

## MAHARASHTRA POLLUTION CONTROL BOARD

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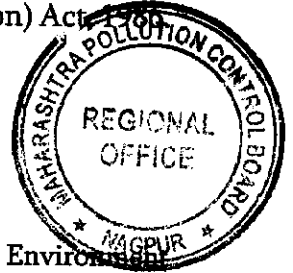
### LETTER OF BIO-MEDICAL WASTE AUTHORISATION

[Authorization for Generation, Storage of Bio-Medical Wastes under Rule 10]

- I. File number of authorization and date of issue  
RO-Nagpur /BMW\_AUTH/2010000276 Dt. 27/11/2020
- II. Your Online BMW Authorization Application through Web-Portal having  
UAN: MPCB-BMW\_AUTH-0000028811 on Dt. 13/02/2020
- III. M/s. Ordnance Factory Hospital Bhandara is hereby granted an authorization  
for generation of biomedical waste on the premises situated A/p-  
Jwaharnagar, Bhandara 441904.
- IV. This authorization shall be in force for a period up to 31/01/2025. An application  
shall be made by the occupier/operator for renewal 3 Months before expiry of  
earlier authorization.
- V. 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.
- VI. No of Beds: 40

#### Terms and Conditions of authorization

1. The authorized Person shall comply with the provisions of the Environment  
(Protection) Act, 1986, and the Rules made there under.
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 Rules, 2016 & amendment thereof 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.

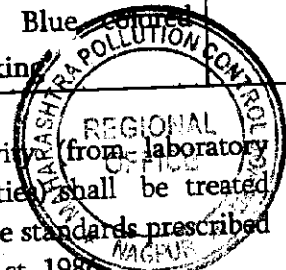
**Schedule-III**

**Treatment and Disposal of Biomedical Waste generated from Hospital to CBMWTSDF**

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

Sr. No.	Category	Type of Waste	Quantity not to exceed (Kg/M)	Segregation coding	Colour	Treatment & Disposal
1	Yellow	a) Human Anatomical waste	5	Yellow non-plastic bags	coloured chlorinated	No onsite treatment of BMW is permitted. The above mentioned Bio medical Waste shall be sent to Common BMW Treatment & Disposal facility authorised by MPCB i.e. CBMWTSDF
		b) Animal Anatomical Waste	NIL			
		c) Soiled Waste	15			
		d) Expired or Discarded Medicines	0.5			
		e) Chemical Waste	Nil	Separate collection system leading to effluent treatment system		
		f) Chemical Liquid Waste	15			
		g) Discarded linens, mattresses, beddings contaminated with blood or body fluid.	Nil		Yellow coloured non-chlorinated plastic bags or suitable packing material	
		h) Microbiology Biotechnology and other clinical laboratory waste	NIL	Autoclave safe plastic bags or containers		
2		Contaminated waste (Recyclable)	7	Red coloured non-chlorinated plastic bags or containers		
3	White (Translucent)	Waste sharps including Metals	9	Puncture proof, Leak proof, tamper proof container		
4	Blue	a) Glassware	NIL	Cardboard boxes with Blue colour marking		

6. The liquid/solid waste generated from the treatment activities 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.



7. (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, 1989 as amended and authorized by MPCB.

8. (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.

09. Standards for waste autoclaving:

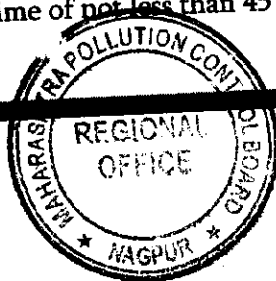
The autoclave should be dedicated for the purposes of disinfecting and treating bio-medical waste,

(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  $1 \times 10^4$  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.

#### 10. CONDITIONS UNDER AIR ACT :

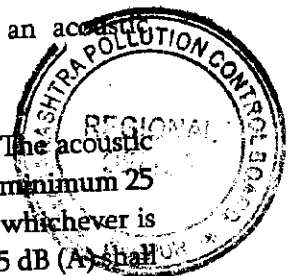
- (i) The applicant shall install a comprehensive control system consisting of control equipments as is warranted with reference to generation of emission and operate and maintain the same continuously so as to achieve the level of pollutants to the following standards:

##### Control Equipment:

Industry shall provide dust collector of sufficient capacity to control the emissions.

##### Conditions for D.G. Set

- a. Noise from the D.G. Set should be controlled by providing an acoustic enclosure or by treating the room acoustically.
- b. Industry should provide acoustic enclosure for control of noise. The acoustic enclosure/ acoustic treatment of the room should be designed for minimum 25 dB (A) insertion loss or for meeting the ambient noise standards, whichever is on higher side. A suitable exhaust muffler with insertion loss of 25 dB (A) shall also be provided. The measurement of insertion loss will be done at different points at 0.5 meters from acoustic enclosure/room and then average.



- c. The industry shall take adequate measures for control of noise levels from its own sources within the premises in respect of noise to less than 55 dB(A) during day time and 45 dB(A) during the night time. Day time is reckoned between 6 a.m. to 10 p.m and night time is reckoned between 10 p.m to 6 a.m.
- d. Industry should make efforts to bring down noise level due to DG set, outside industrial premises, within ambient noise requirements by proper siting and control measures.
- e. Installation of DG Set must be strictly in compliance with recommendations of DG Set manufacturer.
- f. A proper routine and preventive maintenance procedure for DG set should be set and followed in consultation with the DG manufacturer which would help to prevent noise levels of DG set from deteriorating with use.
- g. D.G. Set shall be operated only in case of power failure.
- h. The applicant should not cause any nuisance in the surrounding area due to operation of D.G. Set.

**11. Standards for Stack Emissions:**

- (i) The applicant shall observe the following fuel pattern:-

Sr. No.	Type Of Fuel	Quantity	UOM
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The applicant shall erect the chimney(s) of the following specifications:-

Sr. No.	Chimney Attached To	Height in Mtrs.
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(ii) The applicant shall provide ports in the chimney/(s) and facilities such as ladder, platform etc. for monitoring the air emissions and the same shall be open for inspection to/and for use of the Board's Staff. The chimney(s) vents attached to various sources of emission shall be designated by numbers such as S-1, S-2, etc. and these shall be painted/ displayed to facilitate identification.

- (iii) The industry shall take adequate measures for control of noise levels from its own sources within the premises so as to maintain ambient air quality standard in respect of noise to less than 75 dB(A) during day time and 70 dB(A) during night time. Day time is reckoned in between 6 a.m. and 10 p.m. and night time is reckoned between 10 p.m. and 6 a.m.

*Signature*



(vi) Other Conditions:

- 1) The industry should not cause any nuisance in surrounding area.
- 2) The industry should monitor stack emissions and ambient air quality Regularly.

12. Every 'Authorized Person' shall submit an Annual Report to the prescribed authority in Form-II by 31<sup>st</sup> January every year including information about the categories and quantities of BMW handled during the preceding year.

13. (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.

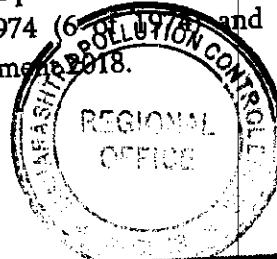
14. You have submitted total Bank Guarantee of Rs. 1,50,000/- valid up to 21/07/2025 Specific conditions with Bank Guarantees along with time bound programme for compliance as per Board's circular no MPCB/PSO/BMW/B-1825 dtd 10/04/2012 is as follows-

Sr. No.	Activity/Condition to be Complied	Compliance Timelime (Months)	Amount of Bank Guarantee (Rs)
<b>I(A) Operation and Maintenance</b>			
1	To segregate and Handle BMW as per rule	Continuous	25,000
2	Operation and Maintenance of ETP to achieve prescribed discharged standards	Continuous	25,000
<b>I(B) Records</b>			
1	To Maintain records of BMW and submission of Annual Report in Form-II before 31st January	Continuous	15,000
2	To Maintain records of BMW material received / delivered to authorized party / CBMWTSDF (Transporters only)	Continuous	10,000
<b>II Performance</b>			
1	To provide Separate BMW storage facility	Six	25,000
2	Effluent treatment plant not provided / need upgradation	Six	50,000
Total			1,50,000/-

Rupees One Lakh Fifty Thousand only

15. You shall submit compliance of each Bank Guarantee conditions every six months to Regional Officer, Nagpur for verification purpose.

16. Ensure treatment and disposal of liquid waste in accordance with Water (Prevention and Control of Pollution) Act, 1974 (6) and comply Bio-Medical Waste Management Rules 2016 & amendments 2018.



16. 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.
17. The Occupier will obey all the lawful instructions issued by the Board Officers from time to time.
18. The hospital shall apply immediately for GCA with required/compulsory document with consent fees.

For and on behalf of the  
Maharashtra Pollution Control Board

(A.M. Kare)  
Regional Officer, Nagpur



To  
M/s. Ordnance Factory Hospital Bhandara,  
Jwaharnagar, Bhandara 441904.

Authorization Fees Received:-

Sr. No.	Amount (Rs.)	e-Payment	Date	Drawn On
1	12500/-	DR No 5459838	13-02-2020	SBI

Copy Submitted to:-

1. The Chief Accounts Officer, MPCB Board Mumbai
2. The Principal Scientific Officer, MPCB, Sion, Mumbai

Copy to:-

Sub-Regional Officer, M.P.C. Board, Bhandara.

For information necessary compliance of condition mentioned in the authorization. He / She is also directed to ensure the timely compliance of condition no. 14.