

**MAHARASHTRA POLLUTION CONTROL BOARD
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LETTER OF BIO-MEDICAL WASTE AUTHORISATION

[Authorisation for Generation, Storage, Disposal of Bio-Medical Wastes under Rule 7(4)]

I. File number of authorisation and date of issue

RO-AURANGABAD/ABD/BMW - AUTH/1804000037 Date:- **3 / 03 /2018**

II. **M/s. Ashtavinayak Hospital and Research Center**, is hereby granted an authorisation for generation of biomedical waste on the premises situated **At Plot No. X-69, Maharana Pratap Chowk, Bajaj Nagar, MIDC Waluj, Aurangabad.**

III. This authorisation shall be in force for a period up to **30.06.2020.**

An application shall be made by the occupier/operator for renewal **3 Months** before expiry of earlier authorisation.

IV. This authorisation is issued subject to compliance of the conditions stated below and to such other conditions as may be specified in the Rules for the time being in force under the Environment (Protection) Act, 1986.

V. No of Beds: **30.**

Terms and Conditions of authorisation

1. The authorized Person shall comply with the provisions of the Environment (Protection) Act, 1986, and the Rules made there under.
2. The authorisation shall be produced for inspection at the request of an officer authorized by the prescribed authority.
3. i) The authorized person shall not rent, lend or sell the biomedical waste or facility.
ii) The authorized person can transfer the BMW generated at above premises to the "Transporter" or "Operator of Facility" authorized by MPCB under Bio-Medical Waste (Management and Handling) Rules, 1998 for collection, transportation, treatment and/or disposal of BMW generated.
4. Any unauthorized change in equipment or working conditions as mentioned in the application by the person authorized shall constitute a breach of this authorisation.
5. It is the duty of the authorized person to take prior permission of the prescribed authority to close down the facility.

6. The authorisation is granted for generation of Bio-Medical Waste (BMW) in waste categories and quantities listed here in below:

Sr. No	Category No.	Description	Quantity (kg/m)	Treatment & disposal
1	1	Human Anatomical Waste	4.0	Bio medical Waste shall be sent to Common BMW Treatment & Disposal facility authorized by MPCB
2	3	Microbiology & Biotechnology Waste	6.0	
3	4	Waste sharps	5.0	
4	6	Solid waste	6.0	
5	7	Solid Waste (item contaminated with blood)	5.0	
6	---	Mercury Waste	---	---
7	---	X-ray hypo solution (fixture)	---	---
8	---	X-ray films & non-infected plastic recycle waste	---	---

* The occupier shall join the common BMW Treatment & Disposal Facility authorized by MPCB ne i.e. M/s – Water Grace Products, Surve No.122, Gevrai tanda, Patoda shivar, Paithan Road, Aurangabad and shall send the BMW regularly to CBMWSTDF for its scientific treatment & disposal.

7. The hospital authority shall proper collection of mercury spillage arising due to breakage of thermometer, pressure gauges & other equipments used in health care facilities as well as the storage shall be in accordance with Hazardous Waste (Management, Handling & Transboundry Movement) Rules, 2008 and returning it to instrument manufacturing apart from taking necessary steps to ensure that the spilled mercury does not become a part of Bio-Medical Waste or other solid waste generated from Health Care Facilities.
8. The liquid/solid waste generated from the treatment activity (from laboratory and washing, cleaning, housekeeping and disinfecting activities) shall be treated suitably by providing effluent treatment facility to conform the standards prescribed in Schedule V of said Rules and the Environment (Protection) Act, 1986.
9. (i) BMW shall be treated and disposed of in accordance with Schedule I; and in compliance with the standards prescribed in Schedule V of said Rules.
10. (i) BMW shall not be mixed with other wastes or reused, recycled or sold in any form.
- (ii) BMW shall be segregated into containers / bags at the point of generation in accordance with Schedule-II prior to storage, treatment and disposal. The containers shall be labeled according to Schedule III.
- (iii) If a container containing BMW is to be transported from the premises where BMW is generated to any waste treatment facility outside the premises, the container shall, apart from the Label prescribed in Schedule III, also carry information prescribed in Schedule IV and shall be transported by authorized Transporter only.
- (iv) Notwithstanding anything contained in the Motor Vehicles Act, 1988 or Rules there under, BMW shall be transported only in such vehicle as may be authorized for the purpose by the competent authority as specified by the Government.
- (v) No untreated BMW shall be kept stored beyond a period of 48 hours.

11. Standards for waste autoclaving:

The autoclave should be dedicated for the purposes of disinfecting and treating Bio-Medical Waste,

- (I) When operating a gravity flow autoclave, medical waste shall be subjected to:
 - (i) A temperature of not less than 121 C° and pressure of 15 pounds per Square inch (psi) for an autoclave residence time of not less than 60 minutes;
or
 - (ii) A temperature of not less than 135 C° and a pressure of 31 psi for an autoclave residence time of not less than 45 minutes;
or
 - (iii) A temperature of not less than 149 C° and a pressure of 52 psi for an autoclave residence time of not less than 30 minutes.
 - (II) When operating a vacuum autoclave, medical waste shall be subjected to a minimum of one pre-vacuum pulse to purge the autoclave of all air. The waste shall be subjected to the following.
 - (i) a temperature of not less than 121 C° and a pressure of 15 psi for an autoclave residence time of not less than 45 minutes;
or
 - (ii) a temperature of not less than 135 C° and a pressure of 31 psi for an autoclave residence time of not less than 30 minutes; or
 - (III) Medical waste shall not be considered properly treated unless the time, temperature and pressure indicators indicate that the required time, temperature and pressure were reached during the autoclave process. If for any reasons, time temperature or pressure indicates that the required temperature, pressure or residence time was not reached, the entire load of medical waste must be autoclaved again until the proper temperature, pressure and residence time were achieved.
 - (IV) *Recording of operational parameters*,- Each autoclave shall have graphic or computer recording devices which will automatically and continuously monitor and record dates, time of day, load identification number and operating parameters throughout the entire length of the autoclave cycle.
 - (V) *Validation test: Spore testing*. – The autoclave should completely and consistently kill the approved biological indicator at the maximum design capacity of each autoclave unit. Biological indicator for autoclave shall be *Bacillus stearothermophilus* spores using vials or spore strips, with at least 1×10^4 spores per milliliter. Under no circumstances will an autoclave have minimum operating parameters less than a residence time of 30 minutes, regardless of temperature and pressure, a temperature less than 121 C° or a pressure, less than 15 psi.
 - (VI) *Routine Test*. — a chemical indicator strip/tape that changes color when a certain temperature is reached can be used to verify that a specific temperature has been achieved. It may be necessary to use more that one strip over the waste package at different location to ensure that the inner content of the package has been adequately autoclaved.
12. Every 'Authorized Person' shall submit an Annual Report to the prescribed authority in Form-II by 31st January every year including information about the categories and quantities of BMW handled during the preceding year.