training to ensure that Foreign Sellers, Importers, qualifying laboratories, and their employees understand their compliance-related obligations
- The creation and maintenance of effective lines of communication, including a process to protect the anonymity of complainants and to protect whistleblowers
- The adoption of processes and procedures for uncovering and addressing noncompliance, misconduct, or conflicts of interest.

As the SIP sponsor and co-sponsor, the Agency and DBPR will assume primary responsibility for overseeing compliance with the program's requirements. Because it will manage the contract or agreement with the importer or its designee, the Agency will monitor performance, while DBPR ensures adherence to state and federal regulations. In addition, the Agency and DBPR will ensure that the foreign seller, qualifying laboratory, relabelers, repackagers, and other subcontractors comply as well. To maintain transparency, all participating entities will routinely submit detailed reports to the Agency on their performance. Additionally, the Agency, in collaboration with DBPR, will conduct routine on-site visits of these entities and their facilities as well as any of those under their subcontractors.

Working together, the Agency and DBPR will use the following strategy for ensuring compliance. The remainder of this section outlines the multiple components of Florida's compliance plan as specified in the FDA's Importation of Prescription Drugs final rule.

Division of Responsibilities Among Sponsor/Co-Sponsors

The Agency, acting as the importation program sponsor, will manage the contract with the importer or its designee and monitor its performance. As the importation program co-sponsor, DBPR will collaborate with the Agency to ensure that the importer or its designee and subcontractors comply with state and federal prescription drug wholesale and distribution regulations, including but not limited to Chapter 499, F.S. and the Drug Supply Chain Security Act (DSCSA).

Identification of Responsible Individual(s) and Their Respective Area(s)

Operationalizing the SIP

The importer or its designee will assume full responsibility for operationalizing the SIP and submits reports to the Agency that describes compliance with all requirements. Since LifeScience Logistics will be performing many duties on behalf of the importer and State (as described earlier), the following is a list of key corporate executive staff and their qualifications:

Key Personnel Name: Richard Beeny	
Key Personnel Position: Chief Executive Officer	
April 2006 - Current	Richard is Co-founder and Chief Executive Officer of LifeScience Logistics. He has more than 25 years of supply chain experience and has held a variety of operations, marketing, and business development roles. Richard has held leadership positions at United Parcel Service and has served in both the U.S. Navy and U.S. Coast Guard. He holds a Bachelor of Arts degree from the University of Texas at Arlington and a Master of Business Administration from Southern Methodist University.

Key Personnel Name: Randy McCollom	
Key Personnel Position: Vice President of Operations	
November 2020 - Current	Randy McCollom, Vice President of Operations, has over 20 years of supply chain experience. Prior to joining LifeScience Logistics, Randy held a variety of roles in operations and inventory management with McKesson Corp, Thermofisher, and ConMed Corporation. He is focused on operational efficiency, process improvement and quality control. Randy attended Troy State University and is a Designated Representative in multiple states.

Key Personnel Name: Joseph Fountaine	
Key Personnel Position: Project Manager	
December 2011 - Current	As LifeScience Logistics' Director of Information Technology & Infrastructure Services. Joseph's responsibilities include oversight of Inventory Control, WMS systems, and all facility infrastructure to maintain operational readiness of five CGMP-compliant facilities. In addition, Joseph is currently the Program Manager for five GSA contracts: GS-00T-11-AJC-0010, GS-00T-11-AJC-0008, GS-00T-12-AJC-0002, 47QFCA20C0002, 47QFCA20C0014.

Key Personnel Name: Paul Hayward	
Key Personnel Position: Vice President of Quality & Compliance	
December 2015 - Current	Paul was named Director of Quality Assurance & Regulatory Affairs in December of 2015. Prior to joining LifeScience Logistics, Paul served as the Vice President of Operations for Azaya Therapeutics, UrgentRx, and Pernix Therapeutics. Paul has over 21 years of experience in quality assurance, manufacturing operations, product development, validation, and supply chain management in the pharmaceutical medical device and biologics industries with organizations that include Reckitt Benckiser, and Allergan Pharmaceuticals. Paul holds a Bachelor of Biology and Chemistry from Southwest Baptist University and a Master of Science in Chemistry from Baylor University. He is a member of the American Society for Quality and Regulatory Affairs Professionals Society.

Key Personnel Name: Chris Mizener	
Key Personnel Position: Head of Drug Importation	
September 2012 - Current	Chris joined LifeScience Logistics in 2012 after 16 years with United Parcel Service. At UPS Chris had several roles within operations, industrial engineering, and finance. He was appointed Director of Client Services in 2016 and is responsible for account management and customer service. Chris holds a Bachelor of Arts degree in Accounting from LeMoyne College in Syracuse, New York.

Individuals Responsible for SIP Oversight

The Agency and DBPR are appointing the following staff members to oversee compliance with the importation program. Each individual will monitor areas specific to their own expertise.

Key Personnel Name: Devona "D.D." Pickle
Key Personnel Position: Program Director, Canadian Prescription Drug Importation
Responsibility: Oversight of the contract with LifeScience Logistics and day to day workflow of the state of Florida Canadian Drug Importation Program.

Key Personnel Name: Walter Copeland
Key Personnel Position: Director, Division of Drugs, Devices, and Cosmetics
Responsibility: Compliance and oversight of prescription products and administering the provisions of the Florida Drug and Cosmetic Act consistent with Florida statutes.

Compliance Policies, Procedures, and Protocols

The Agency will maintain policies that govern how this program will operate and approve the standard operating procedures that are developed by LifeScience Logistics in the operation of the program, on the State's behalf. The contract between the Agency and LifeScience Logistics will also outline delegated duties.

Because Florida is in the process of developing rules specific to prescription drug importation, the contract will provide the detailed requirements to operationalize the SIP and consist of the following requirements. Because implementing the SIP's operational components may require multiple contracts or agreements with third parties, the State may delegate these responsibilities across multiple entities.

- Have an organizational structure that is adequately staffed to operate the SIP
- Have a physical presence in the state of Florida (e.g., corporate office or subsidiary branch dedicated to administering the importation program)
- Have approved agreements in place with a foreign seller registered both in Canada and the U.S., a qualifying laboratory that has ISO 17025 licensing and meets current and good manufacturing practices, a relabeler, and storage facilities that can provide environmental conditions suitable for the imported prescription drugs
- Ensure that the foreign seller has an agreement with a manufacturer to purchase the prescription drugs specified in this proposal
- Ensure adherence with track and trace requirements as specified in the DSCSA by having an electronic tracking system that collects all transaction statements, histories, and information and can document a prescription drug's point of origin through to its distribution
- Provide quality assurance throughout the importation process and monitor all parties involved in the pharmaceutical supply chain
- Ensure that prescription drugs deemed unfit for market in Florida are dispositioned
- Ensure that relabeling and repackaging processes are completed in accordance with FDA guidelines
- Have a procedure in place that requires the following:
 - Submission of pre-import requests at least 30 days prior to shipping prescription drugs into the U.S.
 - Use of the U.S. Customs and Border Protection's Automated Commercial Environment or other approved means of data exchange
- Have a recall and return plan in accordance with that outlined in the Recall and Return section of this proposal
- Have a system for tracking and resolving consumer complaints and an internal quality control plan

In regard to all the above listed aspects, the Agency and DBPR will conduct regular monitoring through yearly on-site visits, weekly and ad hoc calls, and desk reviews. The importer or subcontractors will be required to submit monthly deliverables specifying the number of prescription drugs imported, their testing results (e.g., number of selected samples tested, comparisons to FDA-approved prescription drugs), amounts paid, and number and characteristics of complaints (e.g., open and resolved).

If the importer or subcontractors do not adhere to the contract's terms and conditions, the Agency can impose a corrective action plan, assess liquidated damages, or terminate the agreement.

Provision of Compliance-Related Education and Training

Before entering into any agreement with an importer or its designee, the Agency will ensure that the importer or its designee operationalizing the SIP, as well as its subcontractors, fully

understands its responsibilities regarding state, federal, and Canadian regulations for prescription drug importation. To ensure that the importer or its designee is able to sufficiently train all participating parties and staff involved in the SIP, the State will require proof of the following:

- Educational materials used to train staff and third-party subcontractors regarding the following areas:
 - Storage and handling of prescription drugs
 - How to identify counterfeits or adulterated products based on visual inspections
 - Processes for filing pre-import requests and using the U.S. Customs and Border Protection Automated Commercial Environment
 - Policies and procedures of Health Canada and the Canadian Health Products and Food Branch
 - Processes for recalls and returns
 - Rules regarding relabeling and repackaging
 - Overviews of the FDCA; DSCSA; Chapter 499, F.S.; and the Importation of Prescription Drugs final rule
 - Overview of laboratory testing required for imported prescription drugs and the result thresholds to qualify for entry to Florida's market

Prior to dissemination among staff and subcontractors, the Agency and DBPR will review all educational and training materials to ensure they are aligned with state, federal, and Canadian requirements. In addition, participating entities will not be allowed to begin importing prescription drugs until they have received approval for all educational and training materials.

Lines of Communication

The State will require the importer or its designee to have multiple lines of communication, including a customer service team available to take complaints. In addition, the Agency and DBPR also have separate lines that consumers can use for complaints. In regard to whistleblowers, the Agency, DBPR, and the importer will be compliant with the Federal Whistleblower Protection Act and the Florida Whistleblower Protection Act.

The Florida Whistleblower Protection Act (section 448.102, F.S.) prohibits employers from retaliating against employees who report or threaten to report violations of rules or statutes to a government agency. If an employee makes a complaint alleging that his/her employer is engaged in illegal activities, the investigating government agency will protect their identity during the investigative process.

Regarding violations of Florida's prescription drug wholesaler/distributor regulations, DBPR has a specific online portal for the filing of complaints under the Division of Drugs, Devices, and Cosmetics. Individuals may report concerns with the importer and not have their identities disclosed. In addition to conducting its own investigation, DBPR will communicate the reported issue to the Agency. The Agency will then further investigate as to whether the problem affects compliance with any prescription drug importation regulation that falls beyond DBPR's scope

The Agency also has multiple lines of communication open to the public and employees of the importer, foreign seller, or other subcontracted entity. These include directly contacting the Florida Medicaid Helpline, the Agency's Division of Health Quality Assurance, and/or the Agency's Bureau of Medicaid Program Integrity. All three have full-time teams tasked with

handling and responding to complaints while adhering to federal whistleblower protection statutes. If an individual reports an issue pertaining to the SIP, the receiving team will forward to the Agency's Canadian Prescription Drug Importation Program team to address with the importer.

When communicating and investigating SIP complaints, the Agency will not disclose the reporting individuals' identities. In addition, any employee of the SIP sponsor, SIP co-sponsor, importer, or subcontracted entity that performs tasks related to the SIP is eligible for whistleblower protection as stated under section 448.102, F.S.

Additionally, the importer is contractually required to have a customer service hotline and personnel dedicated to receiving and responding to complaints. As part of its contractual requirements, the importer must maintain a dashboard for complaint tracking and submit a monthly report to the Agency that consists of all complaints reported, dates filed, resolutions, and time spent resolving.

Both the Agency and importer or its designee will also each have a full-time contract manager who will be available to address issues when they arise. The contract managers are dedicated staff with open lines of communication and can quickly receive and disseminate information.

Processes and Procedures for Noncompliance, Misconduct, and Conflicts of Interest

The Agency, in its role as the SIP sponsor, will monitor the importer to ensure full compliance with the FDA's final rule. Although federal regulators will conduct inspections and document reviews, the Agency will take a proactive approach that can identify deficiencies and implement corrective actions before these occur. In addition, DBPR will routinely inspect and report any issues that violate State regulations for prescription drug wholesaler/distributors (Chapter 499, F.S. and Chapter 61N, F.A.C.). Combined, both agencies will oversee the SIP to ensure full compliance with the FDA's final rule, the FDCA, and the Drug Supply Chain Security Act (DSCSA).

Certain aspects of the Program such as the importer's distribution facility fall under existing State regulations. DBPR currently monitors prescription drug wholesalers for adequacy of storage space, temperature controls, security, and quarantine areas.

The State will conduct routine monitoring through monthly, quarterly, and ad hoc reports, as well as on-site reviews for program components that are specific to the FDA's final rule:

- Transaction information, statements, and histories in accordance with the DSCSA
- Pre-import requests prior to their submission to the FDA
- Vendor's methods for ensuring imported prescription drugs are not contaminated or counterfeit
- Labeling of each imported prescription drug
- Transaction documentation between the foreign seller and the vendor
- Laboratory testing documentation
 - Documentation will consist of types of tests performed and the results in addition to identifying whether the manufacturer or a contracted laboratory completed the testing.
- Adverse event reports and individual case safety reports concurrent with submission to the FDA

- Quarterly reports prior to submission to the FDA
 - These reports consist of multiple components that require documentation that verifies purchase from the foreign seller, laboratory testing, and certifications that the prescription drugs are eligible for sale in Canada.

The State reserves the right to inspect all records and facilities operated by the vendor, including those of any subcontractor (if applicable). The State will mirror the FDA's areas of oversight to remain proactive in monitoring, identifying, and correcting deficiencies. The State will conduct onsite visits, both announced and unannounced, to verify that the importation process is compliant with the FDA's final rule. Because the logistics of sending Agency staff to Canada will be costly, the Agency will delegate this to the importer who is contractually responsible for the foreign seller's performance.

The State has established criteria that the importer must use when inspecting the foreign seller's Canadian facility, including providing visual evidence of compliance (e.g., photographs, videos), opportunities for Agency staff to ask questions via livestream. Additionally, the importer's inspections will occur on an annual and ad hoc basis and will observe the following for compliance:

- Separation of inventory that is intended for sale in the U.S.
- Quarantine areas for suspect or illegitimate products
- Records of Canadian prescription drugs purchased that are intended for sale in the U.S.
 - The foreign seller must maintain these records for six years
- Processes and procedures for receiving, storing, placing identifiers, and shipping prescription drugs intended for sale in the U.S.

Following a completed inspection, the importer must submit a written report to the State that specifies the foreign seller's areas of compliance and deficiencies. The State will direct the importer to work with the foreign seller to make changes and corrections as necessary.

The State has the means to dispatch staff to inspect and assess the importer's Florida warehouse for compliance. These onsite inspections will consist of the following:

- Guided and unguided tours of the facility
- Spot checks of randomly selected lots of prescription drugs to verify correct labeling and packaging
- Interviews of randomly selected staff to assess regulatory knowledge as it applies to their individual job duties
- Review of database to examine current inventory information and then verify accuracy by confirming the exact specifications on the warehouse floor
 - The database review will also verify compliance with the DSCSA.

In the course of conducting its monitoring activities, the Agency will work in collaboration with DBPR, taking note of any obvious discrepancies (e.g., unsecure areas, sanitation issues), advising the importer to correct them, and notifying DBPR.

In addition to onsite inspections, the State will conduct desk monitoring on a continuous basis. This will consist of the vendor submitting draft reports for the FDA, shipment information, laboratory testing results, and labeling examples. If deficiencies are present, the Agency will

request an explanation and consider whether an unannounced onsite inspection is necessary to assess severity.

Authenticating Information and the Protection of Trade Secrets

The FDA final rule language requires the SIP proposal to:

Explain how the SIP Sponsor will ensure that any information that the manufacturer supplies to authenticate a prescription drug being tested and confirm that the labeling of the prescription drug complies with labeling requirements under the Federal Food, Drug, and Cosmetic Act, and any trade secrets or commercial or financial information that is privileged or confidential that the manufacturer supplies for the purposes of testing or otherwise complying with the Federal Food, Drug, and Cosmetic Act and this part, are kept in strict confidence and used only for the purposes of testing or otherwise complying with the Federal Food, Drug, and Cosmetic Act and this part.

Supplying Authenticating Information

As co-sponsor, DBPR will ensure that the manufacturer selected to provide imported prescription drugs supplies any necessary information to verify their authenticity and labeling accuracy. This includes not only that specified in the Federal FDCA and DSCSA but also that listed in Florida's rules (Chapter 61N, F.A.C.) and statutes (Chapter 499, F.S.). In addition, the manufacturer will be responsible for providing all information required for the laboratory testing that includes the following:

- Breakdown of the weights and measurements of APIs and excipients per tablet for each medication in accordance with the prescription drugs' FDA-approved NDA or ANDA
 - Note: The qualifying laboratory will conduct testing against the FDA-approved drug as the standard.
- Copies of all Canadian and U.S. labeling, packaging, and instructions
- Images of the U.S. and Canadian prescription drug tablets with identifying marks clearly visible

To verify labeling accuracy, the importer must submit a sample of the proposed label to the State upon request.

Protection of Trade Secrets

Although Florida prides itself on its transparency and access to public records, it has protections in place for withholding trade secrets and proprietary confidential business information. The state's public records statutes (Chapter 119, F.S.) do provide exemptions for the disclosure of trade secrets and proprietary confidential business information. In addition, DBPR has a rule (Rule 61N-1.021, F.A.C.) that provides a procedure for the manufacturer and importer to make known what needs to remain confidential. Provided that information is identified as a trade secret or confidential business information in documents submitted to DBPR or obtained during an inspection, this information will not be disclosed if sought through a public records request. LifeScience Logistics maintains client and vendor confidentiality through mutual NDAs, business contracts and quality agreements.

To ensure that the authenticating information is protected, the importer and the designee will have a written policy regarding confidential information and trade secrets. Additionally, the importer, the designee, and any party receiving confidential information from the manufacturer will provide yearly training to their employees on protecting confidential information and the requirements under the Federal Food, Drug, and Cosmetic Act to protect confidential

information from disclosure, specifically confidential information that the manufacturer provided/supplied regarding the prescription drug(s). The training will also address the penalties associated with failing to maintain the information as confidential.

Attachment A

Florida Department of Health Central Pharmacy Licenses, Disciplinary and Inspection Reports,
and Attestation



Walter Copeland, Director
Drugs, Devices, and Cosmetics
2601 Blair Stone Road
Tallahassee, Florida 32399-1047
Phone: 850.717.1800 • Fax: 850.414.8240

Halsey Beshears, Secretary

Ron DeSantis, Governor

October 26, 2020

RE: Inspections/Disciplinary Actions

To whom it may concern;

Chapter 119.01(1), Florida Statutes, provides that it is the policy of this state, county, and municipal records are open for personal inspection and copying by any person. Providing access to public records is a duty of each agency.

After a review of our Department's records and databases no disciplinary action has been taken against the below entity.

Chapter 499.051, Florida Statutes, provides that the agents of the Department of Business and Professional Regulation and of the Department of Law Enforcement, after they present proper identification, may inspect, monitor, and investigate any establishment permitted pursuant to this chapter during business hours for the purpose of enforcing this chapter and the rules of the department that protect the public health, safety, and welfare.

Rule 61N-1.019(2), Florida Administrative Code, states the department may inspect, monitor, and investigate all drugs, device and cosmetic manufacturers, wholesalers, repackagers, distributors, or other establishments where drugs, devices or cosmetics are made, stored, sold, offered for sale, exposed for sale, or kept for sale or use, for the purpose of determining compliance with the provisions of chapters 499 and 893, Florida Statutes, or any rules adopted thereunder and to secure evidence of any non-compliance.

Inspections are conducted on initial permit applications. Inspections conducted by the division may be announced or unannounced and are not scheduled routinely or on a yearly basis.

The last know inspection conducted at:

DOH Central Pharmacy
104-2 Hamilton Park Drive
Tallahassee, Florida 32304
Permit Number 28:29

was conducted on **October 10, 2019** with a result of **PASS**.

Regards,

A handwritten signature in blue ink that appears to read "Walt Copeland".

Walter Copeland
Division Director

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The State upon completion of the negotiation process and selection of the Foreign Seller will obtain complete history as mandated by federal law. This includes the following:

- All disciplinary actions from the past seven years issued by federal, state, or Canadian authorities
- The Health Canada inspectional history for the foreign seller for the previous 5 years or duration of license if less
- The state and federal inspectional history for the importer for the previous 5 years or duration of license if less

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Lic Type	3328 - Prescription Drug Repackager		Entity #	9637670
Fed Tax #	9637670	Fed Tax Type	FEIN #	Name DOH CENTRAL PHARMACY

 Org Name	DOH CENTRAL PHARMACY			<input type="checkbox"/>	Notes
Org Type					Notes History
Mailing Address	<input checked="" type="checkbox"/>	Private Address	<input type="checkbox"/>	Preferred Name	<input type="checkbox"/>
Street #	104-2	Street	HAMILTON PARK DRIVE		
Line 2					
Line 3					
City	TALLAHASSEE	County	Leon		
Zip	32304	State	Florida		
Routing					
Phone #	850-922-9036	Ext	E-Mail		
Insp Region				Receive Email	<input type="checkbox"/>
Updated	02/10/2009 08:43:17		By bcoates		

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Lic Type 3308 - Product Registration Permit Fed Tax # 9652043 Fed Tax Type PIN #	Entity # 9652043 Name DOH CENTRAL PHARMACY
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Org Name DOH CENTRAL PHARMACY
 Org Type
 Mailing Address ☒ Private Address ☐ Preferred Name ☒

Street # **104-2** Street **HAMILTON PARK DRIVE**
 Line 2
 Line 3
 City **TALLAHASSEE** County **Leon**
 Zip **32304** State **Florida** Country
 Routing
 Phone # **850-922-9036** Ext E-Mail
 Insp Region
 Updated **06/09/2009 08:27:23** By **jhlittle**

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Lic Type 3328 - Prescription Drug Repackager	Entity # 9637670
Fed Tax # 9637670 Fed Tax Type FEIN #	Name DOH CENTRAL PHARMACY

Type	Name	Status	Lic Type	File #	Updated By	Preferred	
Primary	DOH CENTRAL PHARMACY	Current			katkins	<input type="checkbox"/>	

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Lic Type 3328 - Prescription Drug Repackager	Entity # 9637670
Fed Tax # 9637670	Fed Tax Type FEIN # DOH CENTRAL PHARMACY

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LL	104-3 HAMILTON PARK DRIVE, TALLAHASSEE FL 32304 Leon US	850 922-9036	ISIAH.HILL@FLHEALTH.GOV	DOH CENTRAL PHARMACY	Current	<input type="checkbox"/>	<input type="checkbox"/>
EC	1829 SAGEBROOK DR, TALLAHASSEE FL 32303 Leon US	850-445-2215	DARREN.EVANS@FLHEALTH.GOV	EVANS, DARREN	Current	<input type="checkbox"/>	<input type="checkbox"/>
AC	104-2 HAMILTON PARK DRIVE, TALLAHASSEE FL 32304 Leon US	850-922-9036	ISIAH.HILL@FLHEALTH.GOV	HILL, ISIAH E	Current	<input type="checkbox"/>	<input type="checkbox"/>
VEM			kenneth.grice@flhealth.gov		Current	<input type="checkbox"/>	<input type="checkbox"/>

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Department of Health



License Verification

DANYELLE BRIONNE WILLIAMS

Printer Friendly Version

License Number: PS53818

Data As Of 10/26/2020

License Information	Secondary Locations	Discipline/Admin Action	Subordinate Practitioners
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Name	Relationship	Profession	Effective License Date
WILLIAMS, DANYELLE BRIONNE	PHARMACISTSUBORDINATE	CONSULTANT PHARMACIST	8210 6/5/2018

Click on the License Number to view License Details for that Practitioner

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<https://mqa-internet.doh.state.fl.us/MQASearchServices/HealthcareProviders/LicenseVeri...> 10/26/2020



Department of Health



DANYELLE BRIONNE WILLIAMS

License Number: PS53818

Data As Of 10/26/2020

Profession	Pharmacist
License	PS53818
License Status	CLEAR/ACTIVE
Qualifications	Certified To Administer Immunizations
License Expiration Date	9/30/2021
License Original Issue Date	08/05/2015
Address of Record	104-2 HAMILTON PARK DRIVE TALLAHASSEE, FL 32304
Discipline on File	No
Public Complaint	No

The information on this page is a secure, primary source for license verification provided by the Florida Department of Health, Division of Medical Quality Assurance. This website is maintained by Division staff and is updated immediately upon a change to our licensing and enforcement database.



Department of Health



License Verification

NIAZ AHMED SIDDIQUI

Printer Friendly Version

License Number: PS42647

Data As Of 10/26/2020

License Information	Secondary Locations	Discipline/Admin Action	Subordinate Practitioners		
Name	Relationship	Profession	Effective License Date		
SIDDIQUI, NIAZ AHMED	PHARMACISTSUBORDINATE	CONSULTANT PHARMACIST	7204 2/7/2013		

Click on the License Number to view License Details for that Practitioner

[Back](#)

For instructions on how to request a license certification of your Florida license to be sent to another state from the Florida Department of Health, please visit the License Certifications web page.


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Department of Health



NIAZ AHMED SIDDIQUI

License Number: PS42647

Data As Of 10/26/2020

Profession	Pharmacist
License	PS42647
License Status	CLEAR/ACTIVE
Qualifications	Certified To Administer Immunizations
License Expiration Date	9/30/2021
License Original Issue Date	08/01/2007
Address of Record	1100 SW 11TH ST LIVE OAK, FL 32060
Discipline on File	No
Public Complaint	No

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ENTRY NOTICE AND ON-SITE INSPECTION REPORT

DBA Name:
Full Legal Name: DOH CENTRAL PHARMACY
Inspection Reason: Chg Locate - X
Address: 104-3 HAMILTON PARK DRIVE
TALLAHASSEE FL 32304

Form Date: Nov 19, 2019 10:54 - Nov 19, 2019 13:05
Permit Number: 29 Rank: 28
Telephone Number: 850-922-9036
Inspection Visit ID: 7123926

File Number: 5931
Permit Type: 3328

INSPECTION RESULT Completed

Inspection Authority 499.051 F.S., 61N-1.019, F.A.C. (03/10/2017)

Person Receiving Report/Title

Danyelle Williams, Interim Bureau Chief.

Facility Hours 61N-1.015(2)(c), F.A.C.

Hours	Monday	Tuesday	Wednesday	Thursday	Friday	Saturday	Sunday
Open	See notes						
Close							

FACILITY/STORAGE [(Rx drug establishments) FS 499.01(2) and 499.0121(1)(b),(d),(e)]

- Is the establishment a residence? [F.S. 499.012(1)(b)] No
- Lighting adequate? [(OTC drug manufacturers) FS 499.01(2)(n)3 and 21 CFR 211.44] [(Cosmetic manufacturers) 61N-1.010(2)(e), F.A.C.] Yes
- Pest Free? [(OTC drug manufacturers) FS 499.01(2)(n)3 and 21 CFR 211.56] [(Cosmetic manufacturers) 61N-1.010(2)(b), F.A.C.] Yes
- Clean and orderly? [(OTC drug manufacturers) 61N-1.014(4), F.A.C.] [(Cosmetic manufacturers) 61N-1.010(2)(a), F.A.C.] Yes

RECORD KEEPING [F.S. 499.0121(6)] [61N-1.012 F.A.C.]

- All records computer based? No Electronic and Manual.
- Location of hard copy? On site
- All records manual? No Electronic and Manual
- On-site storage? Yes
- Shares facility? No
- If yes to #9, separate records? N/E

PRESCRIPTION DRUG ESTABLISHMENT

- Is the name identical to another person authorized to purchase legend drugs? [F.S. 499.012(1)(c)] [61N-1.015(2)(b), F.A.C.] N/E The firm has not yet been issued the repackager permit for this location
- Is the person a broker only as defined under [61N-1.001(2)(d), F.A.C.]? No

SECURITY [(Rx drug establishments) FS 499.0121(2)(a), (b) and 499.0121(10)(b) or 61N-1.013(1), F.A.C.]

- Access restricted? [(OTC drug manufacturers) FS 499.01(2)(n)3 and 21 CFR 211.28(c)] [(Cosmetic manufacturers) 61N-1.010(4)(f), F.A.C.] Yes Employees have a swipe badge to enter. Visitors have to be buzzed in by the firm.
- Drug facility alarmed? Yes

November 19, 2019 at 1:05:58 PM EST
Location:
License #: 2829
Inspector: Donald Yerbey

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- | | | |
|--|-----|---|
| 15. Telephone monitoring? | Yes | Sonitrol |
| 16. Controlled substance cage (CII-CV)? [21 CFR 1301.72] | N/E | This location will not stock or distribute controlled substances. |
| 17. Controlled Substance CII vault? [21 CFR 1301.72] | N/E | |

STORAGE [(Rx drug establishments) FS 499.0121(3)(5) or 61N-1.013(3)(4), F.A.C.] [(OTC drug manufacturers) 61N-1.014, F.A.C., and/or FS 499.01(2)(n)3 and 21 CFR 211.46] [(Cosmetic manufacturers) 61N-1.010(2), F.A.C.]

- | | | |
|---|-----|------------|
| 18. Quarantine area? | Yes | |
| 19. Air conditioner? | Yes | |
| 20. Method of monitoring? | | Electronic |
| 21. Refrigeration present? | No | |
| 22. Handles drugs or cosmetics requiring refrigeration? | | NA |
| 23. Method of monitoring refrigeration? | | NA |
| 24. Freezer present? | No | |
| 25. Handles drugs or cosmetics requiring freezing? | No | |
| 26. Method of monitoring freezing temperatures? | | NA |

WRITTEN POLICIES AND PROCEDURES [(Rx drug establishments and OTC manufacturers) FS 499.0121(6)(8)(10), 499.01(2) and/or 21 CFR Part 211] [(Cosmetic manufactures) 61N-1.010, F.A.C.]

- | | |
|---|-----|
| 27. Exist? | Yes |
| 28. Rx drug annual examination? [F.S. 499.0121(5)(a)2] | Yes |
| 29. Drug FIFO inventory? | Yes |
| 30. Recalls/emergencies? | Yes |
| 31. Natural disasters for Rx drug establishments? | Yes |
| 32. Receipt and distribution business records? | Yes |
| 33. Transaction Information, History and Statement (T3) for Rx drug products? | N/E |

INBOUND PRESCRIPTION DRUG BUSINESS RECORDS (Purchases and/or Receipts) [FS 499.0121(6)(b)] [61N-1.012(1)(a),(2)(a), F.A.C.]

- | | |
|--|---------|
| 34. What are firm's all-inclusive inbound business records that it receives from its drug source(s) per firm for this establishment? | Invoice |
|--|---------|

OUTBOUND PRESCRIPTION DRUG BUSINESS RECORDS (Sales or Otherwise) [FS 499.0121(6)(b)] [61N-1.012(1)(a),(2)(a), F.A.C.]

- | | |
|--|---|
| 35. What are firm's all-inclusive typical outbound business records (e.g., sales invoice, packing list, etc.) that it provides to the purchaser and recipient for each Rx drug distribution from this establishment according to firm? | Invoice. Firm also provides a repack order and a shortage/overage report. Firm states the shortage/overage report accounts for any discrepancies between the bottle count and the repackaged drug count |
|--|---|

- | | |
|--|-----|
| 36a. Did firm provide its all-inclusive outbound business records for one (1) of its typical Rx drug distribution? | Yes |
|--|-----|

- | | |
|--|-----------------|
| 36b. If yes to 36a, what is the tracking number (e.g., sales invoice or order number, etc.) and distribution date for this provided distribution document? | Invoice number. |
|--|-----------------|

- | | |
|---|---------|
| 37. Of the document(s) provided in 36a, which document contains the minimum required recordkeeping business elements according to firm? | Invoice |
|---|---------|

- | | |
|---|--|
| 38. For the document type referenced in 37 regarding this establishment's outbound Rx drug distributions: | |
|---|--|

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Inspector: Donald Yerbey

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38a. Did firm mark "A" next to the field that lists the seller's name and address?	Yes	
38b. Did firm mark "B" next to the field that lists the seller's Florida permit number?	Yes	
38c. Did firm mark "C" next to the field that lists the shipper's name and address when different from the seller?	Yes	Ship from and sold from will always be the same
38d. Did firm mark "D" next to the field that lists the distribution date?	Yes	
38e. Did firm mark "E" next to the field that lists the name, strength, dosage form, quantity, and (when assigned) the National Drug Code (NDC) of the Rx drug?	Yes	
38f. Did firm mark "F" next to the field that lists the financial data?	Yes	
38g. Did firm mark "G" next to the field that lists the purchaser's name, address, and Florida permit number?	Yes	
38h. Did firm mark "H" next to the field that lists the physical recipient's name, address and permit or license number, when the recipient is different from the purchaser?	Yes	

N/E = Not Evaluated

Inspection Notes

Change of Address inspection Rx Drug Repackager. The Firm is relocating from 104-2 Hamilton Park Drive, Tallahassee, FL to 104-3 Hamilton Park Drive, Tallahassee, FL

Arrived at Facility approximately 1000

Credentials and Bill of Rights presented to Isaiah Hill. Government Consultant II. Also present representing the firm was Carolyn Albaugh, Business manager.

I. PERMITS/LICENSES:

A. FDA REGISTRATION – The Firm currently holds establishment registration at 104-2 Hamilton Park Drive, Tallahassee, FL Registration # 1036356 – Repackager.

B. DEA Registration

The Firm is already registered at this location with the DEA. RF0356015. registration is under the name of FL DOH Bureau of Public Health Pharmacy

Business Activity is listed as Chempack/SNS Distributor
Schedules covered are listed as II, III, IV and V.

C. PHARMACY PERMITS – Firm holds Community Pharmacy permit PH 23821 located at 116 A Hamilton Park Drive, Tallahassee, FL.

D. DBPR PERMITS –

The Firm holds the following permits with DBPR at 104-2 Hamilton Park Drive, Tallahassee, FL
RPK – 2082 - This permit will be transferred to 104-3 Hamilton Park Drive, when issued.

November 19, 2019 at 1:05:58 PM EST
Location:
License #: 2829
Inspector: Donald Yerbey

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50:238

54:33

28:29 – This permit will be transferred to 104-3 Hamilton Park Drive, when issued.

The firm holds the following permits with DBPR at 116-A Hamilton Park Drive, Tallahassee, FL

50:237

54:29

The firm hold DBPR permit 54:258 at this location (104-3 Hamilton Park Drive)

This permit is for Emergency Preparedness. Firm states they will need to keep this permit at this location.

II. DESCRIPTION OF BUSINESS:

Hours of Operation - 0700 - 1700. Monday through Friday

Approximate size of facility - 7000 square feet

Number of employees - 10 -12

Any other business operating at this address? Firm states permit 54:258 will continue to operate from this location.

The firm states the Emergency Preparedness permit is used in order for the facility to store Rx drugs to only be distributed in the event of an in State emergency. Example Cipro for Anthrax, Tamiflu for Influenza outbreaks.

The firm states this permit will also be used in order to distribute Rx drugs to State Prisons.

Any off-site storage and/or warehouses? The firm stores drugs at other permitted locations.

Explain Day to Day Operations

Firm receives Rx drugs from Cardinal. The firm states the majority of the Rx drugs are paid for by the individual prisons and are shipped to this location from Cardinal to be repackaged at this location. Once repackaged the firm will distribute the repackaged Rx drugs to the prisons. The drugs that are intended to be repackaged for the prison is paid for by the prison and shipped to DOH for repackaging. DOH does not purchase these drugs.

The firm states the only reimbursement they receive for the repackaged drugs is the cost of the repackaging.

The firm states they do purchase some Rx drugs from Cardinal pursuant to the STD specialty care program. These drugs are distributed to the DOH Pharmacy located 116 A Hamilton Park Drive. Once received by the pharmacy these drugs are dispensed by the pharmacy to the prison on a patient specific basis.

The Department is reimbursed by the prisons for the STD specialty care program drugs. The firm states the Rx drugs purchased for the STD specialty care program are purchased using their 54 permit.

The firm creates a batch production and control record for each drug that is repackaged. The records provides instructions on how that particular drug is to be repackaged. Also includes elements such as name of drug, strength of drug, lot number and expiration dates.

III. PRODUCT REGISTRATION - F.S. 499.015 and Rule 61N-1.016 FAC

The firm has applied for a product registration permit for this location. The firm previously had a product registration permit for 104-2 Hamilton Park drive

November 19, 2019 at 1:05:58 PM EST
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The firm provided a list of products they have previously registered for review. The list indicates the firm has registered 116 products

IV. CONTROLLED SUBSTANCE REPORTING

A. Has firm registered with the Department's CSR system? The firm states they will not repackage or distribute any controlled substances from this location with either the Restricted Distributor and/or the Repackager permit.

V. VENDOR LIST -

Firm states their only vendor is Cardinal Health. A review of the invoices for 104-2 Hamilton Park Drive indicates the drugs are being received from Cardinal Health, 2045 Interstate Drive. Lakeland, FL Permit 22:913

VI. CUSTOMER LIST - All State correctional facilities. The firm states they distribute to (5) facilities. The firm states these (5) facilities will then further distribute to the rest of the State correctional facilities.

VII. INBOUND DOCUMENTS - Reviewed Cardinal Invoices from 104-2 Hamilton Park Drive. Firm will continue to receive from Cardinal once they have relocated.

VIII. OUTBOUND DOCUMENTS

A. Outbound Business Record Template Provided? Yes

IX. SECURITY - Sonitrol

Firm has Indoor/Outdoor cameras that monitor entry and exit points.
Firm has door badge scanners for entry into building.

X. Equipment

The firm has (3) Autobond repacking machines. The firm states these are used for repackaging smaller quantities. (1) of the Autobonds is used to repackage Penicillin only.

(4) 400 repackaging machines

(1) 500 repackaging machines which is for the largest quantities.

Firm has a self contained clean room located at 104-2 Hamilton Park Drive which has not been relocated. Firm states they no longer provide beta lactam drugs to DOC as such; the firm will not be relocating the clean room.

A) Description

B) Maintenance Performed by? Omnicell

C) Calibration Performed by? Omnicell. Firm states the machines are calibrated annually.

D) Firm states clean room is certified by EOC-1

D) Cleaning Schedule. - Firm states the repacking machines are cleaned every time they change drugs that are being repackaged, as such; the firm states these may be cleaned multiple times daily. The firm states at a minimum the machines will be cleaned once daily.

Firm states they clean with isopropyl alcohol.

November 19, 2019 at 1:05:58 PM EST
Location:
License #: 2829
Inspector: Donald Yerbey

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XI. Temperature/Humidity Monitoring:

A) Room Air Monitored? Number/Location of Sensors/Thermometers

Firm has (1) electronic temperature and humidity monitors installed. Firm states the electronic monitor cards are changed monthly and stored onsite.
The firm has an additional electronic temp/humidity monitor at 104-2 Hamilton Park Drive that they intend to move to this location.

B) Refrigerated Present? No, Firm states they will not stock, repackage and/or distribute any refrigerated drugs.

C) Freezer Present? No, Firm states they will not stock, repackage and/or distribute any refrigerated drugs.

The firm does not have a refrigerator or freezer present at this time.

XII. Policies and Procedures:

The firm states they have P & P's for storage and handling of Rx drugs in their Pharmacy located at 116 A Hamilton Park Drive, however; they state they do not have written P & P's for this location. The firm states the Rx drugs are repackaged and distributed upon receipt, as such; the firm states they do not have P & P's for rotating stock, checking for expiration dates etc.

XIII. MISCELLANEOUS

A. How are Rx Drugs delivered to customers? FedEx and UPS

B. Destructions/Returns handled by? Inmar.

C. Does firm have a copy of FS 499 and Rule 61N-1 FAC onsite? Emailed to facility by inspector

XIV. TOUR OF FACILITY

Firm has a 2 bay doors in the receiving area where drugs are received from Cardinal. Once the firm has verified the contents of the totes from Cardinal the drugs will be transferred to the prep room where they are transferred from the stock bottle into labeled plastic bags. The bags indicate pertinent drug information, i.e. Name of drug, strength, lot # expiration dates etc. from there the drugs are transferred to the actual area where they will be repackaged. After the drugs are repackaged they are transferred to QC (quality control area) where they are checked for accuracy. The drugs are then boxed and ready for shipment.

XV. EXIT INTERVIEW

1) Requested the firm to relocate the 2nd temp/humidity monitor as from 104-2 Hamilton Park Drive to 104-3 as quickly as possible.

email picture to inspector when relocated.

November 19, 2019 at 1:05:58 PM EST
Location:
License #: 2829
Inspector: Donald Yarbey

Page: 6 of 7



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A handwritten signature in black ink, appearing to read "Danyelle Williams".

Licensee or Owner Signature

Danyelle Williams
Interim Bureau Chief
Nov 19, 2019 13:05

A handwritten signature in black ink, appearing to read "Don Yerbey".

Inspector Signature

Donald Yerbey
Drug Inspector
Nov 19, 2019 13:05

November 19, 2019 at 1:05:58 PM EST
Location:
License #: 2829
Inspector: Donald Yerbey

Page: 7 of 7

Update 11/11/21, Page 61

FDA-FLORIDA-004515

Mission:

To protect, promote & improve the health of all people in Florida through integrated state, county & community efforts.



Ron DeSantis
Governor

Scott A. Rivkees, MD
State Surgeon General

Vision: To be the Healthiest State in the Nation

November 18, 2020

Center for Drug Evaluation and Research
Office of Communications
10001 New Hampshire Avenue
Hillandale Building, 4th Floor
Silver Spring, MD 20993

The Florida Department of Health (DOH) attests, to the best of our knowledge, that there are no past criminal convictions or violations of State, Federal, or Canadian laws regarding drugs or devices against or by the responsible individual(s). Further, DOH attests that the responsible individual(s) have not been involved in, or convicted of, any such violations and there have been no disciplinary actions against the responsible individual(s).

The DOH's Pharmacists are Danyelle Brionne Williams, Pharmacist and Consultant Pharmacist License Number PS53818 and Niaz Ahmed Siddiqui, Pharmacist and Consultant Pharmacist License Number PS42647.

Please contact Doug Woodlief, (850) 245-4230 if you have any questions regarding this attestation.

Sincerely,

Doug Woodlief
Division Director
Emergency Preparedness and Community Support

DHW/vj

Florida Department of Health
Division of Emergency Preparedness and Community Support
4052 Bald Cypress Way, Bin A-22 • Tallahassee, FL 32399
PHONE: 850/245-4864 • FAX: 850/921-8162
FloridaHealth.gov



Accredited Health Department
Public Health Accreditation Board

Attachment B

LifeScience Attestations and Inspectional History



Solicitation Number: AHCA ITN 008-19/20
Title: Canadian Prescription Drug Importation Program (CPDIP)

2600 Regent Blvd.
DFW Airport, TX 75261

Telephone: (469) 844-3701
Fax: (972) 456-1832

November 17, 2020
Center for Drug Evaluation and Research
Office of Communications
10001 New Hampshire Avenue
Hillandale Building, 4th Floor
Silver Spring, MD 20993

LifeScience Logistics (LSL) attests that there are no past criminal convictions or violations of State, Federal, or Canadian laws regarding drugs or devices against or by the responsible individual(s), Foreign Seller, or Importer. Further, LSL attests that the responsible individual(s), Foreign Seller or Importer has not been involved in, or convicted of, any such violations. LSL's principal owners (owning 10 percent or greater of outstanding stock) are Richard Beeny and Max Kamhi.

Lastly, there have been no disciplinary actions against the responsible individual(s), Foreign Seller, or Importer by State, Federal, or Canadian regulatory bodies, including any such actions against the principals, owners, directors, officers, quality unit, or any facility manager or designated representative of such manager for the previous 7 years prior to submission of the SIP Proposal, with the exception of the following three administrative fines to LSL from the referenced bodies (continued on Page 2):

- - -



Solicitation Number: AHCA ITN 008-19/20
Title: Canadian Prescription Drug Importation Program (CPDIP)

Detail	Comments
Case # 2016-3687	Colorado State Board of Pharmacy issued an Administrative fine of \$1500 on 19 Jul 2016 for having a change of DR notice outside of the 30-day reporting window for LifeScience Logistics, 1105 E Northfield Drive, Suite 300-400, Brownsburg, IN 46112. This reporting delay occurred in November 2015.
Citation CI 2017 78097	California State Board of Pharmacy issued an Administrative fine of \$900 on 30 Apr 2018 for having a change of DR notice outside of the 30-day reporting window for LifeScience Logistics LLC, 4475 South Fulton Parkway Building 5 Suite C, Atlanta, GA 30349. This reporting delay occurred in September 2016.
Case # 18-L-0075	Alabama State Board of Pharmacy issued a reciprocal administrative fine of \$1000 on 29 Oct 2018 for the same incident in relation to the California State Board of Pharmacy Citation CI 2017 78097.

Warm regards,

Richard Beery
CEO and Co-founder

- - -



**Life Science Logistics, LLC Five Year Regulatory Audit History
(FEI# 968727565)**

17 FEB 2016 - DEA Import/Export - no observations

26 FEB 2016 - Indiana Dept. of Environmental Management - no observations

31 MAR 2016 - ISO 13485:2003 - no observations – ISO Certificate issued

22 AUG 2016 - Verified-Accredited Wholesale Distributors (VAWD) - no observations – Renewed

23 SEP 2016 - Indiana Board of Pharmacy – Pass, no observations

14-16 FEB 2017 -ISO 13485:2003 one-year surveillance audit - 2 minor observations (closed)

14 MAR 2017 - FDA - no observations

25 AUG 2017 - DEA Export - 1 minor observation (closed)

07 MAR 2018 - ISO 13485:2003 one-year surveillance audit - 1 minor observation (closed)

08 MAR 2018 - ISO 13485 transition audit to 2016 standard – ISO 13485:2016 Certificate issued

14 JUN 2018 - DEA Import and Distributor - 3 minor observations (closed)

21 FEB 2019 - ISO 13485:2016 commercial recertification audit – 1 minor observation (closed)


23 May 2019 - Verified-Accredited Wholesale Distributors (VAWD) - no observations – Renewed

14 Jan 2020 - ISO 13485: 2016 corporate one-year surveillance audit – no observations

21 JAN 2020 - ISO 13485: 2016 commercial one-year surveillance audit – no observations

24 APR 2020 - DEA Export – no observations

26 JUN 2020 - FDA Risk Evaluation & Mitigation Strategies program (REMS) – no observations - Certified

 14 Nov 2020
Paul Hayward
Director, Quality Assurance and Regulatory Affairs



**Life Science Logistics, LLC Seven Year Disciplinary History
(FEI# 968727565)**

There have been no disciplinary actions against the responsible individual(s) or Importer by State, Federal, or Canadian regulatory bodies, including any such actions against the principals, owners, directors, officers, quality unit, or any facility manager or designated representative of such manager for the previous 7 years prior to submission of the SIP Proposal, with the exception of the following three administrative fines to the entire Life Science Logistics facility network from the referenced state bodies:

Case # 2016-3687	Colorado State Board of Pharmacy issued an Administrative fine of \$1500 on 19 Jul 2016 for having a change of DR notice outside of the 30-day reporting window for LifeScience Logistics, 1105 E Northfield Drive, Suite 300-400, Brownsburg, IN 46112. This reporting delay occurred in November 2015.
Citation CI 2017 78097	California State Board of Pharmacy issued an Administrative fine of \$900 on 30 Apr 2018 for having a change of DR notice outside of the 30-day reporting window for LifeScience Logistics LLC, 4475 South Fulton Parkway Building 5 Suite C, Atlanta, GA 30349. This reporting delay occurred in September 2016.
Case # 18-L-0075	Alabama State Board of Pharmacy issued a reciprocal administrative fine of \$1000 on 29 Oct 2018 for the same incident in relation to the California State Board of Pharmacy Citation CI 2017 78097.

 14 Nov 2020
 Paul Hayward
 Director, Quality Assurance and Regulatory Affairs

Attachment C

LifeScience Logistics Inspection Records

Drug Establishments Current Registration Site

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[TWEET](#)
[in LINKEDIN](#)
[PIN IT](#)
[EMAIL](#)
[PRINT](#)
[New Search](#)

Search Results for **life science logistics**

[CSV](#)
[Excel](#)

Filter:

Firm Name ▲	FDA Establishment Identifier ◆	DUNS ◆	Business Operations ◆	Address ◆	Expiration Date ◆
Life Science Logistics	3018161030	117963809	RELABEL; REPACK;	3860 S 500 E Suite 200, Whitestown, Indiana (IN) 46075, United States (USA)	12/31/2021



Indiana Department of Environmental Management

We Protect Hoosiers and Our Environment.

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Michael R. Pence
Governor

Carol S. Comer
Commissioner

March 2, 2016

VIA E-MAIL

Mr. Christoph Erdel
Life Science Logistics LLC
1105 E. Northfield Dr.
Brownsburg, IN 46112

Re: Inspection Summary Letter
Life Science Logistics LLC
INR000137521
Brownsburg, Hendricks County

Dear Mr. Erdel:

On 2/26/2016, a representative of the Indiana Department of Environmental Management, Office of Land Quality, conducted an inspection of Life Science Logistics LLC, located at 1105 E. Northfield Dr., Brownsburg, IN. This inspection was conducted pursuant to IC 13-14-2-2. For your information, and in accordance with IC 13-14-5, a summary of the inspection is provided below:

Type of Inspection: Compliance Evaluation Inspection
Multi-Media Screening Evaluation

Results of Inspection: No Violation(s) Discovered

Please direct any response to this letter and any questions to me at (317) 417-7891.

Sincerely,

A handwritten signature in black ink that reads 'Debbie Chesterson'.

Debbie Chesterson
Hazardous Waste Compliance Section
Compliance and Response Branch

Enclosure

cc: Hendricks County Health Department





**INDUSTRIAL/HAZARDOUS
WASTE INSPECTION REPORT**
INDIANA DEPARTMENT OF
ENVIRONMENTAL MANAGEMENT

Inspector's Name:	Debbie Chesterson
Others Present	
Date:	Friday, February 26, 2016
Time In:	8:50 AM
Time Out:	10:00 AM
Inspection Type	Compliance Evaluation Inspection Multi-Media Screening Evaluation

General Information						
Facility Information						
Facility Name	Life Science Logistics LLC					
Facility Location	1105 E. Northfield Dr. Brownsburg, IN 46112 Hendricks County					
Facility Mailing Information	Same Address as Facility					
Facility Contact	Same as Primary Facility Contact					
Primary Facility Contact During Inspection	Christoph Erdel Operations Supervisor 317-456-0257 cerdel@lslog.com					
Other Facility Contact(s) During Inspection	Salutation	First Name	Last Name	Title	Phone Number	Email
	Ms.	Heather	Laurin	QA Specialist	317-489-5036	hlaurin@lslog.com

Facility ID			
EPA ID Number	INR000137521	NAICS Code	49311, 424210

Facility Status			
File Status	Conditionally Exempt Small Quantity Hazardous Waste Generator	Other Activities	

Outstanding Issues	
Last Inspection Date	No prior RCRA inspections
Previous Violations	<input type="radio"/> Yes <input checked="" type="radio"/> No
Details	

Inspection Narrative
Life Science Logistics LLC operates as a third party logistics provider for the healthcare industry. The distribution warehouse stores pharmaceuticals, medical devices and over-the-counter (OTC) items for over twenty clients in Suite 300. Suite 200 is currently unoccupied. Suite 400 warehouses healthcare items for the federal government. They began operations at this location in 2011.
This was the first RCRA inspection at the facility. They began operations at this location in 2011. The facility is notified as a conditionally exempt small quantity generator of hazardous waste. To date, no hazardous waste has been stored or shipped from this location. Hazardous waste generation could occur through damaged or expired products that are stored within the facility; hence they obtained an EPA identification number for the site.

Debbie Chesterson

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Life Science Logistics LLC/Friday, February 26, 2016

No violations were noted during the inspection.

Regulatory Status

Observed Activity	Conditionally Exempt Small Quantity Generator	Other Activities	Universal Waste Handler
Documents Reviewed	Manifests		
Comments			

Waste Management

Comments:

Waste Stream(s) Information

Waste Streams
☒ Yes ☐ No ☐ Not Inspected ☐ Not Applicable

List waste stream(s) information that varies from the most recent Annual Report (Example: additional waste streams, waste streams no longer generated, significant increase/decrease in generation rate, etc.)

EPA Waste Codes	Description	Source	Generation Rate	Disposition
No hazardous waste generation to date				

Exempted/Excluded ☐ Yes ☒ No ☐ Not Inspected ☐ Not Applicable

Explanation

Waste Management Areas

Container Management Area(s) ☒ Yes ☐ No ☐ Not inspected ☐ Not applicable

EPA Waste Codes	Location	Number	Size	Type of Container
No hazardous waste present				

Satellite Area(s) ☐ Yes ☒ No ☐ Not inspected ☐ Not applicable

Tanks, Restricted Waste Sites, and Other Regulated Units
☐ Yes ☒ No ☐ Not inspected ☐ Not applicable

Environmental Releases

Visible Releases/Contamination/Discharges	<input type="radio"/> Yes <input checked="" type="radio"/> No Release Observed
---	--

Compliance Assistance

P2 Information

The following P2 suggestions could possibly save money, reduce waste and/or minimize risk. You might consider having a P2 assessment, or a voluntary technical assistance consultation from IDEM staff. Please visit the agency's P2 web site at <http://www.in.gov/idem/5298.htm> for additional information.

Contact by IDEM OPPTA Requested	<input type="radio"/> Yes <input checked="" type="radio"/> No
P2 Suggestions	

Guidance Materials

Debbie Chesterson

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Life Science Logistics LLC/Friday, February 26, 2016

Guidance Materials Provided to Facility	
--	--

Multi-Media Screening (Checked box indicates a concern)	
Comments:	
Multi-Media Screening Conducted	<input checked="" type="radio"/> Yes <input type="radio"/> No
Water Concerns <input type="checkbox"/> Process wastewater discharge to a POTW collection system (i.e. sewer) without a permit <input type="checkbox"/> Direct discharge (from industrial process, industrial wastewater treatment or non-contact cooling water) to receiving water near the facility without an NPDES permit <input type="checkbox"/> Process materials such as cleaners, solvents, paints, lubricants, etc. are escaping through floor drains	Air Concerns <input type="checkbox"/> Visible emissions from stacks or vents <input type="checkbox"/> Dust crossing property lines <input type="checkbox"/> Open solvent containers <input type="checkbox"/> Spray booth filters not securely in place <input type="checkbox"/> Open burning
Storm Water Concerns <input type="checkbox"/> NOI for Rule 6 (see applicable SIC codes) <input type="checkbox"/> Storm Water Pollution Prevention Plan (must be developed within 365 days of NOI). <input type="checkbox"/> Storm water annual sample <input type="checkbox"/> Measures to ensure contaminants from industrial activities aren't exposed to storm water <input type="checkbox"/> Documented quarterly inspections of storm water run-off conveyances <input type="checkbox"/> Documented annual employee training on SWP3 <input type="checkbox"/> Rule 5 storm water permit for land disturbing activities greater than 1 acre <input type="checkbox"/> Signs of erosion or off-site sedimentation into waters of the state from construction sites	Drinking Water Concerns <input type="checkbox"/> PWSID# (applies to 25 or more employees on self-supplied drinking water system) <input type="checkbox"/> Contamination within 200 ft of wellhead TRI Concerns <input type="checkbox"/> Lack of TRI report (applies to 10 or more employees in applicable SIC codes) UST Concerns <input type="checkbox"/> Unregistered UST containing petroleum or hazardous substance

Checklist (Checked box indicates a compliance concern)	
Standards <input type="checkbox"/> Hazardous Waste Determination <input type="checkbox"/> EPA Identification Number(s) <input type="checkbox"/> Manifest General Requirements <input type="checkbox"/> Use of the Manifest <input type="checkbox"/> Biennial Report	Satellite Accumulation – SQG and LQG <input type="checkbox"/> Satellite Accumulation - 55 Gallon <input type="checkbox"/> Satellite Accumulation - Label <input type="checkbox"/> Satellite Containers Closed
Container Management – LQG <input type="checkbox"/> Accumulated On-site for 90 Days or Less <input type="checkbox"/> Hazardous Waste Container Condition <input type="checkbox"/> Hazardous Waste Container Compatibility <input type="checkbox"/> Hazardous Waste Containers Closed <input type="checkbox"/> Hazardous Waste Container Handling	Container Management – SQG <input type="checkbox"/> Accumulate for 180 Days or Less <input type="checkbox"/> May not Exceed 6000 Kg (13,200 Lbs) <input type="checkbox"/> Hazardous Waste Container Condition <input type="checkbox"/> Hazardous Waste Container Compatibility <input type="checkbox"/> Hazardous Waste Containers Closed

Debbie Chesterson

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Life Science Logistics LLC/Friday, February 26, 2016

<input type="checkbox"/> Hazardous Waste Container Inspections <input type="checkbox"/> Accumulation Start Date Clearly Marked and Visible <input type="checkbox"/> Marked Clearly with Words "Hazardous Waste"	<input type="checkbox"/> Hazardous Waste Container Handling <input type="checkbox"/> Hazardous Waste Container Inspection <input type="checkbox"/> Accumulation Start Date Clearly Marked and Visible <input type="checkbox"/> Marked Clearly with the Words "Hazardous Waste"
Preparedness and Prevention – LQG and SQG <input type="checkbox"/> Maintained and Operated to Minimize Possibility of a Release <input type="checkbox"/> Required Equipment <input type="checkbox"/> Communication & Alarm Access <input type="checkbox"/> Aisle Space	Contingency Plan and Emergency Procedures – LQG <input type="checkbox"/> Contingency Plan Developed <input type="checkbox"/> Contingency Plan Content <input type="checkbox"/> Contingency Plan Maintained at Facility
Personnel Training – LQG <input type="checkbox"/> Personnel Training	Training and Emergency Procedures – SQG <input type="checkbox"/> SQG Emergency Coordinator <input type="checkbox"/> Emergency Information Posted <input type="checkbox"/> Employee Training
Tank Requirements – LQG <input type="checkbox"/> Integrity Assessment <input type="checkbox"/> Containment and Release Detection <input type="checkbox"/> Tank General Requirements <input type="checkbox"/> Tank Inspections <input type="checkbox"/> Subpart BB - Monthly Pump and Valve Monitoring <input type="checkbox"/> Subpart CC - Annual Inspection/Monitoring	Used Oil – All Facilities <input type="checkbox"/> Rebuttable Presumption Applies <input type="checkbox"/> Containers and Tanks in Good Condition <input type="checkbox"/> Containers/Tank Labeling <input type="checkbox"/> Release Clean Up and Containment <input type="checkbox"/> Burning Restrictions - Generated On-site or Dly, .5M BTU
Additional Requirements – LQG and SQG <input type="checkbox"/> Release to the Environment, Disposal of Solid Waste <input type="checkbox"/> Illegal Dumping <input type="checkbox"/> Land-Ban Notification <input type="checkbox"/> Other Violation	Universal Waste – All Facilities <input type="checkbox"/> Universal Waste Labeling <input type="checkbox"/> Containers - Closed, Good Condition, No Evidence of Leaks <input type="checkbox"/> Universal Waste - Bulb Crushing Prohibition

Description of Violation(s)

Inspection Documentation	
Photographs	<input type="radio"/> Yes <input checked="" type="radio"/> No
Map	<input checked="" type="radio"/> Maps
GPS Location Collected	<input type="radio"/> Yes <input checked="" type="radio"/> No
Analytical Screening Conducted	<input type="radio"/> Yes <input checked="" type="radio"/> No

Debbie Chesterson

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Life Science Logistics LLC/Friday, February 26, 2016

Lab Sample	<input type="radio"/> Yes	
	<input checked="" type="radio"/> No	

Inspection Results/Actions	
Comments:	
Inspection Results	
No Violation(s) Discovered	
Multi-Media Screening Results	
No violations	

Finalize Inspection		
Written Summary of Inspection	Notice of Inspection and Verbal Summary Provided	
Inspector Information	Printed/Typed Name	Debbie Chesterson
	Phone Number:	(317) 417-7891
	Email Address:	djcheste@idem.in.gov
	Signature:	Signature obtained on the Notice of Inspection
Facility Representative Signature	Printed/Typed Name:	Christoph Erdel
	Signature:	Signature obtained on the Notice of Inspection


NOTICE OF INSPECTION
 State Form 50890 (R3 / 11-05)

INDIANA DEPARTMENT OF ENVIRONMENTAL MANAGEMENT
 100 N. Senate Avenue
 Indianapolis, IN 46204-2251
 Telephone: (800) 451-6027 or (317) 232-8603

 This is to notify you that on 2/26/16 an inspection of Life Science Logistics LLC
 was conducted by the undersigned representative of the Indiana Department
 of Environmental Management (IDEM), Office of Land Quality
Type of Inspection (may include more than one):

- ☒ Compliance Evaluation Inspection - Hazardous Waste
☐ Complaint
☒ Multi-Media Screening Evaluation
☐ Other

Preliminary Inspection/Screening Findings:

These findings are considered preliminary and identify specific compliance issues discovered during the above-noted inspection that the designated agent of IDEM believes may be a violation of a statute(s), rule(s) or permit(s) issued by IDEM.

Single Media Inspection:

- ☒ No violations were discovered with respect to the particular items observed during the inspection.
☐ Violations were discovered but corrected during the inspection.
☐ Violations were discovered and require a submittal from you and/or follow-up inspection by IDEM.
☐ Violations were discovered and may subject you to an appropriate enforcement response.
☐ Additional information/review is required to evaluate overall compliance.
☐ Other / Comments (attachment may be included)

Multi-Media Screening (Please note that a multi-media screening is not a comprehensive evaluation of the compliance status of the facility):

- ☐ Multi-media screening not conducted.
☒ No violations were discovered with respect to the limited multi-media screening conducted by IDEM.
☐ Potential violations were discovered but corrected during the inspection.
☐ Potential violations were discovered and may be further investigated.

Pollution Prevention:

Pollution prevention is the preferred means of environmental protection in Indiana. The goal of pollution prevention is to promote changes in business and commercial operation, especially manufacturing processes, so that Indiana businesses increase productivity, generate less environmental wastes, reduce their regulatory responsibilities and become more profitable. Your participation in Indiana's pollution prevention program is entirely voluntary. If you have any pollution prevention questions, you may contact our Office of Pollution Prevention and Technical Assistance (OPPTA) at (317) 232-8172 or (800) 988-7901, or visit OPPTA's Web site at www.idem.IN.gov/oppta/p2/. Would your company like to be contacted by IDEM's Office of Pollution Prevention and Technical Assistance? ☐ Yes ☐ No

Compliance Assistance:

In addition to the compliance assistance offered by IDEM's individual programs, IDEM's Compliance and Technical Assistance Program (CTAP) offers free, confidential compliance assistance to regulated entities, including small businesses and municipalities, throughout Indiana. In the future, if you would like to request free, confidential compliance assistance, call (317) 232-8172 or (800) 988-7901, or visit CTAP's Web site at www.idem.IN.gov/ctap.

A summary of violations and concerns noted during the inspection was verbally communicated to the undersigned representative during the inspection. The facility should correct any violations noted as soon as possible. Violations identified and corrected during the inspection may still be cited as violations.

A written inspection summary will be provided within 45 days. In accordance with IC 13-14-5-4, matters not evident to DEM at the time of the inspection might not be included in either the verbal or written inspection summary.

DEM Representative:

Printed Name	Signature	Phone Number	Date	In: Time	Out: Time
Debbie Chesterson	<i>[Signature]</i>	317-417-7891	2/26/16	8:50 am	10:00 am

Owner/Agent Representative:

Printed Name	Signature	Title	Phone Number	Date
CHRISTOPH F. ERDEL	<i>[Signature]</i>	Operations Supervisor	317 456 0357	26 FEB 2016

DISTRIBUTION: White - IDEM Public File; Carbon - Office of Pollution Prevention and Technical Assistance



May 25, 2017

Cory Green, Senior Manager
Commercial Operations
Lifescience Logistics
1105 E. Northfield Drive, Suite 400
Brownsburg, IN 46112-2530

FMD-6981-17

Dear Mr. Green:

The U.S. Food and Drug Administration (FDA) or a state agency contracted by the FDA, conducted an inspection at 1105 E. Northfield Drive, Suite 400, Brownsburg, IN, ending on March 14, 2017. Effective April 1, 1997, when the Agency determines an inspection is closed under 21 C.F.R. 20.64 (d)(3), FDA released a copy of the inspection report to the inspected firm.

You will find a copy of the FDA Establishment Inspection Report or state contracted inspection report attached, FDA may have redacted some information in accordance with the Freedom of Information Act (FOIA) and Title 21, C. F. R., Part 20. Firms may request a copy of their FDA inspections completed prior to April 1, 1997 through FOIA.

FDA is working to make its regulatory process and activities more transparent to the regulated industry. Part of this effort is releasing a copy of your inspection report or summary to you.

Any questions regarding this letter or the release of this report should be directed to FMD Coordinator, U.S. Food and Drug Administration, 300 River Place, Suite 5900, Detroit, Michigan 48207. Telephone 313-393-8110; fax 313-393-8139.

Sincerely,

A handwritten signature in black ink, appearing to read "Kelli L. Wilkinson".

LCDR Kelli L. Wilkinson
Director of Compliance
Detroit District Office

Enclosure: EIR,
ded

U.S. Food and Drug Administration – Detroit District
300 River Place, Suite 5900
Detroit, MI 48207
Telephone: 313-393-8100
FAX: 313-393-8139
www.fda.gov

Establishment Inspection Report
 Lifescience Logistics
 Brownsburg, IN 46112-2530

FEL: 3009057691
 EI Start: 3/14/2017
 EI End: 3/14/2017

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SUMMARY

This was a pre-approval inspection (PAI) of a large contract finished drug storage and distribution warehouse. This firm was identified as a warehousing and distribution facility [REDACTED]

USA Inc.

This inspection was performed in accordance with compliance program 7356.002 *Drug Manufacturing Inspections* as well as 7346.832 *Pre-Approval Inspections/Investigations* and is being reported under FACTS assignment number 11715051.

The firm was previously inspected on 8/21/12 under the biologics program. No inspection observations were issued at the conclusion of the August 2012 inspection.

The current inspection was the firm's first inspection under the drug program and included coverage of the firm's Quality, Facilities & Equipment, and Materials systems. No inspection observations were issued at the conclusion of this inspection.

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 Lifescience Logistics
 Brownsburg, IN 46112-2530

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No refusals were encountered and no samples were collected.

ADMINISTRATIVE DATA

Inspected firm: Lifescience Logistics
 Location: 1105 E Northfield Dr Ste 300-400
 Brownsburg, IN 46112-2530
 Phone: 317-456-0254
 FAX: -
 Mailing address: 1105 E Northfield Dr Ste 300-400
 Brownsburg, IN 46112-2530
 Dates of inspection: 3/14/2017
 Days in the facility: 1
 Participants: Robert M Barbosa, Investigator

Upon arrival I, Investigator Robert M. Barbosa, presented credentials and a prepared FDA 482 Notice of Inspection to Mr. Derek Gates (Supervisor of Operations). Mr. Gates identified himself as the most responsible onsite and accepted the notice. Also in attendance was Mr. Henry Tillman (QA Specialist) and Mr. David Mastromatteo (COO). I relayed the purpose and scope of my inspection to the aforementioned individuals.

An inspection closeout meeting was held with Mr. Tillman and Mr. Mastromatteo later that day. No observations were issued at the conclusion of the inspection.

HISTORY

LifeScience Logistics (abbreviated as LSL) operates as a contract domestic warehousing and distribution facility of finished pharmaceuticals and other regulated products including biotech, medical devices, and biologics. The firm operates four facilities located proximal to Dallas-FW, TX; Baltimore, MD; Indianapolis, IN; and Atlanta, GA. The Indiana site specifically provides contract warehousing and distribution of finished pharmaceuticals (including category I-V) and biologics (animal plasma). The firm also provides storage of finished pharmaceuticals (including category I-V) for [REDACTED]. In addition to controlled room temperature storage (CRT), the facility is equipped for storage of refrigerated (2-8°C) and frozen (-25 to -35°C) products.

The facility was last inspected under the biologics program on 8/21/12. No observations were issued at the conclusion of that inspection.

Please direct all post inspectional correspondence, including FMD-145 to:

Mr. Cory Green,

Establishment Inspection Report

Lifescience Logistics

Brownsburg, IN 46112-2530

FEI: 3009057691

EI Start: 3/14/2017

EI End: 3/14/2017

Sr. Manager of Commercial Operations

1105 E. Northfield Dr., Suite 400

Brownsburg, IN 46112

INTERSTATE (I.S.) COMMERCE

Mr. Tillman stated that approximately 60-70% of all shipments into and out of the Brownsburg facility move in interstate commerce.

JURISDICTION (PRODUCTS MANUFACTURED AND/OR DISTRIBUTED)

The firm provides contract warehousing and distribution services of finished pharmaceuticals (including category I-V) as well as other regulated products including biotech, medical devices, and biologics (animal plasma). The firm also provides storage of finished pharmaceuticals (including category I-V) for [REDACTED] In addition to controlled room temperature storage (CRT), the facility is equipped for storage of refrigerated (2-8°C) and frozen (-25 to -35°C) products. All registrations were verified as current.

INDIVIDUAL RESPONSIBILITY AND PERSONS INTERVIEWED

Mr. Richard Beeny is the firm's current CEO. The site's commercial warehousing and distribution operation is led by Mr. Cory Green (Sr. Manager of Commercial Operations). Mr. Green is temporarily reporting to Mr. Mastromatteo until the vacant position of Operations Manager for the Brownsburg, IN site is filled.

Mr. Tillman is the head of the quality organization at the Brownsburg, IN site. Mr. Tillman reports directly to Ms. Samantha Nash (QA Supervisor). Ms. Nash in-turn reports to Mr. Paul Hayward (Director of Quality and Regulatory Affairs).

Mr. Mastromatteo, Mr. Tillman, and Ms. Clara Walker (QA Coordinator) accompanied me during my walkthrough of the facility. All general operation questions were addressed by Mr. Mastromatteo. All document requests were directed to and fulfilled by Mr. Tillman and Ms. Walker.

A copy of the firm's organizational chart is included here as **Exhibit RMB1**.

FIRM'S TRAINING PROGRAM

The firm's training program is defined in procedure no. SOP 1800 *Training and Qualification*. Training falls into three main categories; read-only, instructor led, and non-procedure training. Training is recorded on hard copy training forms and maintained in a training file. The firm's training program was not reviewed during this inspection.

MANUFACTURING/DESIGN OPERATIONS

Facility Overview:

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Brownsburg, IN 46112-2530

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The LSL facility is located in a large single concrete structure. The facility is divided into two suites identified as 300 and 400. Suite 300 supports the firm's regulated commercial product operation and includes approximately 350,000ft². Suite 400 supports [REDACTED] of the operation and was not evaluated during this inspection. Both suites are managed separately but share a common quality unit.

The firm maintains separate personnel for each suite and access is managed through card key for each area. The commercial product operation consists mainly of a large number of product storage pallet bays, an extensive receiving/shipping area, return/reject cage, a large walk-in refrigeration unit, and smaller upright units for frozen storage.

Quality System Review:

The firm has procedures in place for managing all deviations and change controls. All deviation investigations are performed per procedure number SOP 1350 *Deviation/CAPA-Commercial*. I reviewed a list of deviations initiated between January 1, 2017 and March 2017. Three deviation investigations were selected for review. I noted no issues during my review of these investigations. All reports were observed to have been completed per the procedure

All change controls are managed through procedure number SOP 1002 *Change Control*. I reviewed a list of change controls initiated between January 1, 2016 and March 2017. I reviewed change control number CR-INC-16-0031. I noted no issues during my review of this change control. The processes documented within the report were observed to have been completed per the procedural requirements.

Facilities & Equipment System Review:

As part of my review of the facilities and equipment system, I reviewed the firm's calibration and maintenance of the temperature reporting probes located throughout the ambient storage warehouse expansion area which was qualified for use in December 2016. Initial probe calibration was performed at that time along with the initial temperature mapping of the warehouse expansion area. I reviewed the recent probe calibration data as well as the temperature mapping data without note. Each of the temperature reporting probes were observed to be uniquely identified and within calibration tolerance. I also reviewed a trend of the temperature readings reported throughout the expanded warehouse storage area between December 2016 and March 2017. All reported temperature readings were within acceptable limits.

Materials System Review:

Review of the materials system focused mainly on material receipt and material control through the firm's inventory control system TECSYS Elite Series Warehouse Management System. All materials and material status is controlled using this validated inventory management system. The system is designed to interface with the firm's client's inventory systems as well (specifically [REDACTED]). Orders are transferred into the system electronically by the client and product is rotated

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Brownsburg, IN 46112-2530

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based on oldest expiry date for fulfillment of those orders. Staff will collect the appropriate products, package, and ship them to the designated customer. All products are received into the system in a "PR Hold" status. Product is released for use by quality assurance after notification by the client. Material cannot be allocated to fill orders unless it has been made available for use in the system.

The system validation was documented in a series of six validation documents. A copy of validation summary report no. CSV-WMS-007 was provided and reviewed. All test scripts for the system were included in the firm's IQ and PQ protocols. The summary report stated that the TECSYS system is hosted by the software owner which acts as a 3rd party for tech support and maintenance of inventory data. The software owner was audited as part of the validation effort. The summary report from this audit was provided and reviewed. The audit findings identified several corrective areas but overall no significant issues were noted.

MANUFACTURING CODES

The firm does not assign manufacturing codes to the finished drug products it receives or distributes out of its facilities. All manufacturing codes used are provided by the firm's clients and entered into the inventory tracking system. The manufacturing codes provided vary from client to client.

COMPLAINTS

Complaint investigations responsibilities for LSL are detailed in the firm's signed quality agreement with their client [REDACTED]. The agreement states that LSL will conduct investigations into any complaints due to "missing, overage or damaged product, late delivery etc."

Handling of product complaints is performed per procedure number SOP 4001 *Handling of Product Complaints*. The procedure describes the process for evaluating and investigating the complaints. Additionally all complaints are tracked and trends are reviewed periodically. All observed trends are investigated with CAPAs being initiated when determined to be appropriate.

I reviewed two complaint investigations as well as a recent trend investigation. I noted no issues during my review of these investigation forms.

RECALL PROCEDURES

Mr. Mastromatteo explained that as a contract service provider, LSL does not actively conduct product recalls and that all formal notifications regarding product recalls are issued by the firm's various clients. He did state however that LSL may provide assistance with product recalls per the client's request. This would include providing of distribution information as well as handling of recalled product returns and forwarding for destruction.

Any activities performed in support of a client's recall as well as any responsibilities which the client may require would be stated in writing and/or as part of a quality agreement with the client. The

Establishment Inspection Report

Lifescience Logistics

Brownsburg, IN 46112-2530

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product recall requirements between LSL and [REDACTED] were verified during my review of the firm's Quality agreement.

OBJECTIONABLE CONDITIONS AND MANAGEMENT'S RESPONSE

No inspection observations were issued.

REFUSALS

No refusals were encountered.

GENERAL DISCUSSION WITH MANAGEMENT

None.

ADDITIONAL INFORMATION

None.

SAMPLES COLLECTED

No samples were collected.

VOLUNTARY CORRECTIONS

N/A

EXHIBITS COLLECTED

1 Organizational Chart, 16 pages

ATTACHMENTS

1 FDA 482, 3 pages

Robert M.
Barbosa -S

Digitally signed by Robert M.
Barbosa -S
DN: cn=US, ou=U.S. Government,
ou=HHS, ou=FDA, ou=People,
o=9234219200300.100.1.1-200036
7409, cn=Robert M. Barbosa -S
Date: 2017.05.22 12:08:10 -04'00'



AMCF061(A1252)

**Acceptance of Corrective Action Plan
Revision 2 (March 2013)**

01AUG2017

Paul Hayward
LifeScience Logistics
1105 East Northfield Drive
Suite 300
Brownsburg
Indiana

Dear Mr Hayward,

Acceptance of Corrective Action Plan from ISO 13485:2003

Report No.: 8515524

Thank you for providing your corrective action plan, detailing your actions to resolve the nonconformities raised during the recent audit.

I can confirm that I have now had an opportunity to review and accept the correction, associated root cause analysis and the corrective action, and the actions and timescales specified appear to be appropriate for all nonconformities identified in the audit.

Verification of effectiveness of the nonconformities will be performed at your next scheduled surveillance audit, and the duration of this visit may be increased to allow time for the verification. Any additional time requirements will be confirmed prior to the visit.

If you have any queries on the above, please contact me.

Yours Faithfully,

J Gregory Jones
AVP Healthcare 2



April 26, 2018
Paul Hayward
LifeScience Logistics, LLC
1105 East Northfield Dr
Brownsburg,
Indiana
46112

Dear Paul,

Acceptance of Corrective Action Plan from Report No. 8619861

Thank you for providing your corrective action plan, detailing your actions to resolve the nonconformities raised during the recent audit.

I can confirm that I have now had an opportunity to review and accept the correction, associated root cause analysis and the corrective action, and the actions and timescales specified appear to be appropriate for all nonconformities identified in the audit.

Verification of effectiveness of the nonconformities will be performed at your next scheduled surveillance audit, and the duration of this visit may be increased to allow time for the verification. Any additional time requirements will be confirmed prior to the visit.

If you have any queries on the above, please contact me.

Yours Faithfully,

Jeremy Ward
BSI Client Manager
jeremy.ward@bsigroup.com
571-353-4914



NABP
National Association of
Boards of Pharmacy
www.nabp.pharmacy

1600 Feehanville Drive
Mount Prospect, IL 60056
T) 847/391-4406
F) 847/375-1114

June 4, 2019

Stephen Spelman
LifeScience Logistics, LLC
dba LifeScience Logistics
1105 E Northfield Drive
Brownsburg, IN 46112

Dear Mr Spelman:

On behalf of the National Association of Boards of Pharmacy® (NABP®) and Verified-Accredited Wholesale Distributors® (VAWD®) staff, I would like to take this opportunity to thank you and the team members at your facility for the hospitality extended to Rich Paul during the recent VAWD reaccreditation survey.

Information provided by the surveyor indicates your facility is operating in compliance with program criteria. NABP will conduct a final assessment of all reaccreditation materials, survey findings, and responses to confirm if your facility continues to meet VAWD program criteria. The Accreditation Committee will then render a decision on reaccreditation. NABP would like to assure you that we will move through the process as expediently as possible.

Thank you for your continued support of the VAWD program and for the courtesy you and your staff have extended during the process. If you have questions or concerns, feel free to contact VAWD staff via email at vawd@nabp.pharmacy.

Sincerely,

Dawn Bibbs-Morrissey
Accreditation Manager

Pharmacy
402 W. Washington Street
Indiana Government Center South, Room W072
Indianapolis, IN 46204
317-234-2067(PHONE) • 317-233-4236(FAX)

FACILITY

LifeScience Logistics
 1105 East Northfield Drive Suite 300-400
 Brownsburg IN 46112
 Phone: 3174560250
 Owner: Richard Beeny
 Ownership Type:

LICENSE

License No: 61100944B
Profession: Pharmacy
 License Type: CSR
 1105 East Northfield Drive Suite 300-400
 Brownsburg IN 46112
 Phone: 3174560250

Inspection Type	Controlled Substances Inspection
Scheduled Date:	08/08/2016
Inspection Date:	9/23/2016
Announced?	N
Result:	Pass

Remarks: If you have any questions or concerns please feel free to contact me.

As the undersigned representative of the licensee, I affirm that the information provided during the inspection is true and accurate to the best of my knowledge. I understand that the licensee is responsible for the legal operations of the facility and I am executing this inspection form as, or on behalf of the licensee.

I understand that the licensee is on notice of the violation(s) listed herein and said violation(s) must be corrected within thirty (30) days of notification. Resolution of noncompliance shall occur as stipulated and the licensee shall fully comply with Indiana law, statutes, rules, and professional standards governing their practice. If necessary, a plan of corrective action shall be submitted to the Indiana Professional Licensing Agency within five (5) business days of the initial inspection. This plan of correction shall specify the immediate action to be taken to correct all noted deficiencies.

A plan of corrective action can be either emailed to PLACCompliance@pla.in.gov, or mailed to: Indiana Professional Licensing Agency, Attn: Compliance Division, 402 W. Washington Street, RM W072 Indianapolis, IN 46204. (Postmark must be within five (5) business days after the initial inspection). Failure to timely submit a corrective action plan will result in a personal appearance before the appropriate licensing board. If subsequent inspections identify a failure to address previous violations the facility will "fail" the inspection and disciplinary proceedings will be initiated against licensee.



9/23/2016



Signature of Inspector

Date

Signature of Owner/Licensee/Representative



Assessment Report

LifeScience Logistics, LLC

Assessment dates	02/25/2020 to 02/25/2020 (Please refer to Appendix for details)
Assessment Location(s)	Brownsburg (000)
Report Author	Myles Frohling
Assessment Standard(s)	ISO 13485:2016





Assessment Report.

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Assessment Report.

Executive Summary

The objectives of the assessment were met.

Obstacles, Omissions and Reliability

There were no obstacles encountered during the course of the audit. No factors were encountered during the audit that would affect the reliability of this assessment.

Areas Not Audited

All areas were covered per the assessment plan.

Identification and Dating

Audit report authors are as per the assessment team listed. The recommendation included in this assessment is based on the assessment of the sites documented in the 'assessed locations' table towards the rear of this report. This table also defines the assessment duration.

The report was finalized and issued on 25 Feb 2020.

If this visit is part of a multi-location assessment, the final recommendation will be contingent on the findings from all assessments.

CONTINUING ASSESSMENT

Please note that all recommendations are subject to independent review.

The management system has effectively implemented. The system addresses the scope of registration and is in accordance with the company objectives, applicable requirements of the management standard & BSI Conditions of Contract. The result of this assessment is a recommendation for certification.

ISO 13485:2016

All requirements of ISO13485:2016 are effectively implemented within the management system.



Assessment Report.

Changes in the organization since last assessment

There is no significant change of the organization structure and key personnel involved in the audited management system.

No change in relation to the audited organization's activities, products or services covered by the scope of certification was identified.

There was no change to the reference or normative documents which is related to the scope of certification.

NCR summary graphs

There have been no NCRs raised.

Your next steps

NCR close out process

There were no outstanding nonconformities to review from previous assessments.

No new nonconformities were identified during the assessment. Enhanced detail relating to the overall assessment findings is contained within subsequent sections of the report.

Please refer to Assessment Conclusion and Recommendation section for the required submission and the defined timeline.



Assessment Report.

Assessment objective, scope and criteria

Assessment Scope

The management system processes at the address defined in the 'assessed locations table' towards the rear of this report.

Assessment Objectives

SURVEILLANCE

To conduct a surveillance assessment to determine the continued effectiveness of implementation of the company's management system, in accordance with the company objectives, policies and procedures, the management standards and applicable regulatory requirements from relevant regulatory authorities & BSI Conditions of Contract. To ensure that all requirements are covered during the certification period and to determine whether a recommendation for continuing certification can be made.

ISO 13485:2016

To verify that all of the requirements of ISO13485:2016 are effectively implemented within the management system.

The scope of the assessment is the documented management system with relation to the requirements of ISO 13485:2016 and the defined assessment plan provided in terms of locations and areas of the system and organization to be assessed.

ISO 13485:2016

LifeScience Logistics, LLC. management system documentation



Assessment Report.

Assessment Participants

Name	Position	Opening Meeting	Closing Meeting	Interviewed(processes)
Paul Hayward	Director of Quality & Regulatory Affairs	X	X	X
Teri Johnson	Supervisor of Quality & Regulatory Affairs	X	X	X
Teurayi George	QA Specialist I	X	X	X
Nate Roberts	Director of Operations	X	X	
David Davis	Operations Manager		X	

Assessment conclusion

BSI assessment team

Name	Position
Myles Frohling	Team Leader

Assessment conclusion and recommendation

Audit objectives are met.

RECOMMENDED - The audited organization can be recommended for certification / recertification / continued certification to the above listed standards and has been found in general compliance with the audit criteria as stated in the above-mentioned audit plan.

Use of certification documents, mark / logo or report

The use of the BSI certification documents and mark / logo is effectively controlled.



Assessment Report.

Findings from this assessment

Opening meeting: QMS System Review, QMS, Business, Product & Process Changes and Improvements

Objectives and Targets Management Responsibility and Changes:4.1, 4.2, 5.1, 5.2, 5.3, 5.4, 5.5, 6.1, 8.1, 8.5.1

Opening Meeting, Manufacturer Information and Changes

The opening meeting was conducted with the presence of the Senior staff, Director of operations and Management Representative/Responsible Engineering Manager.

The assessment plan, objectives and scope of the assessment were confirmed.

The opening meeting and full assessment was performed in English.

Scope of Certification:

The registration certificates and scope of the registration were confirmed as follows:

Third Party Logistics for medical device companies, logistic provider performing relabeling, repackaging and distribution activities for medical device companies outside of manufacturing requirements.

Quality Manual Version: Quality Manual – Quality Manual – Rev. 019 – Eff. 24 Jun 2019

Exclusions and Non-Applications of Requirements in the QMS:

Exclusions:

7.3 Design and Development. This registration does not design or develop new products.

Non-Applications:

7.5.2 Cleanliness of Product and Contamination Control

7.5.3 Installation activities

7.5.5 Particular requirements for sterile medical devices (maintain records) 7.5.9.2 Particular requirements for implantable medical devices

Significant Changes:

There have not been any major or significant changes to the QMS, organizational structure, products or process since the last visit.

Adverse Incidents, Field Safety Corrective Actions and Recalls:

There have been no adverse incidents, recalls, or requirement for field safety corrective actions since the last report.

Corporate Identity of the Manufacturer:

Logistics Facilities: ATL - 4475 S. Fulton Pkwy Bldg. 5 Suite C, Atlanta GA 30349 (Approx. 255,000 sq. ft)
BWI – 8901 Snowden River Parkway Suite 150 Columbia, MD 21046 (Approx. 285,000 sq ft)



Assessment Report.

DFW – 2600 Regent Blvd, DFW Airport, TX 75261 (Approx. 405,000 sq ft) IND – 1105 Northfield Drive, Suite 400 Brownsburg, IN 46112 (Approx. 330,000 sq. ft)

Description of the manufacturer:

LifeScience Logistics offers supply chain solutions to customers that range from Big Pharma to small startups. The LSL's cGMP compliant facilities allow Pharmaceutical, medical device, plasma, and biotech clients to distribute frozen, refrigerated and controlled ambient temperature products to customers.

Critical Subcontractors:

LifeScience provides logistical services to government and medical device companies and does not manufacture/produce any medical devices, therefore there are no critical Subcontractors/Suppliers for the this site.

Senior Management of the Assessment Location(s).

[Confirm and record the name and title of the most senior individual]

Audit Duration Rationale:

Staffing and effective staffing numbers were reviewed against IAF MD9 annex D and MDP200 (CP0200). The effective number of staff was stated to be 106. 20% reduction is permissible due to this audit being of a distribution facility, no manufacturing is performed at this facility. Based on the number of effective staff and the recommended 20% reduction, the audit days are appropriate at 1.5 day surveillance and 3 days recertification.

Competency Code:

The competency code T71F is correct for the client and the competency code was covered over the certification cycle.

Quality Manual, Top Management, Management Review, Quality Objectives, Quality Policy, Analysis of data:5.1, 5.6, 5.4, 8.4, 4.2

Quality Manual was reviewed. Quality Manual Quality Manual – Quality Manual – Rev. 019 – Eff. 24 Jun 2019

and The Director of Quality & Regulator Affairs, was interviewed about this document in the Suite 300 conference room. The non-applications are documented and still reference to the older standard for some of the clauses. The non-Applications are documented as 7.5.3 – Installation Activities, 7.5.4 – Servicing Activities and Requirements for Active Implantable Devices.

It is based on the ISO 13485:2016 standard clauses and captures all the requirements per the standard. There are four levels of document structure. The Organization Chart was reviewed.

Quality Policy was reviewed, and it is approved.

Quality Objectives are defined in the Quality Manual and confirmed that they are defined in the Quality Manual.



Assessment Report.

These Quality Objectives are measurable, and they have metrics defined. Based upon the objective evidence reviewed the planned arrangements, implementation, delivery and monitoring for Quality Objectives were found to be effective to meet the needs of the business and compliant with the requirements of ISO 13485:2016.

Management review was assessed. Management review is controlled by SOP 1004 - Management Review – Rev. 007 – 19 Jan 2020. The Director of Quality & Regulatory Affairs and the Supervisor of Quality & Regulatory Affairs was interviewed in the Suite 300 QA conference room and demonstrated implementation of the process. A sample of Management review minutes was taken. The Director of Quality & Regulatory Affairs is the Management Representative. Management Review is performed once a year. Inputs and Outputs are all defined in the SOP. The Management review meeting completed on 12 Feb 2019 was reviewed and Meeting Agenda was reviewed. The meeting minutes and the overall summary was reviewed. The MR process adequately captures all the inputs per Clause 5.6.2 and the outputs per Clause 5.6.3.

Based upon the objective evidence reviewed the planned arrangements, implementation, delivery and monitoring for Management Review were found to be effective to meet the needs of the business and compliant with the requirements of ISO 13485:2016.

Analysis of Data was assessed. Analysis of Data is controlled by SOP 1005 – Statistical Techniques - Rev. 004 – 18 Oct 2019. The Director of Quality & Regulatory Affairs and the Supervisor of Quality & Regulatory Affairs was interviewed in the Suite 300 QA conference room and demonstrated implementation of the process.

Based upon the objective evidence reviewed the planned arrangements, implementation, delivery and monitoring for Analysis of Data process was found to be effective to meet the needs of the business and compliant with the requirements of ISO 13485:2016.

Documents and Records Reviewed:

Quality Manual – Quality Manual – Rev. 019 – Eff. 24 Jun 2019
 SOP 1004 - Management Review – Rev. 007 – 19 Jan 2020
 SOP 1005 – Statistical Techniques - Rev. 004 – 18 Oct 2019
 Organization Chart – Last Refreshed 23 July 2019
 Quality policy reviewed during Management Review (12 Feb 2019)
 Quality objectives
 Management Review Dated: 12 Feb 2019 for review of FY 2018
 Prior Management review Feb 2018
 Next Management Review scheduled for 11 Feb 2020
 Attendance Record – Management Review

Personnel Interviewed:

Paul Hayward – Director of Quality & Regulatory Affairs
 Teri Johnson – Supervisor of Quality & Regulatory Affairs



Assessment Report.

Internal Audit:8.2.2, 4.2.4, 8.2.3, 8.2.4

Internal audit was assessed. Internal audit is controlled by SOP 1501 – Internal Audits – Rev. 008 – Eff. 10 Jan 2020

and an issued schedule which was approved and issued on SOP 1501.01 Rev. 04 – Internal Audit Schedule FY: 2020 (Created 07 Oct 2019). The Director of Quality & Regulatory Affairs and the Supervisor of Quality & Regulatory Affairs was interviewed in the Suite 300 QA Conference Room and implementation of the process was confirmed in a review of the sample that was taken. A sample of internal audit records was taken and no issues were noted in the sample reviewed.

Based upon the objective evidence reviewed the planned arrangements, implementation, delivery and monitoring for internal audit were found to be effective to meet the needs of the business and compliant with the requirements of ISO 13485:2016.

Documents and Records Reviewed:

SOP 1501 – Internal Audits – Rev. 008 – Eff. 10 Jan 2020

SOP 1501.01 Rev. 00 - Internal Audit Schedule FY: 2019 (Created 16 Nov 2018)

SOP 1501.01 Rev. 04 – Internal Audit Schedule FY: 2020 (Created 07 Oct 2019)

Internal Audit Plan

SOP 1501.13 – Internal Audit Quality Checklist – Rev. 004 – Eff. 07 Jan 2020

SOP 1501.08 – Internal Audit Checklist – DEA – Rev. 002 – Eff. 10 Jan 2020

SOP 1501.02 – Internal Audit Cover Page – Rev. 003 – Eff. 15 Dec 2016

Internal Audit Reports sampled: IA-INC-19 (13 May 2019 – 16 May 2019)

- Major – 2

- Minor – 16

- Observations – 0

- K. Ashley (LA), D. Cherry, S. Nash, V. Schmidt

- Report Date: 05 June 2019

Personnel Interviewed:

Paul Hayward – Director of Quality & Regulatory Affairs

Teri Johnson – Supervisor of Quality & Regulatory Affairs

Feedback Processes, Complaints and Vigilance:7.2.3, 8.2.1

Feedback processes, including complaint handling and vigilance procedures were assessed. Complaints are controlled by SOP 4001 – Handling of Product Complaints – Rev. 007 – 01 Nov 2019. The Director of Quality & Regulatory Affairs and the Supervisor of Quality & Regulatory Affairs was interviewed in the Suite 300 QA Conference Room and demonstrated implementation of the process. A sample of complaints was taken, and no issues were noted in the sample reviewed.

Based upon the objective evidence reviewed the planned arrangements, implementation, delivery and monitoring for feedback processes, including complaint handling and vigilance procedures were found to be effective to meet the needs of the business and compliant with the requirements of ISO 13485:2016.



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Documents and Records Reviewed:

SOP 4004 – Client/ Customer Feedback – Commercial – Rev. 004 – Eff. 28 Aug 2018

SOP 4001 – Handling of Product Complaints – Rev. 007 – 01 Nov 2019

SOP 1301 – Risk Management – Rev. 005 – Eff. 20 Feb 2019

Complaint Log (2019): Qty. Open: 1973, Closed: 1817

Complaint Log (2020): Qty. Open: 158, Closed: 10

Reportable(s) (Any injury or death) = N/A

Complaints examined:

- CCF-AA-2020-0155 – Opened 20 Jan 2020 – Shipping Instructions Not Followed – Status: Open

- CCF-AA-2020-0153 – Opened 8 Jan 2020 – Shortage of product – Status: Investigation in Progress

- CCF-AN-2019-1477 – Opened 18 Sept 2019 – External Damage to product – Status: QA Review

- CCF-AN-2020-0035 – Opened: 20 Dec 2019 – Wrong Product – Status: QA Review

- CCF-AA-2020-0157 – Opened: 8 Jan 2020 – Transportation Issue. Order has not moved – Status: Closed a duplicate of CCF-AA-2020-0153 Closed 21 Jan 2020

Personnel Interviewed:

Paul Hayward – Director of Quality & Regulatory Affairs

Teri Johnson – Supervisor of Quality & Regulatory Affairs

Improvement: Corrective and Preventive Action:8.5.1, 8.5.2, 8.5.3

Corrective and preventive action processes were assessed. Corrective and preventive action is controlled by SOP 1350 – Deviation/ CAPA – Commercial – Rev. 003 – Eff. 25 Feb 2019. The Director of Quality & Regulatory Affairs and the Supervisor of Quality & Regulatory Affairs was interviewed in Suite 300 QA Conference Room and demonstrated implementation of the process. A sample of corrective and preventive actions was taken and all issues that were identified in the samples reviewed were addressed in an open CAPA and procedure change request that was circulating for approval.

Based upon the objective evidence reviewed the planned arrangements, implementation, delivery and monitoring for corrective and preventive action were found to be effective to meet the needs of the business and compliant with the requirements of ISO 13485:2016.

Documents and Records Reviewed:

SOP 4004 – Client/ Customer Feedback – Commercial – Rev. 004 – Eff. 28 Aug 2018

SOP 4001 – Handling of Product Complaints – Rev. 007 – 01 Nov 2019

SOP 1301 – Risk Management – Rev. 005 – Eff. 20 Feb 2019

SOP 1350 – Deviation/ CAPA – Commercial – Rev. 003 – Eff. 25 Feb 2019

SOP 1350.02 – Deviation Report e-log

SOP 1350.01 – Deviation Report

Complaint Log (2019): Qty. Open: 1973, Closed: 1817

Complaint Log (2020): Qty. Open: 158, Closed: 10

Reportable(s) (Any injury or death) = N/A

Complaints examined:

- CCF-AA-2020-0155 – Opened 20 Jan 2020 – Shipping Instructions Not Followed – Status: Open



Assessment Report.

- CCF-AA-2020-0153 – Opened 8 Jan 2020 – Shortage of product – Status: Investigation in Progress
- CCF-AN-2019-1477 – Opened 18 Sept 2019 – External Damage to product – Status: QA Review
- CCF-AN-2020-0035 – Opened: 20 Dec 2019 – Wrong Product – Status: QA Review
- CCF-AA-2020-0157 – Opened: 8 Jan 2020 – Transportation Issue. Order has not moved – Status: Closed a duplicate of CCF-AA-2020-0153 Closed 21 Jan 2020

NCs (Deviations)

DEV-INC-19-0008:

SOP 1350.01 – Deviation Report – Commercial – Rev. 004 – Eff. 27 Nov 2017

- Date 24 Feb 2019 – Deviation Type: 3 – Description: There was a temperature excursion in Zone 44 of System 1. – Status: Closed 4 Mar 2019
- DEV-INC-19-0008 Att. A – Detailed Temperature Report
- Att. B – Temp. Report
- Att. C – Explanation of the Preliminary Monthly Climate Data
- Att. D – Zone 44 Primary Probe Photos
- Email of information

DEV-INC-19-0017:

SOP 1350.01 – Deviation Report – Commercial – Rev. 004 – Eff. 27 Nov 2017

- Date 23 May 2019 – Deviation Type: 2 – Description: On 23 May 2019, BF QA communicated via telephone and follow-up email, that a packager received containers from the LSL that have Next Inspection Dates on the label that do not match dates in SAP – Status: Closed 29 Oct 2019
- SOP 1350.05 – Deviation/ CAPA Report Extension Request – Commercial – Rev. 001 – Dated: 26 June 2019 Approved
- SOP 1350.05 – Deviation/ CAPA Report Extension Request – Commercial – Rev. 001 – Dated: 23 July 2019 Approved
- Line Item spreadsheets

DEV-INC-19-0018:

SOP 1350.01 – Deviation Report – Commercial – Rev. 004 – Eff. 27 Nov 2017

- Dated: 24 May 2019 – Deviation Type: 1 & 2 – Description: Operations has a receiving error for items QD507Q and QD509B. These items received and put into the system in the BF Cage when they needed to bin in refrigerator storage 2-8C. – Status: Closed 26 June 2019
- Att. 1 – Spreadsheet of lots with locations
- Att. 2 – BF.001 – Receiving – Rev. 002 – Eff. 01 May 2019 (Training Record) – Date: 18 June 2019

DEV-INC-19-0028:

SOP 1350.01 – Deviation Report – Commercial – Rev. 004 – Eff. 27 Nov 2017

- Dated: 09 Oct 2019 – Deviation Type: 5 – Description: On 9 Oct 2019, AS Team member notified QA , facilities, and operations of a mechanical failure in cooler 20B causing the fans to distribute moisture into the chamber affecting 11 pallets of AS product. – Status: 7 Nov 2019
- Att. 1 – QA hold printout
- Att. 2 – Damaged Product Form – Rev. 002 – Eff. 28 Aug 2017 – Dated 9 Oct 2019
- Att. 3 – WI 100.04.008 – Request for Reactive Maintenance – ALL – Dated: 9 Oct 2019
- Att. 4 – Emails of explanation of incident
- Att. 5 – WI 100.04.037 – Reactive Maintenance Form – ALL – Dated: 28 Oct 2019



Assessment Report.

CAPA:

CAPA-INC-19-0001 – Opened 31 July 2019 – escalated from DEV-INC-19-0017

Personnel Interviewed:

Paul Hayward – Director of Quality & Regulatory Affairs

Teri Johnson – Supervisor of Quality & Regulatory Affairs

Product Realization: Planning and Customer Related Processes: 7.1, 7.2.1, 7.2.2, 4.2.4

Product realization processes were assessed including planning, risk assessment and customer related processes. Product Realization is controlled by:

SOP 2002 – Handling, Storage, Packaging & Distribution – Rev. 009 – Eff. 15 May 2018

WI 400.04 – Commercial Pick Pack and Ship – Rev. 012 – Eff. 26 Aug 2019. The Director of Quality & Regulatory Affairs and the Supervisor of Quality & Regulatory Affairs was interviewed in the Suite 300 QA Conference Room and demonstrated implementation of the process with no apparent issues. A sample of a customer related process was taken and there were no apparent issues noted with the process.

Based upon the objective evidence reviewed the planned arrangements, implementation, delivery and monitoring for product realization were found to be [effective [NO NC]/generally effective [MINOR NC]/not effective [MAJOR NC]] to meet the needs of the business and compliant [NO NC/not fully compliant [MINOR NC]/ not compliant [MAJOR NC]] with the requirements of ISO 13485:2016 / MDD 93/42 EEC.

Documents and Records Reviewed:

WI 400.01 – Commercial Receiving – Rev. 012 – Eff. 24 Oct 2018

SOP 2002 – Handling, Storage, Packaging & Distribution – Rev. 009 – Eff. 15 May 2018

WI 400.04 – Commercial Pick Pack and Ship – Rev. 012 – Eff. 26 Aug 2019

BF.001 – Receiving – Rev. 004 – Eff. 21 Jan 2020

SOP 1301 – Risk Management – Rev. 005 – Eff. 20 Feb 2019

Client Code BQ (name redacted) SOP for Inbound Shipments, Storage, Packing, Outbound Shipping – Version 3.3 – Eff. 3 July 2019

Personnel Interviewed:

Paul Hayward – Director of Quality & Regulatory Affairs

Teri Johnson – Supervisor of Quality & Regulatory Affairs



Assessment Report.

Purchasing and Supplier Management: 7.4.1, 7.4.2, 7.4.3

Purchasing and supplier management was assessed. Purchasing is controlled by SOP 1007 – Purchasing Process and Controls – Rev. 000 – Eff. 16 Jan 2019 and supplier management is controlled by SOP 1031 – Vendor Qualification – Rev. 011 – Eff. 06 Nov 2019. The Director of Quality & Regulatory Affairs and the Supervisor of Quality & Regulatory Affairs was interviewed in the Suite 300 QA Conference Room and demonstrated implementation of the process. A sample of supplier agreements, purchase orders and supplier audits was taken and the only issue that was noted is that purchasing related to product is all completed at the corporate headquarters, not here. There were no issues noted in the samples that were reviewed.

Based upon the objective evidence reviewed the planned arrangements, implementation, delivery and monitoring for purchasing and supplier management were found to be effective to meet the needs of the business and compliant with the requirements of ISO 13485:2016.

Documents and Records Reviewed:

SOP 1007 – Purchasing Process and Controls – Rev. 000 – Eff. 16 Jan 2019
SOP 1031 – Vendor Qualification – Rev. 011 – Eff. 06 Nov 2019
SOP 1031.04 – Controlled Supplier List – All – Rev. 033 – Eff. 13 Jan 2020

Service Agreement

OPC Pest Control – Dated 4/2019 – 3/2020
Ryan Fire Protection, Inc. – 10 Dec 2019
Mold Diagnostics – No Agreement

Certificate of Liability Insurance

OPC Pest Control – Good through Jan 2021
Ryan Fire Protection, Inc. – 30 Dec 2020
Mold Diagnostics – Exp. 1 Jan 2021

Vendor Acknowledgement of Warehouse Policies Form

OPC Pest Control – 18 Sept 2019
Ryan Fire Protection, Inc. – 16 Jan 2020
Mold Diagnostics – No Form at this time

Controlled Vendor Qualification Questionnaire

OPC Pest Control – Established Vendor No Questionnaire
Ryan Fire Protection, Inc. – 23 Dec 2019
Mold Diagnostics – Established Vendor No Questionnaire

Vendor Evaluation

OPC Pest Control – 23 Sept 2019
Mold Diagnostics – 23 Sept 2019
Ryan Diagnostics – Does not have one because vendor is less than a year old

Personnel Interviewed:



Assessment Report.

Paul Hayward – Director of Quality & Regulatory Affairs
Teri Johnson – Supervisor of Quality & Regulatory Affairs

Resources: Human Resources, Infrastructure and Work Environment:6.1, 6.2, 6.3, 6.4

Resource provision including human resources, infrastructure and work environment was assessed. Human resource management is controlled by SOP 1800 – Training and Qualification – Rev. 017 – Eff. 11 Dec 2019.

Maintenance of the work environment and infrastructure is controlled by SOP 204 – Business Continuity Plan – Rev. 005 – Eff. 08 July 2019. The Director of Quality & Regulatory Affairs and Supervisor of Quality & Regulatory Affairs was interviewed in Suite 300 QA Conference Room area and demonstrated implementation of the process and there were no issues noted in the samples taken and reviewed. A sample of training records to provide objective evidence of competency, awareness and training was taken and there were no noted issues with the sample that was reviewed. A sample of planned preventive maintenance records was taken and there were no issues noted with the sample that was taken and reviewed.

Based upon the objective evidence reviewed the planned arrangements, implementation, delivery and monitoring for resource provision including competency, awareness and training and management of infrastructure and work environment were found to be effective to meet the needs of the business and compliant with the requirements of ISO 13485:2016.

Documents and Records Reviewed:

SOP 1800 – Training and Qualification – Rev. 017 – Eff. 11 Dec 2019
Training Role Reports
Employee Training Form
SOP 1800.02 Procedure Training Form
SOP 1800.03 Non-Procedure Training Form
SOP 3001 – Emergency Response Plan – Rev. 008 – Eff. 09 Oct 2019
SOP 3002 – Pest Control – Rev. 007 – Eff. 16 Apr 2018
SOP 204 – Business Continuity Plan – Rev. 005 – Eff. 08 July 2019
WI 100.02 – Facility Sanitation – Rev. 009 – Eff. 20 Dec 2019
WI 100.02.03 – Master Cleaning Schedule-Commercial – Rev. 000 – Eff. 06 Dec 2018
SOP 2002 – Handling, Storage, Packaging & Distribution – Rev. 009 – 15 May 2018
WI 100.03 – Temperature Review – Rev. 012 – Eff. 23 Aug 2019
SOP 1001.08 – Annual cGMP Training – Rev. 003 – Eff. 23 Apr 2019

Training Role Report:

Teurayi George – QA Specialist I
Brook Austin – Warehouse Lead
John Graham – Warehouse Personnel

Training Records were reviewed in the MQ1 system for:

Teurayi George
Brook Austin
John Graham



Assessment Report.

Job Description:
Quality Assurance Specialist I
Warehouse Lead
Warehouse Personnel

WI 100.02.03 – Master Cleaning Schedule – Commercial – Rev. 000 – Eff. 06 Dec 2018

- Jan 2020
- Dec 2019
- Nov 2019

Pest Control Logs

- January – Being conducted today Pest control is on site
- December – 5 Dec 2019 – OPC-4
- November – 8 Nov 2019 – OPC-4
- October – 3 October 2019 – OPC-4

Personnel Interviewed:

Paul Hayward – Director of Quality & Regulatory Affairs
Teri Johnson – Supervisor of Quality & Regulatory Affairs
Teurayi George – QA Specialist I

Control of Nonconforming Product:4.2.5, 8.3

Control of nonconforming product was assessed. Control of nonconforming product is controlled by SOP 1350 – Deviation/ CAPA – Commercial – Rev. 003 – Eff. 25 Feb 2019. The Director of Quality & Regulatory Affairs and Supervisor of Quality & Regulatory Affairs was interviewed in the Suite 300 QA Conference Room area and demonstrated implementation of the process. A sample of internal nonconforming product records was taken and there were no noted issues within the sample reviewed.

Based upon the objective evidence reviewed the planned arrangements, implementation, delivery and monitoring for control of nonconforming product were found to be effective to meet the needs of the business and compliant with the requirements of ISO 13485:2016.

Documents and Records Reviewed:

SOP 4004 – Client/ Customer Feedback – Commercial – Rev. 004 – Eff. 28 Aug 2018
SOP 4001 – Handling of Product Complaints – Rev. 007 – 01 Nov 2019
SOP 1301 – Risk Management – Rev. 005 – Eff. 20 Feb 2019
SOP 1350 – Deviation/ CAPA – Commercial – Rev. 003 – Eff. 25 Feb 2019
SOP 1350.02 – Deviation Report e-log
SOP 1350.01 – Deviation Report

NCs (Deviations)

DEV-INC-19-0008:

SOP 1350.01 – Deviation Report – Commercial – Rev. 004 – Eff. 27 Nov 2017

- Date 24 Feb 2019 – Deviation Type: 3 – Description: There was a temperature excursion in Zone 44 of



Assessment Report.

System 1. – Status: Closed 4 Mar 2019

- DEV-INC-19-0008 Att. A – Detailed Temperature Report
- Att. B – Temp. Report
- Att. C – Explanation of the Preliminary Monthly Climate Data
- Att. D – Zone 44 Primary Probe Photos
- Email of information

DEV-INC-19-0017:

- SOP 1350.01 – Deviation Report – Commercial – Rev. 004 – Eff. 27 Nov 2017
- Date 23 May 2019 – Deviation Type: 2 – Description: On 23 May 2019, BF QA communicated via telephone and follow-up email, that a packager received containers from the LSL that have Next Inspection Dates on the label that do not match dates in SAP – Status: Closed 29 Oct 2019
- SOP 1350.05 – Deviation/ CAPA Report Extension Request – Commercial – Rev. 001 – Dated: 26 June 2019 Approved
- SOP 1350.05 – Deviation/ CAPA Report Extension Request – Commercial – Rev. 001 – Dated: 23 July 2019 Approved
- Line Item spreadsheets

DEV-INC-19-0018:

- SOP 1350.01 – Deviation Report – Commercial – Rev. 004 – Eff. 27 Nov 2017
- Dated: 24 May 2019 – Deviation Type: 1 & 2 – Description: Operations has a receiving error for items QD507Q and QD509B. These items received and put into the system in the BF Cage when they needed to bin in refrigerator storage 2-8C. – Status: Closed 26 June 2019
- Att. 1 – Spreadsheet of lots with locations
- Att. 2 – BF.001 – Receiving – Rev. 002 – Eff. 01 May 2019 (Training Record) – Date: 18 June 2019

DEV-INC-19-0028:

- SOP 1350.01 – Deviation Report – Commercial – Rev. 004 – Eff. 27 Nov 2017
- Dated: 09 Oct 2019 – Deviation Type: 5 – Description: On 9 Oct 2019, AS Team member notified QA , facilities, and operations of a mechanical failure in cooler 20B causing the fans to distribute moisture into the chamber affecting 11 pallets of AS product. – Status: 7 Nov 2019
- Att. 1 – QA hold printout
- Att. 2 – Damaged Product Form – Rev. 002 – Eff. 28 Aug 2017 – Dated 9 Oct 2019
- Att. 3 – WI 100.04.008 – Request for Reactive Maintenance – ALL – Dated: 9 Oct 2019
- Att. 4 – Emails of explanation of incident
- Att. 5 – WI 100.04.037 – Reactive Maintenance Form – ALL – Dated: 28 Oct 2019

Personnel Interviewed:

Paul Hayward – Director of Quality & Regulatory Affairs
Teri Johnson – Supervisor of Quality & Regulatory Affairs
Teurayi George – QA Specialist I



Assessment Report.

Control of Documents and Records:4.2.3, 4.2.4, 4.2.5

Control of documents and records were assessed.

Control of Documents are controlled by SOP 1100 – Document Control – Rev. 012 – Eff. 09 Oct 2019. Control of records processes are controlled by SOP 1101 – Control of Records – Rev. 022 – Eff. 14 Oct 2019. The Director of Quality & Regulatory Affairs and Supervisor of Quality & Regulatory Affairs was interviewed in the Suite 300 QA Conference Room and demonstrated implementation of the process. Based upon the objective evidence reviewed the planned arrangements, implementation, delivery and monitoring for control of documents and records were found to be effective to meet the needs of the business and compliant with the requirements of ISO 13485:2016.

Documents and Records Reviewed:

Document Master List – In MQ1

SOP 1100 – Document Control – Rev. 012 – Eff. 09 Oct 2019

SOP 1101 – Control of Records – Rev. 022 – Eff. 14 Oct 2019

SOP 1002 – Change Control – Rev. 010 – Eff. 11 Apr 2019

WI 100.03 – Temperature Review – Rev. 012 – Eff. 23 Aug 2019 – CR# 0011411

SOP 2002 – Handling, Storage, Packaging & Distribution – Rev. 009 – Eff. 15 May 2018 – CR#001046

WI 100.02 – Facility Sanitation – Rev. 009 – Eff. 20 Dec 2019 – CR# 001423

SOP 3001 – Emergency Response Plan – Rev. 008 – Eff. 09 Oct 2019 – CR#001417

SOP 2004 – Business Continuity Plan – rev. 005 – Eff. 08 Jul 2019 – CR# 001377

Wi 100.19 – Emergency Response Plan – Rev. 006 – Eff. 30 July 2019 – CR# 001390

SOP 3002 – Pest Control – Rev. 007 – Eff. 16 April 2018 – CR#00970

Personnel Interviewed:

Paul Hayward – Director of Quality & Regulatory Affairs

Teri Johnson – Supervisor of Quality & Regulatory Affairs



Assessment Report.

Next visit objectives, scope and criteria

Assessment Scope

The management system processes at the address defined in the 'assessed locations table' towards the rear of this report.

Assessment Objectives

SURVEILLANCE

To conduct a surveillance assessment to determine the continued effectiveness of implementation of the company's management system, in accordance with the company objectives, policies and procedures, the management standards and applicable regulatory requirements from relevant regulatory authorities & BSI Conditions of Contract. To ensure that all requirements are covered during the certification period and to determine whether a recommendation for continuing certification can be made.

ISO 13485:2016

To verify that all of the requirements of ISO13485:2016 are effectively implemented within the management system.

The scope of the assessment is the documented management system with relation to the requirements of ISO 13485:2016 and the defined assessment plan provided in terms of locations and areas of the system and organization to be assessed.

ISO 13485:2016

LifeScience Logistics, LLC. management system documentation

Please refer to BSI terms and conditions regarding cancellation of planned visits.



Assessment Report.

Next Visit Plan

Date	Auditor	Time	Area/Process	Clause
01/19/2021			Opening Meeting	
01/19/2021			Site Tour	
01/19/2021	Assessor 1		Certificate Scope, Quality Manual, Quality Policy and Quality Objectives and Analysis of Data	
01/19/2021			Leadership, Management Commitment, Management Responsibility, Management Review, and Analysis of Data	
01/19/2021			Feedback, Continual Improvement, Complaints, Adverse Event Reporting, Recalls, and Advisory Notices	
01/19/2021			Regulatory, Combined Checklist, Progress Toward Transition to ISO 13485:2016	
01/19/2021			Internal Audits	
01/19/2021			Lunch	
01/19/2021			Corrective Actions and Preventive Actions	
01/19/2021			Preservation of Product, Receiving, Receiving Inspection, and Control of Nonconforming Product	
01/19/2021			Shipping, Distribution, Identification, Traceability, Inspection (including cold storage and ambient temperature), and Control of Nonconforming Product	
01/20/2021			Calibration, Preventive Maintenance, Control of Monitoring and Measuring Devices and Validation of Processes	
01/20/2021			Training and Competency, HR and Provision of Resources	
01/20/2021			Report Preparation	
01/20/2021			Closing Meeting	



Assessment Report.

Appendix: Your certification structure & ongoing assessment program

Scope of Certification

FM 646928 (ISO 13485:2016)

Third Party Logistics for medical device companies, logistic provider performing relabeling, repackaging and distribution activities for medical device companies outside of manufacturing requirements.

Certificate Scheme:

Scheme manager:

Assessed location(s)

The audit has been performed at Permanent Locations.

Brownsburg / FM 646928 (ISO 13485:2016)

Location reference	0047585260-000
Address	LifeScience Logistics, LLC 1105 East Northfield Drive Suite 300 Brownsburg Indiana 46112 USA
Visit type	Special Audit
Assessment reference	3158671
Assessment dates	02/25/2020
Audit Plan (Revision Date)	01/11/2020
Deviation from Audit Plan	No
Total number of Employees	106
Effective number of Employees	106
Scope of activities at the site	Third Party Logistics for medical device companies, logistic provider performing relabeling, repackaging and distribution activities for medical device companies outside of manufacturing requirements.
Assessment duration	0.5 Day(s)



Assessment Report.

Certification assessment program

Certificate Number - FM 646928

Location reference - 0047585260-000

		Audit1	Audit2	Audit3
Business area/Location	Date (mm/yy):	01/19	1/20	1/21
	Duration (days):	2.0	1.5	1.5
Certificate Scope, Quality Manual, Quality Policy and Quality Objectives and Analysis of Data		X	X	X
Leadership, Management Commitment, Management Responsibility, Management Review, and Analysis of Data		X	X	X
Feedback, Continual Improvement, Complaints, Adverse Event Reporting, Recalls, and Advisory Notices		X	X	X
Regulatory, Combined Checklist, Progress Toward Transition to ISO 13485:2016		X	X	X
Internal Audits		X		X
Corrective Actions and Preventive Actions		X	X	X
Preservation of Product, Receiving, Receiving Inspection, and Control of Nonconforming Product		X		X
Purchasing, Supplier Qualification and Supplier Monitoring		X	X	
Shipping, Distribution, Identification, Traceability, Inspection (including cold storage and ambient temperature), and Control of Nonconforming Product		X		X
Planning of Product Realization including Risk Management and Customer Related Processes		X	X	
Calibration, Preventive Maintenance, Control of Monitoring and Measuring Devices and Validation of Processes		X		X
Work Environment including Pest Control, Infrastructure		X	X	
Training and Competency, HR and Provision of Resources		X		X
Control of Documents and Control of Records		X	X	
Stage 1 Assessment				
Stage 2 Assessment				



Assessment Report.

Interview with Top Management	X		
Continuing Surveillance Assessment (CAV)			
Recertification Assessment	X		

Definitions of findings:

Nonconformity:

Non-fulfilment of a requirement.

Major nonconformity:

Nonconformity that affects the capability of the management system to achieve the intended results.

Nonconformities could be classified as major in the following circumstances:

- If there is a significant doubt that effective process control is in place, or that products or services will meet specified requirements;
- A number of minor nonconformities associated with the same requirement or issue could demonstrate a systemic failure and thus constitute a major nonconformity.

Minor nonconformity:

Nonconformity that does not affect the capability of the management system to achieve the intended results.

Opportunity for improvement:

It is a statement of fact made by an assessor during an assessment, and substantiated by objective evidence, referring to a weakness or potential deficiency in a management system which if not improved may lead to nonconformity in the future. We may provide generic information about industrial best practices but no specific solution shall be provided as a part of an opportunity for improvement.

Observation:

It is ONLY applicable for those schemes which prohibit the certification body to issue an opportunity for improvement.

It is a statement of fact made by the assessor referring to a weakness or potential deficiency in a management system which, if not improved, may lead to a nonconformity in the future.

How to contact BSI

'Just for Customers' is the website that we are pleased to offer our clients following successful registration, designed to support you in maximizing the benefits of your BSI registration - please go to www.bsigroup.com/j4c to register. When registering for the first time you will need your client reference number and your certificate number



Assessment Report.

Should you wish to speak with BSI in relation to your certification, please contact your local BSI office – contact details available from the BSI website:

<https://www.bsigroup.com/en-US/contact-us/>

Notes

This report and related documents are prepared for and only for BSI's client and for no other purpose. As such, BSI does not accept or assume any responsibility (legal or otherwise) or accept any liability for or in connection with any other purpose for which the Report may be used, or to any other person to whom the Report is shown or in to whose hands it may come, and no other persons shall be entitled to rely on the Report. If you wish to distribute copies of this report external to your organization, then all pages must be included.

BSI, its staff and agents shall keep confidential all information relating to your organization and shall not disclose any such information to any third party, except that in the public domain or required by law or relevant accreditation bodies. BSI staff, agents and accreditation bodies have signed individual confidentiality undertakings and will only receive confidential information on a 'need to know' basis.

This audit was conducted on-site through document reviews, interviews and observation of activities. The audit method used was based on sampling the organization's activities and it was aimed to evaluate the fulfilment of the audited requirements of the relevant management system standard or other normative document and confirm the conformity and effectiveness of the management system and its continued relevance and applicability for the scope of certification.

As this audit was based on a sample of the organization's activities, the findings reported do not imply to include all issues within the system.

Regulatory compliance

BSI conditions of contract for this visit require that BSI be informed of all relevant regulatory non-compliance or incidents that require notification to any regulatory authority. Acceptance of this report by the client signifies that all such issues have been disclosed as part of the assessment process and agreement that any such non-compliance or incidents occurring after this visit will be notified to the BSI client manager as soon as practical after the event.

Attachment D

FDA-Approved Counterpart of the Eligible Prescription Drug, Copy of the Proposed Labeling for the Eligible Prescription Drug, Side-by-Side Comparison of the FDA-Approved Labeling and the Proposed Labeling, and Copies of the HPFB-Approved Labelling

This attachment incorporates by file a document for each drug identified below. Each file contains:

- **A drug label cover sheet** providing each eligible prescription drug's FDA-approved counterpart, the manufacturer(s) of the finished dosage form and the active ingredient or ingredients of each eligible prescription drug that the SIP Sponsor seeks to import, the legal relationship of these entities to the SIP Sponsor, and information about the HPFB-approved product and its FDA-approved counterpart.
- **The proposed label and side-by-side comparison** of proposed label to FDA-approved drug label, including the Prescribing Information, carton and container labeling, and patient labeling (e.g., Medication Guide, Instructions for Use, patient package inserts), with all differences annotated and explained.
- **The proposed package insert.**
- **The FDA-approved package insert.**
- **The HPFB-approved package insert.**

Drug Name

Biktarvy	Genvoya	Mavyret
Combivent Respimat	Harvoni	Odefsey
Complera Tabs	Incruse Ellipta	Spiriva
Descovy	Intelence	Spiriva Respimat
Edurant	Isentress	Stribild
Emtricitabine Tenofovir	Januvia	Tradjenta
Epclusa	Juluca	Triumeq
Farxiga	Kaletra	Wixela Inhub
Fluphenazine	Latuda	Zepatier

Attachment E

Foreign Seller Attestation, License, FDA Registration, and Inspectional History



March 12, 2021

Steve Solomon
Life Science Logistics, LLC
2600 Regent Boulevard
DFW Airport, Texas 75261
USA

Dear Steve,

**Re: Wholesaler Attestations to qualify as an SIP Foreign Seller
Methapharm Inc. (Canada)**

Methapharm attests that there are no past criminal convictions or violations of State, Federal, or Canadian laws regarding drugs or devices against our company. Further, Methapharm attests that any responsible individual(s) have not been involved in, or convicted of, any such violations. Methapharm is 100% owned by its corporate parent Polar Pharmaceuticals Inc., an Ontario, Canada corporation.

Lastly, there have been no disciplinary actions against the responsible individual(s) or our company by State, Federal, or Canadian regulatory bodies, including any such actions against the principals, owners, directors, officers, quality unit, or any facility manager or designated representative of such manager for the previous 7 years prior to submission of the SIP Proposal, with the exception of the following single administrative fine to our Coral Springs, Florida based affiliate from the referenced state body:

<i>Michigan (2014)</i>	<i>Methapharm, Inc has been licensed as a manufacturer/wholesaler in the State of Michigan since July 2002. In June 2014, we entered into a Consent Order with the Michigan Department of Licensing and Regulatory Affairs whereby we would pay a nominal fine of \$250. The subject matter of the Michigan complaint was <u>not</u> that we were in violation of the Michigan Public Health Code; rather, the State of Michigan took the position that they were able to seek an administrative fine in 2014 because of the administrative actions imposed on us in Colorado (in 2009) and Maine (in 2010) – both of which had been fully disclosed to the State of Michigan throughout the period of 2010-2014).</i>
------------------------	--

If you require further clarification or information with respect to the above, please do not hesitate to contact us.

Sincerely yours,

A handwritten signature in blue ink, appearing to read "Chris Calenti".

Chris Calenti
Director

Direct line: 1-800-287-7686 x 7222

Methapharm Inc. 81 Sinclair Blvd. Brantford Ontario N3S 7X6 Canada
519.751.3601 1.800.287.7686 | Fax 519.751.9149 | www.methapharm.com

Copy of Methapharm's DELHealth
Canada Santé
Canada

This licence is only valid when viewed electronically / Cette licence est seulement valide lorsqu'elle est visionnée électroniquement

Establishment Licence**Licence No. / No. de la licence**
100927-A**Licence d'établissement****Methapharm Inc****81 Sinclair Boulevard
Brantford ON N3S 7X6**This licence is issued in accordance with the *Food and Drugs Act and Regulations* (Division 1A) for the following activities /
Cette licence est délivrée conformément à la *Loi et au Règlement sur les aliments et drogues* (télé 1 A) pour les activités et les catégories de drogues
suivantes :

Category / Catégorie	Activity / Activité	Non-Sterile / Non-Stérile	Sterile / Stérile
Biological / Biologique	Distribute / Distribuer		X
Pharmaceutical / Pharmaceutique	Distribute / Distribuer	X	X
Active Pharmaceutical Ingredients / Ingrédients actifs pharmaceutiques	Import / Importer	X	
Biological / Biologique	Import / Importer		X
Pharmaceutical / Pharmaceutique	Import / Importer	X	X
Pharmaceutical / Pharmaceutique	Label / Étiqueter	X	X
Pharmaceutical / Pharmaceutique	Package / Emballer	X	X
Prescription Drug List, Schedule G, Narcotics, and/or Drug containing Cannabis / Liste des drogues sur ordonnance, l'Annexe G, Stupéfiants et/ou Drogue contenant du Cannabis	Wholesale / Vendre en gros	X	X

1 - if applicable - only a few

• Biologique: includes drugs listed in Schedule D to the Act, other than vaccines or whole blood and its components • Biologique: inclut les drogues visées à l'annexe D de la Loi, autres que les vaccins ou le sang total et ses composants

• Active Pharmaceutical Ingredients: includes drugs listed in Schedule C, to the Act / • Ingrédients pharmaceutiques: inclut les drogues répertoriées à l'annexe C de la Loi

2 - if applicable - only a few

• Distributor: as set out in paragraph C 31.1 A, 303 (4) and/or (6) / • Distributeur: à titre de distributeur au sens de l'alinéa C 31.1 A, 303 (4) et/ou (6)

• Tests include analysis and examinations required under Division 2 / • Analyser: conformément à l'article 2

This licence contains the following additional annex(es) / Cette licence contient les annexes suivantes :

Terms and Conditions / Modalités et conditions


Foreign Building Annex / Annexe concernant les bâtiments étrangers

Alternate Sample Retention Building Annex / Annexe concernant les autres bâtiments de conservation d'échantillons

Warehouse Annex / Annexe des entrepôts

Active Pharmaceutical Ingredient Foreign Building Annex / Annexe concernant les bâtiments étrangers d'ingrédients pharmaceutiques actifs

Date of last GMP inspection / Date de la dernière inspection BPF : 2018-12-10

Authorized Signatory, Regulatory Operations and Regions Branch or designated official Contrôleur/général, Direction générale des opérations réglementaires et des régions ou responsable désigné	
	Godard, Kim O C=CA, O=GC, OU=HC-SC, CN="Godard, Kim O" I attest to the accuracy and integrity of this document

This licence is the property of the Regulatory Operations and Regions Branch and must be returned upon demand
Cet avis de licence appartient à la Direction générale des opérations réglementaires et des régions et doit être retourné sur demande**Canada****Licence No. / No. de la licence : 100927-A****Issued on / Émise le : 2020-12-14****Page 1 of / de 32**

Copy of Methapharm's Registration as SIP Foreign Seller**Drug Establishments Current Registration Site**

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[New Search](#)

Search Results for **methapharm Inc.**

[CSV](#)
[Excel](#)

Filter:

Firm Name	FDA Establishment Identifier	DUNS	Business Operations	Address	Expiration Date
Methapharm Inc.		253526222	SIP FOREIGN SELLER	81 Sindall Boulevard, Brantford, Ontario N3S 7X6, Canada (CAN)	12/31/2021

Showing 1 to 1 of 1 entries

Previous: [1](#) Next

Data Current through: Friday, Mar 5, 2021

3/12/2021

Drug & health product inspections



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→ [Drug and health product inspections](#)

Drug & health product inspections

Licensing information

Establishment name:

Methapharm Inc

Address:

81 Sinclair Boulevard
Brantford, Ontario
Canada
N3S 7X6

Reference number:

503239

Site:

A

Licence number:

100927

Currently licensed:

Yes

Activities(categories):

- Distribute (Biological)
- Distribute (Pharmaceutical)
- Import (Active Pharmaceutical Ingredients)
- Import (Biological)
- Label (Pharmaceutical)
- Package (Pharmaceutical)
- Wholesale (Prescription Drug List, Schedule G, and/or Narcotics or a drug containing cannabis as defined in subsection 2(1) of the Cannabis Act.)
- Import (Pharmaceutical)

3/12/2021

Drug & health product inspections

Terms and conditions:Yes**Inspection information****Inspection:**

Inspection start date	Rating	Type of inspection
2018-12-10	<u>Compliant</u>	GMP Domestic - Regular Inspection
2015-10-07	<u>Compliant</u>	GMP Domestic - Regular Inspection
2013-10-17	Compliant	GMP Domestic - Regular Inspection

Date modified:

2016-11-08

3/12/2021

Drug & health product inspections



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Drug & health product inspections

Inspection report card summary

[Initial inspection deficiencies report](#)

Establishment name	Reference number	Inspection start date	Type of inspection	Inspection rating
Methapharm Inc	503239	2015-10-07	GMP Domestic - Regular Inspection	Compliant

Summary of observations

Filter items







Showing 1 to 8 of 8 entries **Show** **entries**

Observation number	Regulation	Summary of observation
1	C.02.015 - Quality Control	<ul style="list-style-type: none"> Deficiencies were noted with the documentation by the quality control department of process changes.
2	C.02.006 - Personnel	<ul style="list-style-type: none"> Deficiencies were noted with the training of personnel with respect to GMP principles, relevant standard operations procedures and/or their specific job duties.
3	C.02.015 - Quality Control	<ul style="list-style-type: none"> Deficiencies were noted with the implementation of guidelines put in place by the quality control department to ensure that storage conditions would maintain the quality and safe distribution of a drug.

<https://www.drug-inspections.canada.ca/en/full-report/?ref=insp21en0002&inspNumber=503239>

3/12/2021

Drug & health product inspections

Observation number  	Regulation  	Summary of observation  
4	C.02.004 - Premises	<ul style="list-style-type: none"> Deficiencies were noted with aspects of the building design intended to prevent the entry of pests. Deficiencies were noted with the design of premises to avoid mix-ups
5	C.02.012 - Manufacturing Control	<ul style="list-style-type: none"> Active Pharmaceutical Ingredients were manufactured and tested from companies for which GMP compliance information was not available.
6	C.02.011 - Manufacturing Control	<ul style="list-style-type: none"> Deficiencies were noted with the release from quarantine of finished products by the quality control department,
7	C.02.011 - Manufacturing Control	<ul style="list-style-type: none"> Deficiency was noted with the firm's Annual Product Review program.
8	C.02.015 - Quality Control	<ul style="list-style-type: none"> Deficiencies were noted with the investigation of complaints and deviations

1

Inspection outcome

Inspection resulted in a Compliant rating with Terms and Conditions. The site has been determined to be in Compliance with Part C, Division 2 of the Food and Drug Regulations.

Ratings are the result of observations made by Health Canada based on a reasonable belief at a particular point in time during the course of an inspection that the company was conducting the regulated activity / activities in compliance with the Food and Drugs Act or its Regulations.

Measures taken by Health Canada

3/12/2021

Drug & health product inspections

- Drug Establishment Licence (DEL) with Terms & Conditions was maintained.

Date modified:

2016-11-08

3/12/2021

Drug & health product inspections



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Drug & health product inspections

Inspection report card summary

[Initial inspection deficiencies report](#)

Establishment name	Reference number	Inspection start date	Type of inspection	Inspection rating
Methapharm Inc	503239	2018-12-10	GMP Domestic - Regular Inspection	Compliant

Summary of observations

Filter items







Showing 1 to 10 of 10 entries

Show entries

Observation number	Regulation	Summary of observation
1	C.02.011 - Manufacturing control	<ul style="list-style-type: none"> The quality control department did not evaluate the results of the annual product quality review.
2	C.02.014 - Quality control department	<ul style="list-style-type: none"> The assessment to release finished products was inadequate.
3	C.02.014 - Quality control department	<ul style="list-style-type: none"> The assessment, documentation, and/or procedures for considering the resale of returned drugs were inadequate.

3/12/2021

Drug & health product inspections

Observation number  	Regulation  	Summary of observation  
4	C.02.015 - Quality control department	<ul style="list-style-type: none"> The assessment, recording, follow-up, and/or investigation of complaints and/or other information about potentially defective products was inadequate.
5	C.02.015 - Quality control department	<ul style="list-style-type: none"> The premises were not laid out and/or designed for an optimal flow of personnel and/or materials to prevent cross-contamination and/or mix-ups between products.
6	C.02.027 - Stability	<ul style="list-style-type: none"> The company did not determine the stability of the drug before it was marketed or before significant changes were made to the drug formulations, fabrication procedures, or packaging materials.
7	C.02.028 - Stability	<ul style="list-style-type: none"> The ongoing stability program for a drug was inadequate.
8	C.02.003 - Sale	<ul style="list-style-type: none"> The importer did not ensure that the foreign sites use to fabricate, package/label, or test drug products met the requirements described in Guidance on Evidence to Demonstrate Drug GMP Compliance of Foreign Sites (GUI-0080).
9	C.02.015 - Quality control department	<ul style="list-style-type: none"> The quality agreement outlining the responsibilities for fabricating, packaging or labelling, and/or testing was inadequate.
10	C.02.015 - Quality control department	<ul style="list-style-type: none"> The guidelines and/or procedures were inadequate in ensuring storage and/or transportation conditions would maintain the quality and safe distribution of the drug.

3/12/2021

Drug & health product inspections

1

Inspection outcome

Inspection resulted in a Compliant rating. The site has been determined to be in Compliance with Part C, Division 2 of the Food and Drug Regulations. Ratings are the result of observations made by Health Canada based on a reasonable belief at a particular point in time during the course of an inspection that the company was conducting the regulated activity / activities in compliance with the Food and Drugs Act or its Regulations.

Measures taken by Health Canada

- Drug Establishment Licence (DEL) was maintained.

Date modified:

2016-11-08