

MAHARASHTRA POLLUTION CONTROL BOARD

Phone : 0240-2993004



Sub-Regional Office:

Paryavaran Bhavan, Plot No.A-4/1,MIDC
Chikalhana, Behind Dainik Lokpatra, Near Seth
Nandlal Dhoot Hospital Jalna Road,Aurangabad-
431210

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"Your Service is our Duty"

LETTER OF BIO-MEDICAL WASTE AUTHORISATION [Authorisation for Generation, Collection, Reception, Segregation, Storage of Bio-Medical Wastes under Rule 7(4)]

I. File number of authorisation and date of issue

SRO-AURANGABAD/BMW-AUTH/2205000764 Date: 30/05/2022

II. **M/s. Gurukrupa Hospital** is hereby granted an authorization for generation of biomedical waste on the premises situated **At Devgaon Rangari, Tal. Kannad, Dist:- Aurangabad.**

III. This authorization shall be in force for a period up to **30/04/2025**. An application shall be made by the occupier/operator for renewal **3 Months** before expiry of earlier authorization.

IV. This authorization 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: 05 Beds.**

Terms and Conditions of authorized on

1. The authorized Person "**Dr. Sandip Asaram Kadam**, shall comply with the provisions of the Environment (Protection) Act, 1986, and the Rules made thereunder.
2. The authorization 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, 2016 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 authorization.
5. It is the duty of the authorized person to take prior permission of the prescribed authority to close down the facility.

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

Sr. No	Category	Type of Waste	Quantity not to exceed (Kg/M)	Type of Bag or Container to be used	Treatment and Disposal options
1	Yellow	(a) Human Anatomical Waste: Experimental animal carcasses, body parts, organs, tissues, including the waste generated from animals used in experiments or testing in veterinary hospitals or colleges or animal houses.	--	Yellow colored non-chlorinated plastic bags	CBMWSTDF, Aurangabad
		(c) Soiled Waste: Items contaminated with blood, body fluids like dressings, plaster casts, cotton swabs and bags containing residual or discarded blood and blood components.	2.0	Yellow coloured non-chlorinated plastic bags	CBMWSTDF, Aurangabad
		(d) Expired or Discarded Medicines	--	Yellow coloured non-chlorinated plastic bags	CBMWSTDF, Aurangabad
		e) Chemical Waste	..		
		f) Chemical Liquid Waste	--	Separate collection system leading to effluent treatment system	
		h) Microbiology Biotechnology and other clinical laboratory waste	--	Autoclave safe plastic bags or containers	
2	White	Waste sharps including Metals	2.0	Puncture proof, Leak proof, tamper proof container	CBMWSTDF, Aurangabad
3	Red	Contaminated Waste (Recyclable)	2.0	Red Coloured non chlorinated plastic bags or containers	CBMWSTDF, Aurangabad
No onsite treatment of BMW is permitted. The above mentioned Bio medical Waste shall be sent to Common BMW Treatment & Disposal facility authorized by MPCB					

6. 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.
8. (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.
- (ii) You shall setup requisite BMW treatment facilities like incinerator, autoclave / Microwave, shredder etc., at the disposal side in accordance with the BMW rules. You shall disposed of the duly treated BMW and incineration ash in secured land fill site at your own premises / at MSW secured land fill site of Municipal Council authorized by MPCB and duly earmarked for disposal of treated BMW /at common H.W. treatment & disposal facility setup as per the Hazardous Waste (M & H) Rules, 2016 as amended and authorized by MPCB.
9. (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.

10. 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 than one strip over the waste package at different location to ensure that the inner content of the package has been adequately autoclaved.
11. Every 'Authorized Person' shall submit an Annual Report to the prescribed authority in Form-II by 30th June every year including information about the categories and quantities of BMW handled during the preceding year.
12. (i) Every 'Authorized Person' shall maintain records related to the generation, collection, reception, storage, transportation, treatment, disposal and/or any form of handling of BMW in accordance with these Rules and any guidelines issued.
- (ii) All records shall be subject to inspection and verification by the prescribed authority at any time.
13. When any accident occurs at any institution or facility or any other site where BMW is handled or during transportation of such waste, the authorized person shall report the accident in Form III to the prescribed authority forthwith.
14. Authorization is granted subject to obtain valid CBMWTSDF membership and registration under Bombay Nursing Home (Registration) Act 2005 Under Section (5) from Aurangabad Municipal Corporation Aurangabad.
15. Board reserved right to revoke, suspend or amend the BMW Authorization issued.

16. This Authorization is issued subject to conditions as mentioned below:
- Hospital Authority shall submit compliance of authorization conditions every six months to Sub- Regional Office Aurangabad.
 - Hospital Authority shall Provide Annual Report in form -II duly certified by the facility operator.
 - Hospital Authority shall provide valid copy registration under Bombay Nursing Home act 1949.
 - The Occupier will obey all the lawful instruction issued by the Board officer from time to time.
 - The Hospital shall not carry out any expansion, change, addition, modification and modernization in the existing set up till to prior permission from the Board.
 - This Authorization is issued subject to obtain/submission of valid membership of CBMWTSDF, Aurangabad and Certificate of Registration Under section (5) of Bombay Nursing Homes Registration Act & amendment 1949.
17. Hospital Authority shall comply the uniform guidelines issued vide circular No. MPCB/PSO/BMW/B-1825 dtd. 10/04/2013. (Copy available on Board website i.e www.mpcb.gov.in).
18. Hospital Authority shall operate the activity as per nos. of Beds mentioned in Bombay Nursing Home registration certificate issued by concerned Authority (i. e AMC/Health Dept., etc.)
19. The Hospital shall obtain/renew the Bombay Nursing Home Registration Certificate within three months period & submit/upload to MPCB Office, Failing which this authorization shall be treated as null & void (Authorization shall be revoked/cancelled).
20. The BMW authorization is granted subject to comply the BMW Rule & Amended thereof & .to upload BMW Data/Information in COVID APP as per the direction of CPCB, if applicable.
21. The hospital authority shall submit the valid Bombay Nursing Home Act, certificate from the concerned department within a period of one month.

For and on behalf of the
Maharashtra Pollution Control Board

S. S. Rathod
30/08/2022
(S. S. Rathod)

Sub-Regional Officer, Aurangabad.

To,
M/s. Gurukrupa Hospital,
At Devgaon Rangari,
Tal. Kannad, Dist:- Aurangabad.
Authorization Fees Received:-



Sr. No.	Amount (Rs.)	Transaction Number	Trans. Date	Trans. Type	Bank Name
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