- 3. Stericycle's parent company Stericycle, Inc. was founded in 1989. Stericycle, Inc.'s goal then, as now, was providing safe and environmentally responsible management of regulated medical waste. Stericycle, Inc. has grown substantially since its founding. In 1996 it became a publically traded company and in 2000 Stericycle, Inc. and its subsidiaries became the largest provider of medical waste services in North America. Stericycle, Inc. now provides services worldwide, operating in the United States, Argentina, Brazil, Canada, Chile, Ireland, Japan, Mexico, Portugal, Romania, Spain, and the United Kingdom. This rapid growth has been fueled by Stericycle, Inc.'s commitment to its founding goals and its innovations to achieve these goals.
- 4. Stericycle of Washington, Inc. was established in 1993 to provide biomedical waste transportation and collection services to Washington biomedical waste generators. Stericycle offered cradle-to-grave tracking and documentation of biomedical waste from collection to treatment, reusable, leak- and puncture-proof plastic containers for sharps waste, integrated training to help generators correctly classify, segregate, and reduce biomedical waste and improve employee safety, and offered all of these activities state-wide. Stericycle has offered these and other services to Washington generators for almost 20 years without raising its rates for regulated services.
- 5. Stericycle is committed to serving the needs of all medical waste generators. Stericycle has a diverse customer base, serving both large quantity generators such as hospitals, and small-quantity generators, such as outpatient clinics, medical and dental offices, long-term and sub-acute care facilities, and veterinary offices. Stericycle, Inc. and its subsidiaries serve approximately 522,000 customers worldwide. Of these, approximately 16,000 are large-

quantity generators while approximately 506,000 are small-quantity generators.

- 6. Stericycle's parent company opened its biomedical waste treatment facility in Morton, Washington in January 1992. In 1995, Stericycle obtained permanent state-wide authority to collect and transport biomedical waste for treatment at the Morton facility.
- 7. When Stericycle sought Commission authority there was an acute need for any alternative to incineration for the treatment and disposal of biomedical waste. Incineration was viewed by generators as harmful to public health due to toxic air emissions and problems related to the disposal of potentially toxic incinerator ash. Generators also needed services that reduced liability by maintaining tighter control over biomedical waste until it was treated. Specifically, generators expressed a need for service by a single company from collection to treatment and disposal. Finally, generators expressed need for training related to waste segregation and safety.
- 8. Stericycle met these needs by treating biomedical waste at its own treatment facility through a heat treatment process to render it non-infectious. Incineration was limited to pathological wastes, such as bodily tissue and cultures. Stericycle also offered customer audits and training to improve safety and waste segregation.
- 9. Stericycle also spent several years attempting to develop a viable program to recycle raw materials that could be extracted from treated waste. Stericycle's recycling program targeted the sharps waste stream, which contains the highest proportion of plastics, including the disposable plastic containers in use at the time.
- 10. Sharps waste treated at the Morton treatment facility was shipped to another Stericycle, Inc. facility where it was processed to separate recyclable plastics. This process was

imperfect, however, and it was difficult to extract usable plastic material from a mixed waste stream that was often poorly segregated. As a result, the yield of recyclable plastics from this sharps waste stream was low. Stericycle estimates that in general only 10% of the sharps waste was recoverable as recyclable plastics. The plastics recovered by Stericycle were then sent to Sage, Inc. to be used in manufacturing new disposable sharps containers.

- 11. Stericycle continued this arrangement for several years but it proved not to be economically viable. The price of sharps containers manufactured using reclaimed plastics was appreciably higher than that of containers manufactured with virgin plastic stock. Customers were unwilling to buy sufficient quantities of the higher priced containers with recycled materials. Ultimately, Sage, Inc. stopped taking the plastics reclaimed by Stericycle. Stericycle stored some of the reclaimed plastics in a warehouse for several years but was unable to find a company willing to purchase the material.
- 12. Through these efforts, Stericycle concluded that there was no commercially viable use for recycled plastics from the sharps waste stream. However, Stericycle did not abandon its commitment to environmentally beneficial biomedical waste collection services. Through comprehensive use of reusable sharps containers, Stericycle has since created a method of diverting plastic waste from landfills that is far superior to the recycling of plastics from sharps waste.
- 13. Stericycle has maintained stable biomedical waste collection rates since it filed its first rate tariff in 1993. Stericycle now serves more than 7,600 customers. It has served customers in every Washington county, urban and rural, and currently serves customers in every county except one.

- 14. Stericycle offers a comprehensive and customizable suite of services to Washington biomedical waste generators. Stericycle's services are intended to improve safety and promote environmental sustainability.
- 15. Stericycle's commitment to safety takes many forms, from offering on-site waste management to employing automated technology to safely handle sharps waste during treatment. Stericycle's commitment to environmental sustainability is longstanding, from its pioneering use of heat treatment rather than incineration to its current leadership in providing 100 percent reusable sharps containers.
- Washington generators since 1995. Stericycle offers training programs to generators on the OSHA bloodborne pathogens rule, proper waste segregation to reduce the amount of biomedical waste, and safe handling of waste. Stericycle provides its biomedical waste customers with compliance training on U.S. Department of Transportation (DOT) regulations governing shipment of medical waste. Stericycle also trains its customers on Stericycle's waste acceptance protocol. Stericycle requires every new customer to sign an acknowledgement of its Waste Acceptance Protocol. Exhibit A is Stericycle's Waste Acceptance Protocol.
- 17. Stericycle provides its customers with red bags, recyclable corrugated cardboard containers, and reusable plastic collection containers in which to discard biomedical waste. Stericycle offers a variety of container options to customers in a variety of sizes. Exhibit B is a list of containers in use by Stericycle at the end of 2011, when Waste Management's application was filed. These containers include reusable plastic tubs ranging in size from 10 gallons to 48 gallons manufactured by Ballbounce, Inc., Rubbermaid, and Rehrig.

- During collection, transportation, and treatment all biomedical waste is handled exclusively by Stericycle personnel who are trained to safely perform their work. Exhibit C is a list of employee trainings and certifications currently required by Stericycle. These employee trainings include: bloodborne pathogens, waste acceptance protocol, hazardous materials, spill response, and accident and injury reporting.
- 19. At pick-up, drivers visually inspect each waste container to ensure that it is correctly filled and closed. Overfilled containers and containers in which red bags or corrugated boxes are not properly closed are not accepted. Visually soiled containers are not accepted.
- 20. Stericycle also begins its proprietary, cradle-to-grave tracking and documentation program at pick-up. This program is designed to provide customers with accurate tracking, billing, waste stream information, and proof of waste treatment.
- 21. This program includes Stericycle's bar code tracking system known as Steriworks (formerly known as Bio Track). Prior to pick-up, customers place customerspecific bar code labels on each sealed container. Exhibit D is an example of the customerspecific bar code labels used in the tracking system. Stericycle's drivers use hand-held scanners to scan each container's bar code, registering the waste as collected. At transfer yards, drivers use the hand-held devices to record that they have transferred the containers to trailers that will be hauled to the processing facility. Finally, containers are scanned again just prior to treatment at the processing plant. This data provides a record that every container collected from a generator has been successfully transferred to Stericycle's processing facility and treated. Exhibit E is an example of a report from the tracking system indicating the movement

and delivery of containers. Stericycle employees reconcile each container billed to a customer against this data to confirm that it was treated.

- 22. All collections are also documented on shipping manifests filled out by
 Stericycle's drivers and signed by generators, in either electronic or hard copy. Customers sign
 electronic shipping manifests on the driver's hand-held scanner, which prints a copy retained
 by the generator. These manifests document the containers that were picked up and
 Stericycle's processing facility as their shipping destination. The customers' signature verifies
 that each shipment is properly packed for transportation in accordance with DOT regulations.
 Exhibit F is an example of Stericycle shipping manifests.
- 23. Stericycle has developed an on-line Customer Manifest Archive System available to Washington generators that stores all biomedical waste manifests in a secure, on-line data base. Generators can view and print these records at any time. This archive system ensures that generators will always have proof of shipment for treatment at their fingertips.
- 24. Stericycle's tracking and documentation services are also used to provide generators with important information about their waste. If requested, Stericycle provides generators with monthly reports that show the waste that has been collected and treated. These reports can also be configured to provide a generator with department level information, tracking monthly waste collection by specific departments at the generator's facility. These reports are tools for generators to use in reducing the biomedical waste generated in their facilities and reducing improper disposal of non-infectious waste in the biomedical waste stream. Exhibit G contains an example of a waste report.
 - 25. Stericycle maintains a fleet of approximately 30 box trucks and tractor/trailers to

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25 26 serve its 7,600 customers in Washington. Each transportation vehicle is equipped with an impermeable floor, either an overlay aluminum floor or a floor made impermeable through treatment with a sealant. Every truck and tractor/trailer is equipped with a spill kit to allow for rapid containment and cleaning of any spills. Exhibit H is a list of the contents of Stericycle spill kits. All drivers are trained in emergency response protocols specific to biomedical waste. See Exhibit C. Most Stericycle box trucks used on collection routes are equipped with hydraulic lifts to reduce driver injuries and the chance of accidental spills while loading.

- Stericycle's collection and transportation network in Washington is managed 26. locally to maximize transportation efficiency by keeping routes as short and dense as possible. This active management keeps Stericycle's costs low and its rates stable. Stericycle's routes are maintained in a hub-and-spoke pattern surrounding five transfer yards in Woodinville, Kent, Spokane, Pasco, and Portland, Oregon. Exhibit I presents two maps indicating the locations of Stericycle's transfer yards and its processing facility.
- Stericycle's route drivers follow a daily schedule of collections and terminate 27. their collection routes at one of the transportation yards. At the yard the driver transfers collected waste containers to a waiting trailer, an event that is recorded in the tracking system. Stericycle's transfer yards are secure facilities.
- The trailers to which the route drivers transfer the waste collected are driven to 28. the Morton processing facility on a planned schedule that varies at each of the transfer yards, from six 28 foot trailers and one 53 foot trailer per day to one 53 foot trailer per week. This process allows shipment by the most efficient means, reducing costs and reducing Stericycle's use of fossil fuels.

- 29. Ultimately, all biomedical waste collected in Washington is transported to Stericycle's treatment facility in Morton, Washington. Upon arrival containers are off-loaded and the trailers are swept and sprayed with bleach to disinfect the interior.
- 30. There are separate processing lines at the Morton processing facility for general biomedical waste (typically called "RMW waste") and sharps waste. RMW waste collection containers and sharps waste containers are moved to staging areas near their respective processing lines. All pathological waste, such as anatomical waste and tissue samples, is stored in a refrigerated trailer at the Morton facility before being shipped to Stericycle's Salt Lake City, Utah incinerator for disposal by incineration. The refrigerated pathological waste is transported to Salt Lake City approximately two to three times per week.
- 31. The RMW waste processing line begins with the containers being weighed and bar-code scanned in the final stage of the tracking system. These containers are also scanned for radioactive emissions to ensure that radioactive materials have not been improperly discarded with biomedical waste. Next, the containers are emptied and their contents placed into large bins. The containers are inspected for debris and damage and then proceed by conveyer to the RMW container wash line.
- 32. The RMW container wash line consists of an automated wash machine with two washing zones. In both zones the containers are washed by jets of water heated to at least 180 degrees. If the temperature falls below 180 degrees, Quaternary Ammonium is added to the wash water, at a minimum concentration of 400 ppm. In the second zone, an antimicrobial solution is added to the final wash water. At the end of the wash protocol, containers are again inspected for cleanliness. Finally, clean reusable containers are collected and returned to

customers for reuse.

- 33. The bins filled with bagged biomedical waste are transferred to a steam autoclave and processed to render the waste non-infectious. In the autoclave, biomedical waste is treated with boiler generated steam at a minimum temperature of 290 degrees and 40 psi for a minimum of 30 minutes. This treatment protocol was developed through testing by Stericycle and is tested every week by processing a spore sample with the biomedical waste. That sample is sent for testing at an independent lab to ensure the continued effectiveness of Stericycle's process.
- 34. At the end of the autoclave process, the non-infectious waste is loaded into a trailer and compacted. At least once a day a full trailer of treated, compacted waste is transported from the Morton facility to the Allied Waste landfill in Roosevelt, Washington for final disposal.
- 35. Stericycle offers a specialized program to manage and dispose of sharps waste. This program is known as the BioSystems Sharps Management program. This sharps program shares many features with Stericycle's handling and treatment of non-sharps waste but also contains additional service features and utilizes separate processing and container wash lines at the Morton facility.
- 36. Stericycle offers this unique sharps program for several reasons. First, infectious sharps present the most acute danger to hospital personnel. Reducing needle sticks through professional and proactive management of sharps waste is a large benefit to generators and their employees. Second, sharps treatment and disposal has traditionally generated a large proportion of waste simply from the use of disposable plastic sharps containers. Eliminating

this waste by employing reusable sharps containers helps generators reduce their costs and the volume of disposable waste by over half.

- 37. When a customer begins using Stericycle's sharps program, Stericycle provides an initial facility profile that documents sharps disposal locations and ensures compliance with regulations related to disposal containers and spill response procedures. Stericycle technicians then install brackets for Stericycle's proprietary reusable sharps containers at all disposal locations and create monitoring schedules tailored to generators' operational and clinical hours. Stericycle trains the facility's primary contact on proper sharps segregation and offers sharps segregation training to a facility's staff upon request.
- 38. For customers who request the service, Stericycle personnel monitor and exchange reusable sharps containers at the generator's facility as the containers are filled. Stericycle's technicians conduct routine monitoring of sharps containers so that they can be replaced before they are over-full. Exchanging sharps containers before they are over-filled dramatically reduces the risk of needle sticks resulting from use, closing and transporting sharps containers. Each generator is provided with a store of contingency containers in the event a container fills between scheduled inspections by Stericycle personnel.
- 39. Stericycle's technicians collect the sharps containers within the generator's facility on specially designed wheeled racks with locking bars. DOT regulations do not permit the sharps containers to be transported by truck without additional restraint or containment. Stericycle, therefore, developed a rack system that could be used both to safely collect sharps containers within a facility and, consistent with DOT requirements, to safely transport the sharps containers to Stericycle's processing plant.

- 40. At the time of collection, Stericycle's drivers inspect each sharps container to ensure that it is closed and correctly stored. Each rack of reusable containers is given a bar code label, scanned into the tracking system, and entered on the shipment manifest. From there, Stericycle's tracking and documentation system tracks and reports sharps waste in the same manner as non-sharps "RMW waste."
- 41. When the racks of sharps containers arrive at the Morton processing facility they are off-loaded to the staging area for the sharps processing line. Just before processing, the racks of sharps containers are weighed (the weight of the rack is automatically subtracted) and bar code scanned in the final stage of the tracking system. The containers are then individually bar code scanned to subtract the base weight of the containers and to track the number of times each container has been used. After scanning, the containers are removed from the racks and put on a conveyer system. These containers are also scanned for radioactive emissions.
- 42. The lids of all sharps containers except the largest 17 gallon containers are removed by a robotic arm and placed on a conveyer to the sharps container wash line. All opened sharps containers are decanted by an automated process. These two processes are performed by machine to remove the risk of needle sticks during processing. The rigid plastic lids of the 17 gallon containers are removed manually by sliding them from the container. This process does not require workers to reach under the lid or into the containers.
- 43. The sharps waste is loaded into large autoclave bins and then transferred to a steam autoclave and processed using the same protocol described above. The treated waste is then loaded into a trailer, compacted, and disposed of at the Allied Waste landfill.
 - 44. Empty reusable sharps containers are transferred to the sharps container wash

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25 26 line. The containers are first inspected for debris and damage and then placed on a conveyer to pass through the automated wash machine for the sharps containers. Over the course of seven minutes the sharps containers pass through a wash, rinse, and dry zone. In the wash zone the containers are sprayed with many jets of water heated to approximately 149 degrees and mixed with bleach. In the rinse zone the containers are sprayed with jets of pure water heated to approximately 189 degrees. A blow drier removes water in the dry zone. This wash machine is monitored hourly for correct water temperature, chlorine concentration, and conveyer speed.

- Clean containers are inspected again to ensure they are not soiled or damaged, 45. reassembled and returned to the sharps racks, and placed in a quality assurance staging area. Every hour, a sample of the containers in this area are quality control inspected. If any evidence of debris or damage is discovered, the past hour's batch of cleaned containers will be rejected and either the entire batch must be cleaned again, or each container is inspected and individually cleared. On a monthly basis cleaned sharps containers are tested by swab sent to an outside laboratory to ensure appropriate elimination of microorganisms.
- 46. After quality control inspection, the clean reusable sharps containers are returned to customers for reuse. Any sharps container that has been damaged or that has reached the end of its useful life is discarded after being washed.
- 47. Stericycle's proprietary reusable sharps containers are the first and only reusable sharps containers offered to Washington generators. These containers are FDA approved medical devices and, as such, the FDA maintains regulatory oversight over their use. Exhibit J is the FDA's approval of the BioSystems containers. Each BioSystems container is rated for reuse up to 600 times and Stericycle monitors each container to ensure that its reuse does not

exceed FDA approval.

- 48. On average, the reusable containers account for approximately 51% of the weight of filled containers collected by Stericycle. Exhibit K is a report of sharps container weight and sharps waste weight from January through August, 2012 indicating the percentage of the waste stream that is reused. Thus, by reusing the containers, Stericycle effectively recycles over 50% of its customers' sharps waste.
- 49. I understand from reviewing Waste Management's responses to data requests in this proceeding that in two recent months Waste Management calculated that between 17% and 28% of the sharps and sharps containers it collects under its "ecoFinity" program are reclaimed as recyclable materials. Exhibit L contains excerpts of Waste Management's discovery responses. Thus, Stericycle's reusable sharps container program diverts approximately twice as much waste from landfills as Waste Management's ecoFinity program. Moreover, the use of reusable sharps containers saves the large expenditures of energy (with associated air emissions) necessary to transport sharps waste to distant recycling facilities, reclaim plastic materials, convert them to pelletized plastics, transport the reclaimed plastics to a container maker, and process those plastics into new containers. I understand that Waste Management's sharps waste is treated and partially recycled in Southern California more than 1,200 miles from where it is collected and that reclaimed plastics are then shipped again to Becton Dickinson for incorporation into new disposable sharps containers.
- 50. Stericycle's sharps customers receive customized reports that demonstrate the amount of plastics they have diverted from landfills by using reusable sharps containers.

 Attached as Exhibit M is an example of a sharps waste diversion report. In just 2012 through

August, the sharps program has diverted 464,763 sharps containers from Washington generators, completely reusing more than 1,423,392 pounds of plastics. *See* Exhibit K.

I declare under penalty of perjury under the laws of the State of Washington and the United States that the foregoing is true and correct to the best of my knowledge and belief.

EXECUTED this 15T day of October, 2012 at Kent, Washington

Mike Philpott

GARVEY SCHUBERT
BARER
A PARTNERSHIP OF PROFESSIONAL
CORPORATIONS
eightcenth floor
1191 **second ayenue

CERTIFICATE OF SERVICE

2	I, Vickie L. Owen, certify under penalty of	perjury	under the laws of the State of
3	Washington that, on October 10, 2012, I caused to	be serv	ved on the person(s) listed below in
4	the manner shown a copy of PREFILED TESTIMO	ONY C	OF MIKE PHILPOTT:
5	Washington Utilities and		Via Legal Messenger
6	Transportation Commission 1300 S. Evergreen Park Dr. SW		Via Facsimile
7	PO Box 47250 Olympia, WA 98504-7250	X	Via FedEx
8	(360) 664-1160 records@utc.wa.gov	X	Via Email
9			
10	Administrative Law Judge	X	Via Email
11	Gregory Kopta gkopta@utc.wa.gov		
12			
13	Jessica Goldman Polly L. McNeill		Via Legal Messenger
14	Summit Law Group 315 – 5 th Avenue South		Via Facsimile
15	Seattle, WA 98104 jessicag@summitlaw.com	L	Via U.S. Mail, First Class, Postage Prepaid
16	pollym@summitlaw.com kathym@summitlaw.com	×	Via Email
17	deannas@summitlaw.com		
18	James K. Sells	П	Y7' Y 1 N
19	Attorney at Law PMB 22, 3110 Judson Street		Via Legal Messenger Via Facsimile
20	Gig Harbor, WA 98335 jamessells@comcast.net		Via U.S. Mail, First Class,
21	cheryls@rsulaw.com Attorney for Protestant WRRA, Rubatino,	×	Postage Prepaid Via Email
22	Consolidated, Murrey's and Pullman		· -·· -
23			
24			
25			

GARVEY SCHUBERT BARER

A PARTNERSHIP OF PROFESSIONAL CORPORATIONS eighteenth floor 1191 second avenue seattle, washington 98101-2939 206 464-3939

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1 2 3 4 5 6	Fronda Woods Office of the Attorney General Utilities and Transportation Division 1400 S. Evergreen Park Drive SW PO Box 40128 Olympia, WA 98504-0128 (360) 664-1225 (360) 586-5522 Fax fwoods@utc.wa.gov BDeMarco@utc.wa.gov		Via Legal Messenger Via Facsimile Via U.S. Mail, First Class, Postage Prepaid Via Email	
7				
8	Dated at Seattle, Washington this 10 th	day of Octo	bber, 2012.	
9		<i>(</i> -)		
10	•	Vickie L. O	e L Omen	***
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STERICYCLE, INC. WASTE ACCEPTANCE PROTOCOL PACIFIC NORTHWEST DISTRICT

The information contained within this STERICYCLE Protocol reflects the requirements necessary to comply with U.S. Department of Transportation (DOT) *Hazardous Materials Regulated Medical Waste* regulations and all other applicable laws and regulations governing regulated medical waste for generators, transporters, and/or treatment facilities.

It is very important for this document to be signed and returned to Stericycle, Inc. where it will be maintained in the appropriate customer file. This action will also serve to acknowledge receipt and understanding of the Stericycle, Inc. Waste Acceptance Protocol.

INSTRUCTIONS:

- Read Stericycle, Inc. Waste Acceptance Protocol information.
- Sign top page and detach from the rest of the Protocol.
- Retain Protocol for your files and reference with your "Medical Waste Management Plan".
- Return detached top page immediately to your Stericycle, Inc. District or Sales Representative.

If you have any questions or concerns regarding this document or its contents, please do not hesitate to call our Customer Service Department or your Sales Representative.

Customer		
Pacility:		
Address:		
City:	State:	Zip:
Telephone		
Name of Authorized Representative:		W.,
Citle of Authorized Representative:		
Signature of Authorized Representative:		
Date:		

Please mail signed cover sheet to:

0r

fax signed cover sheet to:

Stericycle, Inc 20320 80th Ave S Kent, WA 98032 888-234-3866



STERICYCLE, INC. WASTE ACCEPTANCE PROTOCOL PACIFIC NORTHWEST DISTRICT

CONTENT OUTLINE

- 1.0 WASTE ACCEPTED BY STERICYCLE
- 2.0 PHARMACEUTICAL WASTE
- 3.0 NON-CONFORMING WASTES NOT ACCEPTED BY STERICYCLE
- 4.0 SEGREGATION AND PACKAGING OF WASTE
- 5.0 LABELING AND MARKING OF MEDICAL WASTE CONTAINERS/BAGS
- 6.0 REUSABLE VS DISPOSABLE CONTAINERS AND DECONTAMINATION
- 7.0 STORAGE OF MEDICAL WASTE
- 8.0 TRACKING DOCUMENTS FOR MEDICAL WASTE
- 9.0 TRANSPORTATION OF WASTE
- 10.0 TREATMENT OF WASTE
- 11.0 LAWS, REGULATIONS AND POLICIES FOR MEDICAL WASTE

STERICYCLE CONFIDENTIAL INFORMATION FOR CUSTOMERS

This document is provided for your education and notification of procedures required to assure compliance with applicable regulations.

This document shall not be reproduced, copied, loaned, or transferred to another person directly or indirectly, in whole or in part, nor used for any purpose other than that for which it is specifically furnished, without the prior written consent of STERICYCLE.



STERICYCLE, INC. WASTE ACCEPTANCE PROTOCOL

1.0 WASTE ACCEPTED BY STERICYCLE

- 1.1 Regulated Medical Waste (Biohazardous and Sharps Waste)
 - 1.1.1 Stericycle, Inc. accepts those wastes generated in the diagnosis, treatment, and immunization of humans and animals or related research, in the production and testing of biologicals, and in the preparation and administration of antineoplastic/cytotoxic agents. Stericycle's waste acceptance practices and policies are based upon federal, state, and local laws for regulated medical waste. For the purposes of this document, the term regulated medical waste also means biohazardous, biomedical, or infectious waste.
 - 1.1.2 Laboratory wastes including, but not limited to:
 - 1.1.2.1 Human or animal specimen cultures from medical and pathology laboratories.
 - 1.1.2.2 Cultures and stocks of infectious agents from clinical, research, and industrial laboratories (CDC Biohazard Levels or Risk Groups I, II, III).
 - 1.1.2.3 Wastes from production of bacteria, viruses, spores, discarded vaccines and biologicals from healthcare or research, and dishes or devices used to transfer, inoculate, and mix cultures.
 - 1.1.3 Human surgical specimens, tissues, organs, placentas, and limbs (pathology waste only, exclusive of preservative agents).
 - 1.1.4 Animal parts, tissues, fluids, and carcasses from both research and veterinary facilities (pathology waste only, exclusive of preservative agents).
 - 1.1.5 Fluid blood, fluid blood products, and containers, equipment, or articles with fluid blood/blood products/body fluids.
 - 1.1.6 Medical waste contaminated with excretions, exudates, secretions, or body fluids including, but not limited to, isolation waste, or other medical waste as determined by the infection control staff, physician, veterinarian, or local health officer to be isolated and handled as such.
 - 1.1.7 Chemotherapy waste including sharps, syringes, IV tubing/bags/bottles, vials, and other discarded contaminated items generated in the preparation and

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administration of cytotoxic/antineoplastic drugs. Only *empty* containers/bags are acceptable with residue not to exceed 3% of total volume.

- 1.1.8 Sharps waste including, but not limited to:
 - 1.1.8.1 Suture needles, hypodermic needles, syringes, needles with attached tubing, scalpel and razor blades, dental wires, disposable surgical instruments, and electrosurgical needles/blades.
 - 1.1.8.2 Medical/laboratory glassware or plasticware that may be contaminated with an infectious substance and is able to cut or penetrate skin or packaging such as needles, syringes, scalpels, broken glass, slides, pipettes, blood tubes, vials, bottles, culture slides or dishes, broken rigid plastic, exposed ends of dental wire and contaminated unbroken glass articles that could be broken during handling and transportation, thus rendering them sharps waste.
- 1.2 International waste from ocean liners, ships, and planes which is not otherwise hazardous. The U.S. Department of Agriculture (USDA) regulates this waste.
- 1.3 Medical Records and Confidential Documents

2.0 PHARMACEUTICAL WASTE

2.1 Definitions of pharmaceutical waste (aborted doses, illegal drugs, controlled, legend or OTC drugs, expired or unopened pharmaceuticals, samples) have undergone increased scrutiny at several agency levels and are typically being re-defined as dangerous waste by state regulations in conjunction with RCRA regulations and criteria. As such, this type of material precludes inclusion into the normal regulated medical waste stream. For disposal of pharmaceutical wastes, please contact your Stericycle representative for more information regarding the Mail Back Program for Pharmaceutical Waste.

3.0 NON-CONFORMING WASTES NOT ACCEPTED BY STERICYCLE

- 3.1 Improper Packaging: The generator shall not tender and Stericycle, Inc. shall not knowingly accept for transportation any container which:
 - a. is not sealed and properly labeled;
 - b. is punctured or materially damaged;
 - c. is overfilled or overweight (see below);
 - d. contains anything other than biomedical waste;
 - e. contains radioactive materials as defined by the U.S. Nuclear Regulatory Commission; or
 - f. requires special handling specified by the generator.

Container Maximum Weights:

35 pounds	50 pounds	<u>60 pounds</u>
Small Tub (10 gal)	Small/Med Tub	Med/Large Tubs
	(21 gal)	(28 gal)
40 Pounds	Medium Tub	(32 gal)
Small Box (15 gal)	(20 gal)	Large Tubs
Med/Large Box		(40 gal)
(33 gal)		(48 gal)

The following charges will be assessed for improperly packaged containers or for containers requiring special handling:

\$2.00 PER GALLON IN ADDITION TO ALL OTHER RATES AND CHARGES

3.1 Radioactive Waste: UNDER NO CIRCUMSTANCES, will Stericycle, Inc. knowingly accept any waste emitting radiation in levels greater than regulatory limits. Prior to treatment, all containers will be inspected by means of a radiation monitor. Any container above regulatory limits will be rejected for treatment, and arrangements will be made for return to the generating facility. This procedure may be result in added service charges.

3.2 Hazardous Waste, including, but not limited to:

- 3.2.1 Solvents, paints, paint thinner
- 3.2.2 Drums or other containers with hazard warning sign
- 3.2.3 Batteries of any kind
- 3.2.4 Glass thermometers, sphygmomanometers, florescent light tubes, or other medical equipment or devices containing mercury such as bougie dilators and GI tubes with mercury pouches. This includes any dental material or equipment that may contain mercury.
- 3.2.5 Chemicals such as formaldehyde/formalin, ova-parasite fixative, acids, alcohol, acetone, waste oil, items preserved in thimerosal in concentrations exceeding 0.002%, and mercury-containing reagents.
- 3.2.6 Bulk chemotherapy waste (antineoplastic/cytotoxic drugs) or other RCRA listed hazardous pharmaceuticals full or partially full IV bottles/bags and vials of chemotherapy agents constitute hazardous waste and must be managed accordingly. Stericycle, Inc. will not accept any outdated or unused chemotherapy drugs. It is recommended that such agents be returned to the pharmaceutical company.
- 3.2.7 Any item listed as being hazardous in federal, state, or local regulations.
- 3.2.8. In order to comply with state and federal regulations, all hazardous waste must be managed by a licensed hazardous waste contractor. Stericycle, Inc. does not provide service for hazardous waste disposal. Each facility should contact their state or local regulatory agency for hazardous waste regulations and information.
- 3.3 Compressed Gas Cylinders, Canisters, Inhalers, and Aerosol Cans
- 3.4 Human Remains: Stericycle, Inc. requires human remains, fetuses/products of conception and cadavers (intact and otherwise) are segregated from the medical waste stream. Stericycle, Inc. will not accept these materials.

4.0 SEGREGATION AND PACKAGING OF WASTE

4.1 Non-Sharp Biohazardous Medical Waste

4.1.1 All non-sharp biohazardous waste must be segregated at the point of origin and placed into at least one biohazard bag which is certified by the manufacturer as LLDPE with a minimum thickness of 1.5 mil, or equivalent, meeting ASTM D

1709-97 (Free-falling dart test) and ASTM D 1922-94a (Tear Resistance Test). Bags must be red in color, except for chemotherapy and pharmaceutical biohazardous waste. Bags must be twisted and tied in a single knot to prevent leakage or expulsion of contents.

4.1.2 The containment of waste in "autoclavable" bags is acceptable, but not required for waste that is treated via ETD or autoclaved off-site by Stericycle. If used, these bags must be red in color.

4.2 Sharps Waste - Needles, Syringes, Blades, Medical Glassware

- 4.2.1 All sharps waste must be segregated at the point of use and placed in rigid, puncture-resistant containers which when sealed are leak resistant and cannot be easily opened. Sharps containers must be certified by the Food and Drug Administration (FDA). Sharps containers are then placed in a red bag, tied in a single knot and placed into a STERICYCLE container.
- 4.2.2 Care should be taken not to overfill sharps containers in order to avoid associated hazards. Sharps containers to be removed and exchanged when they are 3/4 full.

4.3 Body Fluids, Suctioned Fluids, and Other Non-chemical Fluids

- 4.3.1 Fluids not absorbed within other waste materials, such as sponges or dressings, must be placed within leak-resistant, break-resistant containers that are tightly lidded or capped for fluids in quantities greater than 20 cubic centimeters to prevent leakage during handling and transportation.
- 4.3.2 The discharge of liquid and semi-liquid wastes to a public sewage system is permissible where not prohibited by local ordinance, except for hazardous waste, laboratory waste, and microbiological specimens.

4.4 Containment of Biohazardous and Sharps Waste Prior to STERICYCLE Collection

Medical waste contained as described in 4.1 through 4.3 must be placed by the generator into properly lidded and secured STERICYCLE container for transport off site, depending upon the types of waste and specific requirements of the Stericycle, Inc. district providing the service. At a minimum, all medical waste, including sharps containers, must be secured in at least one red biohazard bag or liner and placed into a reusable plastic tub or disposable fiberboard container with the top of the container secured. Note: Cultures and stocks cannot be discarded into sharps containers unless the container is then placed into a securely tied (single knot) red bag and contained inside a lidded container certified for DOT Packaging Group II requirements.

4.5 Segregation and Containment for Specific Treatment Requirements

- 4.5.1 Chemotherapy waste, pathology waste (human organs, limbs, and surgical specimens, and animal parts, tissues, and carcasses), and non-hazardous pharmaceuticals require incineration, whereas other medical waste and sharps waste are acceptable to either ETD, steam autoclave processing or incineration.
- 4.5.2 Chemotherapy waste Pathology waste, and non-hazardous pharmaceuticals must be segregated and packaged into designated containers, separate from other biohazardous waste and sharps waste, in order to ensure appropriate treatment methods for specific wastes.

- 4.5.2.1 In order to ensure safe handling, treatment by incineration, and proper spill clean-up techniques of chemotherapy waste must be segregated from other medical waste into containers labeled "CHEMOTHERAPY WASTE".
- 4.5.2.2 Pathology waste must be segregated from other biohazardous waste into containers labeled "PATHOLOGY WASTE" and incinerated. It is necessary for preservative agents to be decanted from pathology waste prior to being packaged for collection and treatment, as preservatives are specifically excluded from biohazardous waste and must be regarded as hazardous.
- 4.6 International Waste: Shipboard and airline waste must be secured in trash bags and placed into leak-resistant STERICYCLE containers prior to transport as directed by local port authority Compliance Agreement. All containers must be labeled "FOREIGN GARBAGE" to ensure the waste is processed as required by steam autoclave

5.0 LABELING AND MARKING OF MEDICAL WASTE BAGS AND CONTAINERS

- 5.1 Biohazard Bags: Bags must be red in color and labeled with the word "BIOHAZARD" and the international biohazard symbol OR with the words "BIOHAZARD WASTE."
- **Sharps Containers:** Containers must be labeled with the words "SHARPS WASTE" or with the word "BIOHAZARD" and the international biohazard symbol.
- 5.3 Secondary Containers provided by STERICYCLE
 - 5.3.1 All containers provided by STERICYCLE will be labeled in accordance with applicable federal, state, and local regulations, with the word "BIOHAZARD", the international biohazard symbol, the words "REGULATED MEDICAL WASTE, UN 3291", and the name and address for the medical waste facility collecting the waste. In addition, generators are required to use appropriate containers for chemotherapy waste, pathology waste, and non-hazardous pharmaceutical waste as described in section 4.5.
 - 5.3.2 If applicable, STERICYCLE BIOTRACK® bar codes may be placed on each container providing identification, tracking, billing and proper treatment of medical waste transported off site by each generating facility serviced by Stericycle, Inc.
- Non-conforming Labels and Markings
 Any container which bears a label with the words or symbols reflecting "HAZARDOUS CHEMICALS", "HAZARDOUS DRUGS", "HAZARDOUS WASTE",
 "RADIOACTIVE MATERIAL", or "RADIOACTIVE WASTE" cannot be transported, accepted, or treated by Stericycle, Inc, no matter what the contents.

6.0 REUSABLE VERSUS DISPOSABLE WASTE CONTAINERS - DECONTAMINATION

- 6.1 Reusable Plastic Tubs and Lids: Used containers and lids are decontaminated by exposure to a tub washing process utilizing approved disinfectants and hot water at the STERICYCLE treatment facility before being returned to customers as required.
- **Disposable Fiberboard Boxes:** These containers and their contents are incinerated, autoclaved or processed by ETD. The treatment process is dictated by the type of waste (autoclave, ETD versus incineration).

6.3 Disposable Sharps Containers: All containers and their contents are incinerated, autoclaved or ETD. Treatment is dictated by the type of waste (chemotherapy sharps waste is incinerated).

7.0 STORAGE OF MEDICAL WASTE

- 7.1 Dedicated Storage Enclosure Customer /Generator Site
 - 7.1.1 Medical waste to be collected by Stericycle, Inc. shall be maintained in an enclosure or designated accumulation area that is secured to deny access to unauthorized persons, marked with warning signs, and provides protection from animals, rodents, insects, and natural elements.
 - 7.1.2 Warning signs shall read in English and in Spanish:

"CAUTION - BIOHAZARDOUS WASTE STORAGE AREA -UNAUTHORIZED PERSONS KEEP OUT" "CUIDADO - ZONA DE RESIDUOS - BIOLOGICOS PELIGROSOS -PROHIBIDA LA ENTRADA A PERSONAS NO AUTORIZADAS"

7.1.3 Waste to be transported off site shall not be subjected to trash chutes, compaction, or grinding prior to collection/treatment by Stericycle, unless prior approval has been obtained from local enforcement officials.

8.0 TRACKING DOCUMENTS FOR MEDICAL WASTE

- 8.1 STERICYCLE BIOTRACK® System
 - 8.1.1 All waste transported from the generating facility/person for treatment will be accompanied by a tracking document.
 - 8.1.2 STERICYCLE may use a unique approach to the tracking process known as BIOTRACK®. Bar codes and optical scanners or "readers" record all pertinent data.
 - 8.1.3 Tracking documents will include, but are not limited to, the following:
 - 8.1.3.1 Name, address, telephone number, and registration number of Hazardous Waste Transporter
 - 8.1.3.2 Type and quantity of medical waste transported
 - 8.1.3.3 "Regulated Medical Waste, 6.2, UN3291, PG II"
 - 8.1.3.4 Name, address, and telephone number of generator
 - 8.1.3.5 Generator Certification: "This is to certify that the above-name materials are properly classified, described, packaged, marked, and labeled, and are in proper condition for transportation, according to the applicable regulations of the Department of Transportation." with the signature of an authorized representative, or printed name on signature line with original signature card/certification statement on file as defined on the cover page of this document.

- 8.1.3.6 Name, address, telephone number, permit number, and the signature of an authorized representative of the permitted medical waste treatment facility receiving the waste.
- 8.1.3.7 The date the medical waste is collected from the generator's facility, the date the waste is received by the transfer station (as applicable), and the date the waste is received by the treatment facility.
- 8.1.3.8 Stericycle, Inc. Emergency Response Telephone Number (800) 424-9300
- 8.1.4 A copy of the tracking document(s) will be provided to the customer at the time of waste collection bearing signatures of the generator and the driver.
- 8.1.5 The tracking document(s) will be in the custody of the STERICYCLE driver hauling the medical waste to its treatment destination at all times.
- 8.1.6 Documentation will be mailed to the customer by Stericycle, Inc. on a monthly basis detailing receipt/treatment of medical waste collected.
- 8.1.7 Stericycle, Inc. will keep signed copies of all tracking documents for at least 3 years.

9.0 TRANSPORTATION OF WASTE

9.1 Registered Vehicles

9.1.1 Stericycle, Inc. maintains a Hazardous Waste Hauler's Registration for transportation of all medical waste collected and/or transferred and/or treated

9.2 Responsibility and Authority of STERICYCLE Drivers

- 9.2.1 STERICYCLE drivers are responsible for the collection and tracking of all waste containers generated on their assigned routes on any given day. They are responsible for monitoring the proper containment, closure, and labeling of each container prior to scanning/entering the specific data into the STERICYCLE BIOTRACK® system. The driver will also leave the appropriate number of empty clean waste containers/lids for each customer serviced, as well as a copy of the signed tracking document(s).
- 9.2.2 STERICYCLE drivers are authorized to reject any containers that do not meet DOT specifications. Odor, leakage, bulging or damaged containers, improper packaging, incorrect labels, non-conforming waste, and improper segregation are some of the causes for rejection of medical waste containers.
- 9.2.3 Containers may be subject to an off-specification charge for repackaging and special handling, if such is required.

9.3 Emergency Spill Response

9.3.1 Stericycle, Inc. Hazardous Waste Registered vehicles are equipped with emergency spill kits, and drivers are trained in emergency response spill procedures and U.S. DOT regulations. Written Emergency Response Spill and Hazardous Materials Procedures are available in the cab of each vehicle.

- 9.3.2 Stericycle, Inc. provides and maintains an Emergency Spill Response Telephone Number 24 hours a day at (800) 424-9300.
- 9.4 All policies and practices for transportation of medical waste provided by Stericycle, Inc. are in full compliance with applicable U.S. DOT and local laws and regulations.

10.0 TREATMENT OF WASTE

10.1 Permitted Waste Treatment Facilities: All waste collected by Stericycle, Inc. is transported to a permitted facility for proper treatment by ETD, autoclave or incineration for subsequent disposal.

10.2 Waste Treatment Methods/Parameters

- Pathology waste (human tissue specimens, organs, limbs, and contaminated animal carcasses, parts and specimens) is incinerated.
- 10.2.2 Trace chemotherapy contaminated waste is incinerated.
- 10.2.3 Non-hazardous pharmaceutical waste is incinerated.
- 10.2.4 Biohazardous waste and sharps waste is subjected to steam autoclave processing, ETD or incineration. Chemotherapy sharps waste is incinerated.
- 10.2.5 International waste is subjected to ETD, autoclave or incineration processing per Compliance Agreement with the local port authority or U.S. Dept. of Agriculture.
- 10.3 Waste treatment facilities operate in compliance with all applicable federal, state, and local laws/regulations and maintain all required permits and licenses.

11.0 LAWS, REGULATIONS AND POLICIES FOR MEDICAL WASTE

11.1 Transportation Authority - W.U.T.C.:

Washington Administrative Code WAC 480-70 regulates the packaging and containment, transportation to treatment and disposal facilities, shipping papers, insurance and operational requirements as enforced by the Washington Utilities and Transportation Commission

11.2 Transportation and Collection Authority – Local:

Many county health departments regulate biomedical waste collection and transportation as part of the county solid waste regulations. Contact your local County Health Department for specific County regulations.

11.3 Bloodborne Pathogen Standard -WISHA/OR-OSHA/OSHA

11.3.1 29 CFR 1910.1030 defines the Bloodborne Pathogen Standard which became effective in 1992. In the state of Washington, the Bloodborne Pathogen Standard is defined in Part J, WAC 296-62-08001, and in Oregon, it is found in OAR 437, Division 2, Subdivision Z, Bloodborne Pathogens (1920-1030). WISHA/OR-OSHA/OSHA regulate "blood and potentially infectious material" and the handling, containment, labeling and storage of "regulated waste" through the Bloodborne Pathogens Standard. The standard requires the utilization of "Universal Precautions" (Center for Disease Control Guideline—1989) in managing all blood, certain body fluids and containment materials as "potentially infectious". It includes specific requirements pertaining to personal

protective equipment (PPE), houskeeping, exposure control, engineering and work practice controls, recordkeeping, signs and labels, training, Hepatitis B vaccination, and exposure follow up, all of which must be defined in a written "Exposure Control Plan". Stericycle is regulated by the "Bloodborne Pathogen Standard", as are all healthcare providers and certain other employers with potential for exposure to blood and body fluids in the workplace. Stericycle maintains compliance with the standard under a comprehensive "Exposure Control Plan" and training program developed specifically for our medical waste drivers, medical waste handlers and equipment operators.

- "Potentially infectious materials" includes human blood/blood components and products, as well as semen, vaginal secretions, cerebrospinal, pleural, peritoneal, pericardial and amniotic fluids, saliva in dental procedures, body fluids visibly contaminated with blood such as saliva or vomitis, unfixed tissues or organs, and all body fluids where it is difficult or impossible to differentiate.
- 11.3.3 "Regulated waste", as defined in the standard, includes contaminated sharps, and liquid/semi-liquid blood or other potentially infectious material (OPIM), contaminated items that would release blood or OPIM in a liquid or semi-liquid state if compressed, items that are caked with dried blood or OPIM and are capable of releasing these materials during handling, and pathological and microbiological wastes containing blood or OPIM.
- 11.3.4 Copies of the *Bloodborne Pathogen Standard* are available through WISHA, OR-OSHA or OSHA or from your local Stericycle district providing medical waste service.
- 11.4 Chemotherapy waste is regulated by the Federal Resource Conservation and Recovery Act (RCRA), along with state of Washington Department of Ecology (WDOE) Hazardous Waste Rules.
- 11.5 Hazardous Materials Regulations for Regulated Medical Waste US Department of Transportation

U.S. Department of Transportation (DOT) Hazardous Waste Regulations (HMR; 49 CFR Parts 171-180) regulate the transportation, packaging, labeling, tracking, and training for drivers of "Regulated Medical Waste" ("RMW"). In the state of Washington, W.U.T.C. is the regulatory authority for DOT HMR for "RMW". Stericycle medical waste tracking forms and training standards reflect these requirements. The waste categories listed in section 1.0 of this Protocol include wastes defined as "RMW".

Stericycle Containers Currently in Use

- 15 gal. corrugated box (WS/WY19) Manufacturer RockTenn box dimension used since at least 1995
- 33 gal. corrugated box (WS/WY43) Manufacturer RockTenn box dimension used since at least 1995
- 30 gal. (approx.) corrugated pharma box (RXBI) Manufacturer RockTenn used since 2008
- 30 gal. (approx.) corrugated sharps box (KRB3) Manufacturer RockTenn used since 2011
- 21 gal. reusable plastic SteriTub (ST32) Manufacturer Ballbounce Inc. used since 1991
- 48 gal. reusable plastic SteriTub (ST75) Manufacturer Ballbounce Inc. used since 1991
- 10 gal. reusable plastic tub (TB20) Manufacturer Rubbermaid used since 1993
- 20 gal. reusable plastic tub (TB15) Manufacturer Rubbermaid used since 1993
- 28 gal. reusable plastic tub (TB04) Manufacturer Rubbermaid used since 1993
- 40 gal. reusable plastic tub (TB05) Manufacturer Rubbermaid used since 1993
- 28 gal. reusable plastic tub (TY04) Manufacturer Rubbermaid used since 2002
- 32 gal. reusable plastic tub (TH32) Manufacturer Rehrig used since 2011
- 43 gal. reusable plastic tub (TH43) Manufacturer Rehrig used since 2011
- 2 gal. reusable red plastic sharps container Manufacturer Xten used since 2004
- 3 gal. reusable red plastic sharps container Manufacturer Xten used since 2004
- 4 gal. reusable red plastic sharps container Manufacturer Xten used since 2009
- 8 gal. reusable red plastic sharps container Manufacturer Xten used since 2004
- 17 gal. reusable red plastic sharps container Manufacturer Xten used since 2004
- Aluminum sharps container rack (KR59/KR65) Manufacturer Xten used since 2004 2 gal. reusable blue plastic pharma container Manufacturer Xten used since 2004
- 8 gal. reusable blue plastic pharma container Manufacturer Xten used since 2004

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			Hazard Communication
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			Drivers Certification of Violations/Suspension
			MVR/Driving Record Copy
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			Accident and Injury Reporting Requirements
			Racking Properties
			Bloodborne Pathogens
			DOT CSA Training
			DOT Hazardous Materials

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Driver Qualifications Emergency Action Plans Fire Extinguishers - Portable Hazard Communication Hours of Service Personal Protective Equipment - PPE Proper Lifting Receipt of FMC Safety Regulations - Drivers Slip, Trip, Fall or Walking/Working Surfaces Waste Acceptance Protocol Wellness Training for Drivers Whistleblower Policy and Procedures	Blood Chemistry CBC Hepatitis B Compliance Hepatitis C Status Physical Exam/History Urinalysis	Access to Exposure and Medical Records Accident and Injury Reporting Requirements Bloodborne Pathogens DOT Hazardous Materials Electrical Arc Flash Training Emergency Action Plans Eyewash and Emergency Shower Fire Extinguishers - Portable Hand and Power Tools
	Medical	Training

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Hazard Communication	Lockout/Tagout Control Haz Energy - Authorized	Machine Guarding	Personal Protective Equipment - PPE	Proper Lifting	Slip, Trip, Fall or Walking/Working Surfaces	Spill Response	Blood Chemistry	CBC	Hepatitis B Compliance	Hepatitis C Status	Physical Exam/History	Urinalysis	Access to Exposure and Medical Records	Accident and Injury Reporting Requirements	Bloodborne Pathogens	DOT Hazardous Materials	Electrical Arc Flash Training	Emergency Action Plans	Eyewash and Emergency Shower	Fire Extinguishers - Portable	Hazard Communication		Lockout/Tagout Control Haz Energy - Affected	Machine Guarding	Permit Required Confined Spaces - Affected
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Proper Lifting	Respiratory Protection	Slip, Trip, Fall or Walking/Working Surfaces	Tuberculosis Training	Waste Acceptance Protocol	Annual Review of MVR / Driving Record	DOT Employment Application	DOT Medical Examiners Certificate	Drivers Certification of Violations/Suspension	MVR/Driving Record Copy	Past Employer Safety Performance History	Photo Capy of Drivers License	Road/Driving Test	Valid CDL (all classes)	DOT Substance Abuse Testing	Physical Exam/History	DOT Alcohol/Controlled Substance Abuse - Drivers	Driver Qualifications	Hours of Service	Receipt of FMC Safety Regulations - Drivers	Vehicle Condition Reports - VCRs	Wellness Training for Drivers	Whistleblower Policy and Procedures
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10 Days Of 10 Days Of ion Date uployment 2 Years 10 Days Of 10 Day	
Once Every Year Within 30 Days Of Hire Every Year Once	e Date

Date

Date

Date

Date:

Date

6029350-001-00A53LD

Generator

Stericycle, Inc. 20320 80th Ave S Kent, WA 98032-1258

Ph: (425)291-9324 Fx: (425)291-9329

Transporter

574411

#15

CTR-11

C1411

Stericycle, Inc. 20320 80th Ave South Kent, WA 98032 (425) 291-9322

Thank You for Choosing Stericycle

UN 3291, Regulated Medical Waste, n.o.s

6029350-001-00A53LG

Generator

Stericycle, Inc. 20320 80th Ave S Kent, WA 98032-1258 Ph: (425)291-9324 Fx: (425)291-9329

Transporter

Stericycle, Inc. 20320 80th Ave South Kent, WA 98032 (425) 291-9322

Thank You for Choosing Stericycle

UN 3291, Regulated Medical Waste, n.o.s

6029350-001-00A53LJ

Generator

Stericycle, Inc. 20320 80th Ave S Kent, WA 98032-1258

Ph: (425)291-9324 Fx: (425)291-9329

Transporter

Stericycle, Inc. 20320 80th Ave South Kent, WA 98032 (425) 291-9322

Thank You for Choosing Stericycle

UN 3291, Regulated Medical Waste, n.o.s

6029350-001-00A53LM

Generator

Stericycle, Inc. 20320 80th Ave S Kent, WA 98032-1258 Ph: (425)291-9324 Fx: (425)291-9329

Transporter

Stericycle, Inc. 20320 80th Ave South Kent, WA 98032 (425) 291-9322

Thank You for Choosing Stericycle

EXHIBIT D

UN 3291, Regulated Medical Waste, n.o.s

6029350-001-00A53LE

Generator

Stericycle, Inc. 20320 80th Ave S Kent. WA 98032-1258

Ph: (425)291-9324 Fx: (425)291-9329

CT-19-11

CTR-11

CT-R-11

CT-R-11

Date

Date

Date

Transporter Stericycle, Inc. 20320 80th Ave South Kent, WA 98032 (425) 291-9322

Thank You for Choosing Stericycle

UN 3291, Regulated Medical Waste, n.o.s

Generator

Stericycle, Inc. 20320 80th Ave S Kent, WA 98032-1258 Ph: (425)291-9324 Fx: (425)291-9329

Transporter

Stericycle, Inc. 20320 80th Ave South Kent, WA 98032 (425) 291-9322 Thank You for Choosing Stericycle

UN 3291, Regulated Medical Waste, n.o.s

6029350-001-00A53LK

Generator

Stericycle, Inc. 20320 80th Ave S Kent WA 98032-1258 Ph: (425)291-9324 Fx: (425)291-9329

Transporter

Date Stericycle, Inc. 20320 80th Ave South Kent, WA 98032 (425) 291-9322 Thank You for Choosing Stericycle

UN 3291, Regulated Medical Waste, n.o.s



6029350-001-00A53LN

Generator

Stericycle, Inc. 20320 80th Ave S Kent, WA 98032-1258 Ph: (425)291-9324 Fx: (425)291-9329

Transporter

Stericycle, Inc. 20320 80th Ave South Kent, WA 98032 (425) 291-9322

Thank You for Choosing Stericycle

Date

UN 3291, Regulated Medical Waste, n.o.s

6029350-001-00A53LF

Generator

Stericycle, Inc. 20320 80th Ave S Kent, WA 98032-1258

Ph (425)291-9324 Fx (425)291-9329

Transporter

Stericycle, Inc. 20320 80th Ave South Kent, WA 98032 (425) 291-9322

Thank You for Choosing Stericycle

UN 3291, Regulated Medical Waste, n.o.s

6029350-001-00A53LI

Generator Stericycle, Inc.

20320 80th Ave S Kent, WA 98032-1258

Ph: (425)291-9324 Fx: (425)291-9329

Transporter

4

CHRII

CFF11

Stericycle, Inc. 20320 80th Ave South Kent, WA 98032 (425) 291-9322

Thank You for Choosing Stericycle

UN 3291, Regulated Medical Waste, n.o.s

6029350-001-00A53LL

Generator

Stericycle, Inc. 20320 80th Ave S Kent, WA 98032-1258

Ph (425)291-9324 Fx (425)291-9329

Transporter

Stericycle, Inc. 20320 80th Ave South Kent, WA 98032 (425) 291-9322

Thank You for Choosing Stericycle

UN 3291, Regulated Medical Waste, n.o.s

6029350-001-00A53L0 Generator

Stericycle, Inc. 20320 80th Ave S Kent, WA 98032-1258

Ph: (425)291-9324 Fx: (425)291-9329

Transporter

Stericycle, Inc. 20320 80th Ave South Kent. WA 98032 (425) 291-9322

Thank You for Choosing Stericycle



Generator

Stericycle, Inc. 20320 80th Ave S Kent, WA 98032

Ph: (425)291-9324 Fx: (425)291-9329

Transporter

Stericycle, Inc. 20320 80th Ave South Kent, WA 98032 (425) 291-9322

Thank You for Choosing Stericycle

Date

Date

Date

Incinerate Only



Generator

Stericycle, Inc. 20320 80th Ave S Kent, WA 98032

Ph: (425)291-9324 Fx: (425)291-9329

Transporter

Stericycle, Inc. 20320 80th Ave South Kent, WA 98032 (425) 291-9322

Thank You for Choosing Stericycle

Incinerate Only



Generator

Stericycle, Inc. 20320 80th Ave S Kent, WA 98032

Ph: (425)291-9324 Fx: (425)291-9329

Transporter

Stericycle, Inc. 20320 80th Ave South Kent, WA 98032 (425) 291-9322

Thank You for Choosing Stericycle

Incinerate Only



Generator

Stericycle, Inc. 20320 80th Ave S Kent, WA 98032 Rb: (435)301 9334

Ph: (425)291-9324 Fx: (425)291-9329

Transporter

Stericycle, Inc. Date 20320 80th Ave South Kent, WA 98032 (425) 291-9322

Thank You for Choosing Stericycle

Incinerate Only



6029350-001-IYA53K2

Generator

Stericycle, Inc. 20320 80th Ave S Kent, WA 98032

Ph: (425)291-9324 Fx: (425)291-9329

Date

Date

Date

Date

Transporter

Stericycle, Inc. 20320 80th Ave South Kent, WA 98032 (425) 291-9322

Thank You for Choosing Stericycle

Incinerate Only



6029350-001-IYA53K5

Generator

Stericycle, Inc. 20320 80th Ave S Kent, WA 98032 Ph: (425)291-9324 Fx: (425)291-9329

Transporter

12-Y-CT-R

005-

12-Y-CT-R

12-Y-CT-R

Stericycle, Inc. 20320 80th Ave South Kent, WA 98032 (425) 291-9322

Thank You for Choosing Stericycle

Incinerate Only



6029350-001-IYA53K8

Generator

Stericycle, Inc. 20320 80th Ave S Kent, WA 98032 Ph; (425)291-9324 Fx; (425)291-9329

Transporter

Stericycle, Inc. 20320 80th Ave South Kent, WA 98032 (425) 291-9322

Thank You for Choosing Stericycle

Incinerate Only



6029350-001-IYA53KB

Generator

Stericycle, Inc. 20320 80th Ave S Kent, WA 98032

Ph: (425)291-9324 Fx: (425)291-9329

Transporter

Stericycle, Inc. 20320 80th Ave South Kent, WA 98032 (425) 291-9322

Thank You for Choosing Stericycle

Incinerate Only



6029350-001-IYA53K3

Generator

Stericycle, Inc. 20320 80th Ave S Kent, WA 98032

Ph: (425)291-9324 Fx: (425)291-9329

Date

Date

Date

Date

Transporter

12-Y-CT-R

002-

Stericycle, Inc. 20320 80th Ave South Kent, WA 98032 (425) 291-9322

Thank You for Choosing Stericycle

Incinerate Only



6029350-001-IYA53K6

Generator

Stericycle, Inc. 20320 80th Ave S Kent, WA 98032

Ph: (425)291-9324 Fx: (425)291-9329

Transporter

Stericycle, Inc. 20320 80th Ave South Kent, WA 98032 (425) 291-9322

Thank You for Choosing Stericycle

Incinerate Only



6029350-001-IYA53K9

Generator

Stericycle, Inc. 20320 80th Ave S Kent, WA 98032

Ph: (425)291-9324 Fx: (425)291-9329

Transporter

002-12-Y-CT

Stericycle, Inc. 20320 80th Ave South Kent, WA 98032 (425) 291-9322

Thank You for Choosing Stericycle

Incinerate Only



6029350-001-IYA53KC

Generator

Stericycle, Inc. 20320 80th Ave S Kent, WA 98032 Ph: (425)291-9324

Ph: (425)291-9324 Fx: (425)291-9329

Transporter

Stericycle, Inc. 20320 80th Ave South Kent, WA 98032 (425) 291-9322

Thank You for Choosing Stericycle





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REGULATED WASTE MANIFEST

CERTIFICATE OF DESTRUCTION: This is to certify that the Regulated Medical Wastes described above were treated in accordance with state and federal guidelines

Part/Type Name

DISCREPANCY INDICATION

OF MATERIALS

Print/Type Name___

DISCREPANCY INDICATION

Signature ___

OF MATERIALS

Print/Type Name

DISCREPANCY INDICATION

Signature_

OF MATERIALS

Signature_

Date

EMATERIALS

nt/Type Name

SCREPANCY INDICATION

OFMATERIALS

Prof/Type Name

DISCRIEFANCY INDICATION

HAZARDOUS MATERIAL SHIPPING DOCUMENT TRANSPORTER: Stericycle, Inc. 20320 80th Ave South Kent, WA 98032 (425) 291-9322 For Stericycle Customer Care Call 1-866-783-7422 Stericycle Customer # 6029350 Site # 001 stericycle REGULATORY #: Phone 11: Contact : SERVICE DATE: 9/24/12 10:23:45 AM SHIPPING DOCUMENT N: MOSE001233 UN3291, REGULATED HEDICAL HASTE, N.O.S... 6.2, PGI 6.2, PG11
For DOT HAZMAT Emergency Response Call:
CHEMTREC 1-800-424-9300
Customer No. 21132
TOTAL CONTAINERS COLLECTED: 1
TOTAL VOLUME COLLECTED: 4.300 CU FT SUMMARY(Cont Type) QTY 4.300 BXO6 30 Gal Box Disposal 1 00A0987 BX06 I hereby declare that the contents of this consignment are fully and accurately described above by the proper shipping name, and are classified, packaged, marked and labelled/placarded, and are in all respects in proper condition for transport according to applicable international and national governmental regulations. GENERATOR PRINT NAME Generator Signature Transporter Certification: Driver, Rel Transporter Signature [X] THIS IS A THROUGH SHIPHENT. [X] LOCAL TRANS CENTER Stericycle-Kent, WA DESTINATION FACILITY: Incinerate Only Stericycle-North Salt Lake,UT 3A-448/JA-36 Pharmaceutical Stericycle-Dacono,CO Standard X Stericycle-Morton,WA DATE OF RECEIPT AT TREATMENT FACILITY: DELIVERY DOCUMENT #: PDSE001233 TOTAL DELIVERED ITEMS: 2 ITEM QTY 8M32 32 Gal Red Bag 2 DRIVER: Driver, Relief FREQUENCY: N/A NEXT PICKUP: N/A

CUSTOMER SERVICE: (866) 783-7422 Thank you for choosing Stericycle

EXHIBIT G



Sample Container Detail Report

Run Date: 2012-07-16

Right Straine) Area: West	CustiD: Be	egin Period: 11206	End Period: 201206	Unit of Measure: LB
Name WEST	Address 1 Cl	lty	State	Zip :
6029188	Name Ac Customer Name and Address	ddress 1	City	State Zip
Sides ID 002 003	Name Ar Customer Sites with Names an	ddress 1 nd Addresses	City	State Zip
004 008 009				

tivo cers		
SiteID	Number	Date
002	3001911138	2012-06-30
003	3001911138	2012-06-30
004	3001911139	2012-06-30
008	3001911139	2012-06-30
009	3001911139	2012-06-30

Report Release #:

StARS.Reports 1.3.1105.0

Report BuildDate:

10/17/2006

Sample

Container Detail-Weight Summary

2012-06-30 300191138 2012-06-04 MDSE00DFJO 2012-06-30 300191138 2012-06-07 MDSE00DGJP 2012-06-30 300191138 2012-06-07 MDSE00DGJP 2012-06-30 300191138 2012-06-17 MDSE00DGJP 2012-06-30 300191138 2012-06-11 MDSE00DGZQ 2012-06-30 300191138 2012-06-14 MDSE00DGZQ 2012-06-30 300191138 2012-06-14 MDSE00DJCB 2012-06-30 300191138 2012-06-18 MDSE00DJCB 2012-06-30 300191138 2012-06-28 MDSE00DJWS 2012-06-30 300191138 2012-06-28 MDSE00DJWS 2012-06-30 300191138 2012-06-28 MDSE00DJWS 2012-06-30 300191138 2012-06-24 MDSE00DJWS 2012-06-30 300191138 2012-06-44 MDSE00DJWG 2012-06-30 300191138 2012-06-44 MDSE00DJWG 2012-06-30 300191138 2012-06-41 MDSE00DJWG 2012-06-30 300191139 2012-06-41 MDSE00DJWG 2012-06-30 300191139 2012-06-44 MDSE00DJWG	Site Involce Date	Invoice #	Service Date	Manifest ID	Containers	Total Weight
2012-06-30 3001911138 2012-06-04 MDSE00DFJO 2012-06-30 3001911138 2012-06-07 MDSE00DGIP 2012-06-30 3001911138 2012-06-07 MDSE00DGIP 2012-06-30 3001911138 2012-06-11 MDSE00DGZQ 2012-06-30 3001911138 2012-06-14 MDSE00DGZQ 2012-06-30 3001911138 2012-06-14 MDSE00DHYY 2012-06-30 3001911138 2012-06-14 MDSE00DHYY 2012-06-30 3001911138 2012-06-14 MDSE00DICB 2012-06-30 3001911138 2012-06-21 MDSE00DJK8 2012-06-30 3001911138 2012-06-21 MDSE00DJK8 2012-06-30 3001911138 2012-06-28 MDSE00DJK8 2012-06-30 3001911138 2012-06-04 MDSE00DJK8 2012-06-30 3001911138 2012-06-04 MDSE00DJK8 2012-06-30 3001911138 2012-06-04 MDSE00DJK8 2012-06-30 3001911138 2012-06-11 MDSE00DJK9 2012-06-30 3001911138 2012-06-11 MDSE00DJHG 2012-06-30 3001911138 2012-06-11 MDSE00DJHG 2012-06-30 3001911138 2012-06-11 MDSE00DJHG 2012-06-30 3001911138 2012-06-14 MDSE00DJHG 2012-06-30 3001911139 2012-06-14 MDSE00DJHG 2012-06-30 300191139 2012-06-14 MDSE00DJHG 2012-06-30 300191139 2012-06-14 MDSE00DJHG 2012-06-30 300191139 2012-06-14 MDSE00DJHG 2012-06-30 3001911139 2012-06-14 MDSE00DJHG 2012-06-30 300191139 2012-06-30 30		3001911138	2012-06-04	MDSE00DFJO	27.00	886.40
2012-06-30 3001911138 2012-06-07 MDSE00DGIP 2012-06-30 3001911138 2012-06-71 MDSE00DGZQ 2012-06-30 3001911138 2012-06-11 MDSE00DGZQ 2012-06-30 3001911138 2012-06-14 MDSE00DGZQ 2012-06-30 3001911138 2012-06-14 MDSE00DHYY 2012-06-30 3001911138 2012-06-18 MDSE00DHYY 2012-06-30 3001911138 2012-06-18 MDSE00DJ02 2012-06-30 3001911138 2012-06-21 MDSE00DJ02 2012-06-30 3001911138 2012-06-21 MDSE00DJ08 2012-06-30 3001911138 2012-06-28 MDSE00DJ08 2012-06-30 3001911138 2012-06-28 MDSE00DJ08 2012-06-30 3001911138 2012-06-44 MDSE00DJ09 2012-06-30 3001911138 2012-06-41 MDSE00DJ09 2012-06-30 3001911139 2012-06-41 MDSE00DJ09 2012-06-41 MDSE00DJ09 2012-06-30 3001911139 2012-06-41 MDSE00DJ09 2012-06-41 MDSE00DJ09 2012-06-30 3001911139 2012-06-41 MDSE00DJ09		3001911138	2012-06-04	MDSE00DFJO	1.00	9.80
2012-06-30 3001911138 2012-06-07 MDSE00DGIP 2012-06-30 3001911138 2012-06-11 MDSE00DGZQ 2012-06-30 3001911138 2012-06-14 MDSE00DGZQ 2012-06-30 3001911138 2012-06-14 MDSE00DHYY 2012-06-30 3001911138 2012-06-18 MDSE00DHYY 2012-06-30 3001911138 2012-06-18 MDSE00DHYY 2012-06-30 3001911138 2012-06-21 MDSE00DJZ 2012-06-30 3001911138 2012-06-21 MDSE00DJZ 2012-06-30 3001911138 2012-06-25 MDSE00DJX8 2012-06-30 3001911138 2012-06-25 MDSE00DJX8 2012-06-30 3001911138 2012-06-28 MDSE00DJX8 2012-06-30 3001911138 2012-06-30 MDSE00DJX8 2012-06-04 MDSE00DJX8 2012-06-30 3001911138 2012-06-04 MDSE00DJYG 2012-06-30 3001911139 2012-06-04 MDSE00DJYG 20		3001911138	2012-06-07	MDSE00DGIP	37.00	1,139.40
2012-06-30 3001911138 2012-06-11 MDSE00DGZQ 2012-06-30 3001911138 2012-06-14 MDSE00DHYY 2012-06-30 3001911138 2012-06-14 MDSE00DHYY 2012-06-30 3001911138 2012-06-18 MDSE00DHYY 2012-06-30 3001911138 2012-06-21 MDSE00DJ02 2012-06-30 3001911138 2012-06-21 MDSE00DJ02 2012-06-30 3001911138 2012-06-21 MDSE00DJX8 2012-06-30 3001911138 2012-06-25 MDSE00DJX8 2012-06-30 3001911138 2012-06-26 MDSE00DJX8 2012-06-30 3001911138 2012-06-26 MDSE00DJX8 2012-06-30 3001911138 2012-06-04 MDSE00DJY8 2012-06-30 3001911138 2012-06-11 MDSE00DGWA 2012-06-30 3001911138 2012-06-18 MDSE00DJHG 2012-06-30 3001911139 2012-06-25 MDSE00DJHG 2012-06-30 3001911139 2012-06-18 MDSE00DJHG		3001911138	2012-06-07	MDSE00DGIP	2.00	54.60
2012-06-30 3001911138 2012-06-11 MDSE00DGZQ 2012-06-30 3001911138 2012-06-14 MDSE00DHYY 2012-06-30 3001911138 2012-06-18 MDSE00DICB 2012-06-30 3001911138 2012-06-21 MDSE00DJ02 2012-06-30 3001911138 2012-06-21 MDSE00DJ02 2012-06-30 3001911138 2012-06-25 MDSE00DJK8 2012-06-30 3001911138 2012-06-26 MDSE00DJK8 2012-06-30 3001911138 2012-06-26 MDSE00DJK8 2012-06-30 3001911138 2012-06-04 MDSE00DJK8 2012-06-30 3001911138 2012-06-04 MDSE00DJK8 2012-06-30 3001911138 2012-06-04 MDSE00DJHG 2012-06-30 3001911138 2012-06-04 MDSE00DJHG 2012-06-30 3001911138 2012-06-18 MDSE00DJHG 2012-06-30 3001911138 2012-06-18 MDSE00DJHG 2012-06-30 3001911138 2012-06-18 MDSE00DJHG 2012-06-30 3001911138 2012-06-18 MDSE00DJHG 2012-06-30 3001911138 2012-06-25 MDSE00DJHG 2012-06-30 3001911138 2012-06-26 MDSE00DJHG 2012-06-30 3001911138 2012-06-18 MDSE00DJHG 2012-06-30 3001911139 2012-06-18 MDSE00DJHG		3001911138	2012-06-11	MDSE00DGZQ	29.00	824.80
2012-06-30 3001911138 2012-06-14 MDSE00DHYY 2012-06-30 3001911138 2012-06-18 MDSE00DICB 2012-06-30 3001911138 2012-06-18 MDSE00DICB 2012-06-30 3001911138 2012-06-21 MDSE00DJ02 2012-06-30 3001911138 2012-06-25 MDSE00DJ02 2012-06-30 3001911138 2012-06-26 MDSE00DJK8 2012-06-30 3001911138 2012-06-28 MDSE00DJK8 2012-06-30 3001911138 2012-06-04 MDSE00DJK8 2012-06-30 3001911138 2012-06-04 MDSE00DJK8 2012-06-30 3001911138 2012-06-11 MDSE00DGWA 2012-06-30 3001911138 2012-06-11 MDSE00DJHG 2012-06-30 3001911138 2012-06-18 MDSE00DJHG 2012-06-30 3001911138 2012-06-18 MDSE00DJHG 2012-06-30 3001911138 2012-06-18 MDSE00DJHG 2012-06-30 3001911138 2012-06-18 MDSE00DJHG 2012-06-30 3001911139 2012-06-19 MDSE00DJHG 2012-06-30 3001911139 2012-06-19 MDSE00DJHG 2012-06-30 3001911139 2012-06-11 MDSE00DJHG 2012-06-30 3001911139 2012-06-12 MDSE00DJHO 2012-06-30 3001911139 2012-06-19 MDSE00DJHO 2012-06-30 3001911139 2012-06-19 MDSE00DJHO 2012-06-30 3001911139 2012-06-19 MDSE00DJHO 2012-06-30 3001911139 2012-06-19 MDSE00DJHO 2012-06-30 3001911139 2012-06-26 MDSE00DJHO 2012-06-30 3001911139 2012-06-19 MDSE00DJHO 2012-06-30 3001911139 2012-06-26 MDSE00DJHO 2012-06-30 3001911139 2012-06-19 MDSE00DJHO 2012-06-30 3001911139 2012-06-26 MDSE00DJHO 2012-06-30 3001911139 2012-06-11 MDSE00DJHO 2012-06-30 3001911139 2012-06-26 MDSE00DJHO 2012-06-30 3001911139 2012-06-11 MDSE00DJHO 2012-06-30 300191139 2012-06-26 MDSE00DJHO		3001911138	2012-06-11	MDSE00DGZQ	3.00	47.20
2012-06-30 3001911138 2012-06-14 MDSE00DHYY 2012-06-30 3001911138 2012-06-18 MDSE00DICB 2012-06-30 3001911138 2012-06-21 MDSE00DJ02 2012-06-30 3001911138 2012-06-21 MDSE00DJK8 2012-06-30 3001911138 2012-06-25 MDSE00DJK8 2012-06-30 3001911138 2012-06-28 MDSE00DJK8 2012-06-30 3001911138 2012-06-28 MDSE00DJK8 2012-06-30 3001911138 2012-06-04 MDSE00DJK8 2012-06-30 3001911138 2012-06-04 MDSE00DGWA 2012-06-30 3001911138 2012-06-11 MDSE00DGWA 2012-06-30 3001911138 2012-06-18 MDSE00DJHG 2012-06-30 3001911138 2012-06-18 MDSE00DJHG 2012-06-30 3001911138 2012-06-18 MDSE00DJHG 2012-06-30 3001911139 2012-06-18 MDSE00DJHG 2012-06-30 300191139 2012-06-18 MDSE00DJHG		3001911138	2012-06-14	MDSE00DHYY	46.00	1,309.00
2012-06-30 3001911138 2012-06-18 MDSE00DICB 2012-06-30 3001911138 2012-06-21 MDSE00DJ02 2012-06-30 3001911138 2012-06-21 MDSE00DJ02 2012-06-30 3001911138 2012-06-25 MDSE00DJK8 2012-06-30 3001911138 2012-06-28 MDSE00DJK8 2012-06-30 3001911138 2012-06-28 MDSE00DJK8 2012-06-30 3001911138 2012-06-04 MDSE00DJK8 2012-06-30 3001911138 2012-06-11 MDSE00DJHG 2012-06-30 3001911138 2012-06-11 MDSE00DJHG 2012-06-30 3001911138 2012-06-11 MDSE00DJHG 2012-06-30 3001911138 2012-06-11 MDSE00DJHG 2012-06-30 3001911138 2012-06-18 MDSE00DJHG 2012-06-30 3001911138 2012-06-11 MDSE00DJHG 2012-06-30 3001911139 2012-06-12 MDSE00DJHO 2012-06-13 MDSE00DJHO 2012-06-14 MDSE00DJHO 2012-06-15 MDSE00DJHO 2012-06-15 MDSE00DJHO 2012-06-16 MDSE00DJHO 2012-06-17 MDSE00DJHO 2012-06-18 MDSE00DJHO 2012-06-19 MDSE00DJHO	•	3001911138	2012-06-14	MDSE00DHYY	3.00	75.00
2012-06-30 3001911138 2012-06-21 MDSE00DJ02 2012-06-30 3001911138 2012-06-21 MDSE00DJ02 2012-06-30 3001911138 2012-06-25 MDSE00DJK8 2012-06-30 3001911138 2012-06-28 MDSE00DJK8 2012-06-30 3001911138 2012-06-28 MDSE00DJK8 2012-06-30 3001911138 2012-06-04 MDSE00DJK8 2012-06-30 3001911138 2012-06-04 MDSE00DJHG 2012-06-30 3001911138 2012-06-11 MDSE00DJHG 2012-06-30 3001911138 2012-06-11 MDSE00DJHG 2012-06-30 3001911138 2012-06-18 MDSE00DJHG 2012-06-30 3001911138 2012-06-18 MDSE00DJHG 2012-06-30 3001911138 2012-06-11 MDSE00DJHG 2012-06-30 3001911138 2012-06-14 MDSE00DJHG 2012-06-30 3001911139 2012-06-14 MDSE00DJHG 2012-06-30 3001911139 2012-06-14 MDSE00DJHG 2012-06-30 3001911139 2012-06-18 MDSE00DJHG 2012-06-30 3001911139 2012-06-18 MDSE00DJHG 2012-06-30 3001911139 2012-06-18 MDSE00DJHG		3001911138	2012-06-18	MDSE00DICB	3.00	24.00
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Totals...

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Sample Container Detail-Details

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(a) (b) (c) (c) (c) (c) (c) (c) (c) (c) (c) (c		50280029188002IY 28 Gal Tub-Incinerate	502800291880020t Red 43 Gallon Hinged Lid	502800291880020(Red 43 Gallon Hinged Lid		Hinged	502800291880020/ Red 43 Gallon Hinged Lid		502800291880020t Red 43 Gallon Hinged Lid			502800291880020(Red 43 Gallon Hinged Lid			502800291880020(Red 43 Gallon Hinged Lid	502800291880020(Red 43 Gallon Hinged Lid	502800291880020(Red 43 Gallon Hinged Lid	502800291880020t Red 43 Gallon Hinged Lid	502800291880020t Red 43 Gallon Hinged Lid	502800291880020x Red 43 Gallon Hinged Lid	502800291880020(Red 43 Gallon Hinged Lid	502800291880020t Red 43 Gallon Hinged Lid	502800291880020/ Red 43 Gallon Hinged Lid	502800291880020t Red 43 Gallon Hinged Lid	5028002918800201 Red 43 Gallon Hinged Lid	502800291880020t Red 43 Gallon Hinged Lid	502800291880020/ Red 43 Gallon Hinged Lid		502800291880020t Red 43 Gallon Hinged Lid	50280029188002lY 28 Gal Tub-Incinerate	50280029188002lY 28 Gal Tub-Incinerate	502800291880020(Red 43 Gallon Hinged Lid	502800291880020t Red 43 Gallon Hinged Lid	502800291880020(Red 43 Gallon Hinged Lid	502800291880020(Red 43 Gallon Hinged Lid	Deguin.	502800291880020(Red 43 Gallon Hinged Lid	502800291880020(Red 43 Gallon Hinged Lid	502800291880020rRed 43 Gallon Hinged Lid	5028002918800201 Red 43 Gallon Hinged Lid	502800291880020(Red 43 Gallon Hinged Lid	502800291880020(Red 43 Gallon Hinged Lid	502800291880020(Red 43 Gallon Hinged Lid	502800291880020(Red 43 Gallon Hinged Lid	502800291880020(Red 43 Gallon Minged Lid	
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Sample Container Details

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Principal Control of the Control of	N 28 Gal Tub-In	SUCCOSTANTANTANTANTANTANTANTANTANTANTANTANTANT	SOCOOLS I COOLS I LED TENTE CONTRACTOR CONTR	002018800200211 Red 43 Gallon Hinged Lid	002918800200A2JI Red 43 Gallon Hinged Lid	002918800200A2JI Red 43 Gallon Hinged Lid	202918800200A2JI Red 43 Gallon Hinged Lid	M02918800200A2JIRed 43 Gallon Hinged Lid	002918800200A2JI Red 43 Gallon Hinged Lid	002918800200A2Jl Red 43 Gallon Hinged Lid	002918800200A2JI Red 43 Gallon Hinged Lid	002918800200A2JI Red 43 Gallon Hinged Lid	002918800200A2JI Red 43 Gallon Hinged Lid			002918800200A2JIRed 43 Gallon Hinged Lid	002918800200A2JI Red 43 Gallon Hinged Lid	002918800200A2Jl Red 43 Gallon Hinged Lid	002918800200A2JI Red 43 Gallon Hinged Lid	002918800200A2JI Red 43 Gailon Hinged Lid	002918800200AZJI Red 43 Gallon Hinger Lid	5028002918800204 Red 43 Gallon Hinged Lid	5028(0251860020) Red 43 Gallon milged Lid	50/250/0/29 100/0/20 Not 43 Callott Tilliged Etc.	50/26/0/29 100/0/2/ Ned 43 Callon Hinged Lid	SUZGOVEZ 10000ZCC 100, 43 Callon Hinded Lid	502002100002000000000000000000000000000	502800291880020(Red 43 Gallon Hinged Lid	502800291880020/ Red 43 Gallon Hinged Lid	502800291880020/ Red 43 Gallon Hinged Lid	502800291880020t Red 43 Gallon Hinged Lid	502800291880020(Red 43 Gallon Hinged Lid	502800291880020t Red 43 Gallon Hinged Lid	5028002918800201 Red 43 Gallon Hinged Lid	502800291880020/ Red 43 Gallon Hinged Lid	502800291880020(Red 43 Gallon Hinged Lid	502800291880020t Red 43 Gallon Hinged Lid	502800291880020/ Red 43 Gallon Hinged Lid	502800291880020(Red 43 Gallon Hinged Lid	502800291880020(Red 43 Gallon Hinged Lid				
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Sample Container Detail-Details

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Container Detail-Details Sample

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Sample Container Detail-Details

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Stericycle, Inc. - (800) 643-0240 - www.stericycle.com

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EXHIBIT H

Stericycle Exhibit No. ____ (MP-9)

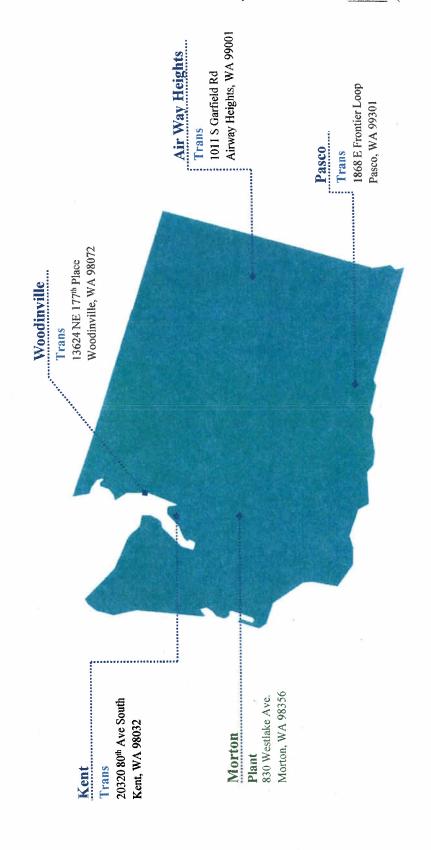
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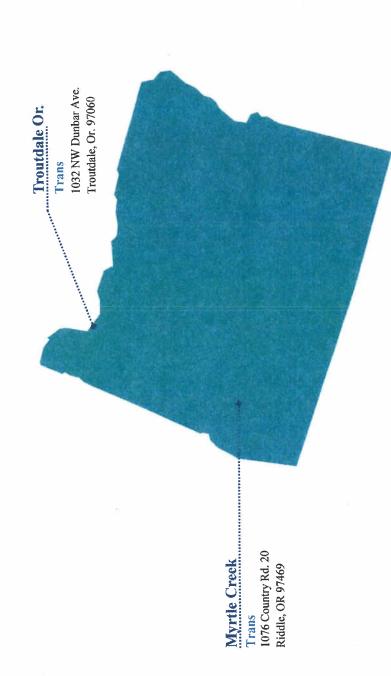
Stericycle of Washington, Inc. Spill Kit Inventory

The list below represents items that are present in a vehicle spill kit.

- Tyvek Suit / Boot covers
- Impervious Gloves (Latex or Nitrel)
- Disposable Surgical Mask
- Bleach or Other Approved Disinfectant
- Absorbent Materials (Snakes, Pads, Solidifier)
- Extra Red Bags
- Duct Tape & Caution Tape
- Safety Goggles / Glasses & Face Shield
- Tongs
- Germicidal Wipes
- Whisk Broom and Dust Pan
- Paper Towels







DEPARTMENT OF HEALTH & HUMAN SERVICES



Public Health Service

Food and Drug Administration 9200 Corporate Boulevard Rockville MD 20850

AUG 24 1995

Ms. Jane E. Rubinstein Director Environmental Affairs Biosystems 210 Sherwood Avenue Farmingdale, New York 11735

Re: K950897

Trade Name: Biobox Traptop (Small, Medium, Large

and X-Large)

Regulatory Class: II Product Code: FMI Dated: August 2, 1995 Received: August 4, 1995

Dear Ms. Rubinstein:

We have reviewed your Section 510(k) notification of intent to market the device referenced above and we have determined the device is substantially equivalent to devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act). You may, therefore, market the device, subject to the general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (Premarket Approval) it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 895. A substantially equivalent determination assumes compliance with the Good Manufacturing Practice for Medical Devices: General (GMP) regulation (21 CFR Part 820) and that, through periodic GMP inspections, FDA will verify such assumptions. Failure to comply with the GMP regulation may result in regulatory action. In addition, the Food and Drug Administration (FDA) may publish further announcements concerning your device in the Federal Register. Please note: this response to your premarket notification submission does not affect any obligation you might have under sections 531 through 542 of the Act for devices under the Electronic Product Radiation Control provisions, or other Federal Laws or Regulations.

Page 2 - Mr. Rubinstein

This letter immediately will allow you to begin marketing your device as described in your 510(k) premarket notification. An FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and permits your device to proceed to the market, but it does not mean that FDA approves your device. Therefore, you may not promote or in any way represent your device or its labeling as being approved by FDA. If you desire specific advice for your device on our labeling regulation (21 CFR Part 801 and additionally 809.10 for in vitro diagnostic devices), promotion, or advertising please contact the Office of Compliance, Promotion and Advertising Policy Staff (HFZ-300) at (301) 594-4639. Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers Assistance at their toll free number (800) 638-2041 or at (301) 443-6597.

Sincerely yours,

Timothy A. Ulatowski

Acting Director Pilot Division

Office of Device Evaluation

Center for Devices and Radiological Health

DEPARTMENT OF HEALTH & HUMAN SERVICES



Public Health Service

ALIG 2 4 1995

Food and Drug Administration 9200 Corporate Boulevard Rockville MD 20850

Ms. Jane E. Rubinstein Director Biosystems 210 Sherwood Avenue Farmingdale, New York 11735

Re: K950898

Trade Name: Biobox with Funnel Top (Small, Medium, Large, and X-Large)
Regulatory Class: II
Product Code: FMI
Dated: August 16, 1995
Received: August 16, 1995

Dear Ms. Rubinstein:

We have reviewed your Section 510(k) notification of intent to market the device referenced above and we have determined the device is substantially equivalent to devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (Premarket Approval) it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 895. A substantially equivalent determination assumes compliance with the Good Manufacturing Practice for Medical Devices: General (GMP) regulation (21 CFR Part 820) and that, through periodic GMP inspections, FDA will verify such assumptions. Failure to comply with the GMP regulation may result in regulatory action. In addition, the Food and Drug Administration (FDA) may publish further announcements concerning your device in the Federal Register. Please note: this response to your premarket notification submission does not affect any obligation you might have under sections 531 through 542 of the Act for devices under the Electronic Product Radiation Control provisions, or other Federal Laws or Regulations.

Page 2 - Mr. Rubinstein

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Sincerely yours,

A Timothy A. Ulatowski

Acting Director Pilot Division

Office of Device Evaluation Center for Devices and Radiological Health

EXHIBIT K

Stericycle Exhibit No. ____ (MP-12)

Sharps waste and container weight report (Jan-Aug 2012)

Service Line	i iotal Carts I	tal Reusable Containers	Total Net Waste Weight
BioSystems Sharps	11,322	464,763	1,319,342.9

Service Line	Reusable Container Size	Container Tare Weight (In Pounds)	Number of Containers	Total Container Tare Weight
BioSystems Sharps	2 Gal	1.9	138.843	263.801.7
BioSystems Sharps	3 Gal	2.0	146,557	293,114.0
BioSystems Sharps	4 Gal	3.4	96,560	330,235,2
BioSystems Sharps	8 Gal	6.0	43.380	260,280.0
BioSystems Sharps	17 Gal	7.0	39,423	275,961.0

Total Container Tare Weight: Total Net Waste Weight :	1,423,391.9 1,319,342.9
Total Gross	2,742,734.8
Recycle %	51.9%

EXHIBIT L

Stericycle Exhibit No. (MP-13)

BEFORE THE WASHINGTON STATE UTILITIES AND TRANSPORTATION COMMISSION

In Re Application of

WASTE MANAGEMENT OF WASHINGTON, INC. d/b/a WM Healthcare Solutions of Washington 720 4th Ave. Ste 400 Kirkland, WA 98033-8136 Docket No. TG-120033

PROTESTANT STERICYCLE OF WASHINGTON, INC.'S FIRST DATA REQUESTS TO APPLICANT WASTE MANAGEMENT OF WASHINGTON, INC. AND SUPPLEMENTAL RESPONSES THERETO

Subject to and without waving its previously stated objections, Waste Management of Washington, Inc. ("Waste Management") supplements its July 5, 2012 responses to Stericycle's First Data Requests as follows.

DATA REQUESTS

DATA REQUEST NO. 8:

Identify and Describe all offers and/or solicitations Relating to Your current or proposed Biomedical Waste Services that You have made to any existing or prospective customer from January 1, 2010 to the present.

Produce all Documents and Communications with actual or potential customers Relating to the offers and/or solicitations identified and described in response to this Data Request No. 8.

<u>Supplemental Response</u>: Marketing materials Waste Management has used in soliciting new business and advertizing its biomedical waste services are produced herewith. Jeff Norton and Jeff Daub have knowledge regarding this response.

DATA REQUEST NO. 9:

Identify and Describe all Documents and other materials you have used to advertise, promote, or otherwise make known Your current Biomedical Waste Services and all materials you intend to use for such purposes should Your Application be approved.

Produce copies of all such advertising and promotional Documents and materials.

<u>Supplemental Response</u>: Marketing materials Waste Management has used in soliciting new business and advertizing its biomedical waste services are produced herewith. Jeff Norton and Jeff Daub have knowledge regarding this response.

PROTESTANT STERICYCLE OF WASHINGTON, INC.'S FIRST DATA REQUESTS TO APPLICANT WASTE MANAGEMENT OF WASHINGTON, INC. AND SUPPLEMENTAL RESPONSES THERETO - 1

DATA REQUEST NO. 14:

Identify and Describe (a) each vehicle and any other transportation equipment You, Your Affiliates, or any independent contractor or other third party currently use in providing Biomedical Waste Services to Washington State customers and (b) any additional vehicles and other transportation equipment you intend to use in providing such Services if your Application is granted. Include in Your response a description of all features, design elements or modifications to such vehicles or equipment made for the purpose of preparing them for use to store or transport Biomedical Waste, DOT numbers, registration numbers, licensing information, signage, and vehicle markings, and state whether You own, lease, or rent the vehicle or other equipment and, if leased or rented, the name and address of the title holder.

<u>Supplemental Response</u>: A description of Waste Management's vehicles is produced herewith. Waste Management marks its vehicles in compliance with US DOT regulations. Jeff Norton and Jeff Daub have knowledge regarding this response.

DATA REQUEST NO. 18:

Describe any Services You offer involving the collection and transportation of sharps or sharps waste, including but not limited to any Service Relating to the BD ecoFinity Life Cycle Solution sharps program. Your answer must include, without limitation:

- (a) A statement characterizing Your Service either as involving the collection and transportation of recyclable materials unregulated by the WUTC or as involving the collection and transportation of Biomedical Waste regulated by the WUTC;
 - (b) A description of the material collected and transported;
- (c) A description of the sharps containers used, including all specifications, their manufacturers, and all manufacturer's information;
- (d) A description of all treatment and disposal methods employed for the material, including sharps containers;
- (e) If You contend that any portion of the sharps, sharps waste or sharps containers are recycled, a description of the material recycled, the methods used in such recycling, the percentage of the sharps or sharps containers that is recycled, and the methods used in tracking, calculating, and/or documenting the amounts recycled;
 - (f) The rates You charge for any such Services; and
- (g) The Washington State generators of Biomedical Waste to whom you provide any such Services.

Produce copies of all contracts, agreements, purchase orders, invoices, Communications, or other Documents describing or otherwise Related to any Services Identified in response to this Data Request No. 18.

Supplemental Response: Waste Management autoclaves the majority of sharps waste collected. Approved sharps containers are deposited into Waste Management's lined, reusable tubs and are transported to the Seattle processing plant for autoclaving. The sterilized sharps are then transported to Columbia Ridge or Greater Wenatchee Landfill for final disposal. BD ecoFinity is a sharps recycling program rolled out to hospitals in 2011 by Waste Management and Becton Dickenson. Waste Management collects full sharps containers weekly from St. Joseph Medical Center in Bellingham. The contract with St. Joseph Medical Center is produced herewith. The sharps containers are delivered to the Seattle processing facility and are loaded to 1-yard Gaylord's, placed on a 53' trailer and transported to Vernon, California for processing in a Red Bag Solutions machine. The sterilized, washed and shredded sharps containers and their contents are then sent to Talco Corporation where the material is separated utilizing float/sink technology. The plastics recovered in this process are pelletized and used in the remanufacturing of sharps containers. In May and June 2012, recycled sharps and sharps containers yielded between 17% and 28% of the recycled product. Waste Management accepts all approved sharps and sharps containers under both its BD ecoFinity program and its regulated biomedical waste program. Waste Management charges competitive market rates for its BD ecoFinity program and tariff rates for its regulated biomedical waste program. Jeff Daub, Jeff Norton and Tim Tucker have knowledge regarding this response.

DATA REQUEST NO. 20:

Describe each offer, solicitation, meeting, negotiation, or other Communication, and any agreement, contract, or other understanding reached or in effect, within the past 24 months Related to (1) Your Biomedical Waste Services, (2) Your Services Related to the collection, transportation or recycling of recyclable materials, and (3) Your rates or charges for any of such Services, with or involving any representative of each of the following:

- (a) Skagit Valley Hospital (Mt. Vernon)
- (b) Northwest Hospital (Seattle)
- (c) St. Joseph's Hospital (Bellingham)
- (d) Sacred Heart Hospital (Spokane)
- (e) Holy Family Hospital (Spokane)
- (f) Pathology Associates Medical Laboratories.

Produce all Communications, notes, reports, contracts, agreements, or other Documents Related to any offer, solicitation, meeting, negotiation, or other Communication, or any agreement, contract or understanding referenced in this Data Request No. 20.

<u>Supplemental Response</u>: The contracts with these entities along with the one email exchange which is responsive are produced herewith. Waste Management does not perform recycling services for Sacred Heart Medical Center, Holy Family Hospital, or Pathology Associates

PROTESTANT STERICYCLE OF WASHINGTON, INC.'S FIRST DATA REQUESTS TO APPLICANT WASTE MANAGEMENT OF WASHINGTON, INC. AND SUPPLEMENTAL RESPONSES THERETO - 6

SUPPLEMENTAL RESPONSES DATED this 27th day of July, 2012.

SUMMIT LAW GROUP PLLC

Polly L. McNeill, WSBA #17437

Jessica L. Goldman, WSBA #21856

pollym@summitlaw.com jessicag@summitlaw.com

Attorneys for Waste Management of Washington, Inc.

BEFORE THE WASHINGTON STATE UTILITIES AND TRANSPORTATION COMMISSION

In Re Application of

WASTE MANAGEMENT OF WASHINGTON, INC. d/b/a WM Healthcare Solutions of Washington 720 4th Ave. Ste 400 Kirkland, WA 98033-8136 Docket No. TG-120033

PROTESTANT STERICYCLE OF WASHINGTON, INC.'S FIRST DATA REQUESTS TO APPLICANT WASTE MANAGEMENT OF WASHINGTON, INC. <u>AND OBJECTIONS AND</u> RESPONSES THERETO

TO:

WASTE MANAGEMENT OF WASHINGTON, INC.

Pursuant to WAC 480-07-400 and 480-07-410, Protestant Stericycle of Washington, Inc. ("Stericycle") propounds the following data requests to Applicant Waste Management of Washington, Inc. ("Waste Management" or "Applicant").

INSTRUCTIONS AND DEFINITIONS

These data requests are continuing in nature, and if you obtain additional or different information after responding to them, you are required to file a supplemental response through the date of hearing. Each document requested in these data requests must be produced for inspection and copying at the offices of Garvey Schubert Barer, 1191 Second Avenue, 18th Floor, Seattle, Washington, or provided by some other mutually agreed method. Any electronic record requested in these data requests must be produced in a form and manner that is readable by conventional means and that preserves the record's metadata, including but not limited to title and subject, creation and modification dates, authors and editors, and sent and received dates. Any electronic records must be produced on a CD-ROM, DVD, or a portable hard drive.

If you object to answering any data request, in whole or in part, state your objections and state with particularity all of the factual and legal reasons supporting your objection in lieu of your answer. If you object on the ground of privilege, also state with particularity the nature and extent of all allegedly privileged matters and identify with specificity all allegedly privileged

PROTESTANT STERICYCLE OF WASHINGTON, INC.'S FIRST DATA REQUESTS TO APPLICANT WASTE MANAGEMENT OF WASHINGTON, INC. AND OBJECTIONS & RESPONSES THERETO.- 1

DATA REQUEST NO. 18:

Describe any Services You offer involving the collection and transportation of sharps or sharps waste, including but not limited to any Service Relating to the BD ecoFinity Life Cycle Solution sharps program. Your answer must include, without limitation:

Response: Waste Management objects that this Data Request is not reasonably calculated to lead to the discovery of admissible evidence, is overly broad and unduly burdensome and seeks Waste Management's trade secrets and confidential business information. Without waiving these objections, Waste Management generally describes below the recycling services it has offered since the filing of its biomedical waste tariff.

(a) A statement characterizing Your Service either as involving the collection and transportation of recyclable materials unregulated by the WUTC or as involving the collection and transportation of Biomedical Waste regulated by the WUTC:

<u>Response</u>: Collection services offer to BD ecoFinity customers are performed in the same manner as medical waste customers with the exception of the uniquely labeled tubs filled with sharps containers. Once these tubs are received at the Seattle processing plant, the tubs are loaded onto trailers and transported to Vernon, California for processing. Waste Management performs this as a commercial recycling collection service.

(b) A description of the material collected and transported;

Response: Tubs filled with sharps and sharps containers.

(c) A description of the sharps containers used, including all specifications, their manufacturers, and all manufacturer's information:

Response: Becton, Dickinson and Company sharps containers of various sizes.

(d) A description of all treatment and disposal methods employed for the material, including sharps containers;

Response: Sharps and sharps containers are either disposed of at a landfill or processed for recycling.

(e) If You contend that any portion of the sharps, sharps waste or sharps containers are recycled, a description of the material recycled, the methods used in such recycling, the percentage of the sharps or sharps containers that is recycled, and the methods used in tracking, calculating, and/or documenting the amounts recycled;

Response: Tubs are transported to Waste Management's facility at Vernon, California, and processed in a Red Bag Solutions (RBS) hardware/software system designed to safely,

PROTESTANT STERICYCLE OF WASHINGTON, INC.'S FIRST DATA REQUESTS TO APPLICANT WASTE MANAGEMENT OF WASHINGTON, INC. AND OBJECTIONS & RESPONSES THERETO.- 18

efficiently, and effectively sterilize and grind medical waste. By exposing infectious medical waste to superheated water and steam (272°F / 133°C) and simultaneously employing a proprietary cutting system, the RBS renders infectious medical waste non-infectious, non-hazardous, and non-recognizable. Once processed through the RBS, the non-infectious medical waste is sent to Talco Plastics in Corona, California where the non-infectious ground sharps are processed and the metals and plastics separated. The recovered plastics are pelletized at Talco and sent to BD to be manufactured into BD Recykleen Products.

(f) The rates You charge for any such Services; and

Response: Waste Management objects to this Data Request as being prohibited by Order 01 ¶ 8, Order 03 ¶ 24, and Order 04 ¶ 10. Waste Management further objects that this Data Request is not reasonably calculated to lead to the discovery of admissible evidence, is overly broad, unduly burdensome, and seeks Waste Management's trade secrets and confidential business information.

(g) The Washington State generators of Biomedical Waste to whom you provide any such Services.

Response: Waste Management objects to this Data Request as being prohibited by Order 01 ¶ 8, Order 03 ¶ 24, and Order 04 ¶ 10. Waste Management further objects that this Data Request is not reasonably calculated to lead to the discovery of admissible evidence, is overly broad, unduly burdensome, and seeks Waste Management's trade secrets and confidential business information.

Produce copies of all contracts, agreements, purchase orders, invoices, Communications, or other Documents describing or otherwise Related to any Services Identified in response to this Data Request No. 18.

<u>Response</u>: Subject to, and without waiving, the above-stated objections, Waste Management will produce documents at a mutually agreeable time.

DATA REQUEST No. 19:

Describe Your existing Biomedical Waste operations in Washington, including collection, transportation, treatment, and disposal, and provide an itemized description of all personnel, vehicles, transportation equipment, transportation yards, transportation routes, storage facilities, transfer facilities, treatment facilities, disposal facilities and other facilities or equipment currently used in providing such Services, and Describe all changes to existing operations, including personnel, vehicles, transportation equipment, transportation yards, transportation routes, storage facilities, transfer facilities, treatment facilities, disposal facilities and other facilities or equipment You intend to implement to provide Biomedical Waste Services in the additional territory covered by the Application.

Response: Waste Management objects to this Data Request as being prohibited by Order 01 ¶ 8, Order 03 ¶ 24, and Order 04 ¶ 10. Waste Management further objects that this Data Request is

PROTESTANT STERICYCLE OF WASHINGTON, INC.'S FIRST DATA REQUESTS TO APPLICANT WASTE MANAGEMENT OF WASHINGTON, INC. AND OBJECTIONS & RESPONSES THERETO. 19

By

Stephen B. Johnson, WSBA #6196 Jared Van Kirk, WSBA #37029 Attorneys for Protestant Stericycle of Washington, Inc.

1191 Second Avenue, 18th Floor Seattle, WA 98101 206-464-3939

Direct: 206-816-1385 Fax: 206-464-0125

Email: sjohnson@gsblaw.com jvankirk@gsblaw.com

RESPONSES AND OBJECTIONS DATED this 5th day of July, 2012.

SUMMIT LAW GROUP PLLC

y ___[_

Polly L. McNell, WSBA #17437 Jessica L. Goldman, WSBA #21856 pollym@summitlaw.com jessicag@summitlaw.com

Attorneys for Waste Management of Washington, Inc.

StARS.Reports 1.15.1223.0

Bio Systems Container Volume Report Sample

Bio Systems Container Volume Report 06/27/2007 Report Release #: Report BulldDate: Sample

Run Date: 2012-07-16

Report Parameters:

2012-06-30 BeginDate: 2012-06-01 EndDate:

6082432 001 Customer

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Click on the tabs at the bottom of the workbook to view report.

Page 1 of 3

Sample BioSystems Container Volume Report

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