कार्यालय आदेश

पर्यावरण मंडल की 36वीं सैंटो दिनांक 21.12.2018 के क्रम संख्या 05 (अध्यक्ष के आदेश से) में लिये गये निर्णय के अनुसार 05 कलोद तक के लागत घूमनी के लाल, नागनी एवं हरी श्रेणी की इकाईयाँ का सी.टी.ओ इस आदेश के निर्विरोध तिथि से आरम्भ होगी। अन्य सर्व निर्देश होगी:-

1. Auto renewal के लिए इकाई अपने current CTO के समापति अवधि के स्था पत्तम 69 दिन पहले आवेदन को Upload करना होगा।

2. Auto Renewal की अवधि current CTO के समापति अवधि के अगले दिन से प्रभावी होगा।

3. सी.टी.ओ के लिए निर्धारित शुल्क जमा करने पर ही Auto renewal का आवेदन पत्र स्वीकार होगा।

4. सी.टी.ओ के लिए निर्धारित सभी अभिलेख जमा करना होगा।

5. इकाई के प्रोप्राइटर अथवा अधिकृत प्रतिनिधि द्वारा एक Self Declaration देना होगा जिसमें अंकित तिथि के अंतर्गत पत्र होने पर इसकी कानूनी रूप से जाहिराती हुक्की होगी।

6. पर्यावरण CTO आवेदन के निर्णय के लिए निर्धारित समय सीमा के अंतर्गत आवेदन का निर्णय करना होगा।

7. Auto renewal की राहत एक वर्ष की होगी तथा स्थायी सीटीओ निर्माता/ रद्द होने पर Auto renewal की नीति स्वतंत्र संपादन हो जाएगी।

इसमें सक्षम प्राधिकार का आदेश निहित है।

जापाम: ...D,C.Y.

प्रतिलिपि- सभी संबंधित को सूचनार्थ एवं आवश्यक कारावाद हेतु प्रेषित।

(राजीव लौचन बक्सी) सदस्य सचिव

(राजीव लौचन बक्सी) सदस्य सचिव
Ranchi, Dated 11.12.15

Notification

The validity of the Consent to operate was notified as 05 (five) years, 10 (ten) years and 15 (fifteen) years for the industries falling under Red Category, Orange Category and Green Category, respectively vide Notification No.-B-06 dated 15.06.2015 of the Jharkhand State Pollution Control Board. The frequency of the inspection of the industries to be carried out was also notified vide aforesaid Notification.

In continuation to the said Notification, it has been decided that the same Inspector of the Jharkhand State Pollution Control Board will not inspect an industry twice consecutively.

Sd/-
(Sanjay Kr. Suman)
Member secretary

Memo No. 5327
Ranchi, Dated 11.12.15

Copy to: Principal Secretary, Forest, Environment and Climate Change, Govt. of Jharkhand/ Principal Secretary, Industry Deptt., Govt. of Jharkhand/ Principal Secretary, Mines and Geology, Deptt., Govt.of Jharkhand/ Director, Industries, Govt. of Jharkhand for information and necessary action.

Sd/-
(Sanjay Kr. Suman)
Member secretary

Memo No. 5327
Ranchi, Dated 11.12.15

Copy to: Section Head, Head Office/All RO's JSPC Board for information and necessary action.

(Sanjay Kr. Suman)
Member secretary
Ranchi, Dated - 11/12/15

Notification

In partial modification to Notification No.-05 dated 25.02.2012, the guidelines for validity of NOC (Consent to Establish) are revised to the extent as follows:

<table>
<thead>
<tr>
<th>S.N.</th>
<th>Item</th>
<th>As notified earlier</th>
<th>Validity period</th>
<th>Revised</th>
</tr>
</thead>
<tbody>
<tr>
<td>1.</td>
<td>New Project</td>
<td>01 year or till validity period of Environment Clearance</td>
<td>05 years for the project which requires environment and forest clearance</td>
<td></td>
</tr>
<tr>
<td>2.</td>
<td>Expansion of existing unit</td>
<td>01 year or till validity period of Environment Clearance</td>
<td>05 years for the project which require environment and forest clearance</td>
<td></td>
</tr>
</tbody>
</table>

Sd/-
(Sanjay Kr. Suman)
Member secretary

Ranchi, Dated - 11/12/15

Memo No.-B - 5328

Copy to: Principal Secretary, Forest, Environment and Climate Change, Govt. of Jharkhand / Principal Secretary, Industry Deptt., Govt. of Jharkhand/ Principal Secretary, Mines and Geology, Deptt., Govt. of Jharkhand/Director, Industries, Govt. of Jharkhand for information and necessary action.

Sd/-
(Sanjay Kr. Suman)
Member secretary

Ranchi, Dated - 11/12/15

Memo No.-B - 5328

Copy to: Section Head, Head Office/All RO’s JSPC Board for information and necessary action.

(Sanjay Kr. Suman)
Member secretary
NOTIFICATION

In order to ensure the quick disposal of applications for obtaining CTE, CTO under Water (Prevention & Control of Pollution) Act, 1974 and Air (Prevention & Control of Pollution) Act, 1981, authorization under Bio-medical Waste Management Rules, 2016 and Hazardous Waste and Trans-boundary Movement Rules, 2016, Board has decided to publish list of documents to be uploaded online along with the applications through "www.advantage.jharkhand.gov.in" or "www.jspecb.nic.in". (List is enclosed).

This order shall be effective from the immediate effect and will supersede all the previous orders.

(Rajeev Lochan Bakshi)
Member Secretary

Memo No.- B - 1195

Copy to Sr Anil Kumar, Sr. Technical Director, NIC, Delhi/ all section heads, Head office, JSPCB/ concerned officials, Head office, JSPCB/All Regional Officers, JSPCB/ for information and necessary action

(Rajeev Lochan Bakshi)
Member Secretary
Check-list of documents for submission of applications for authorization under

Documents to be upload:

1. Authorization to file application.
2. Duly filled up application form.
3. Copy of valid Consent to Operate (CTO) [for bedded HCF].
4. Copy of annual return.
5. Upload the names of committee members for reviewing and monitoring the activities related to bio-medical waste management and also upload minutes of committee meeting (for the HCFs beds more than 29).
6. Upload the names of designated qualified person to review and monitor the activities relating to bio-medical waste management (for the HCFs beds up to 29).
7. Upload name of website on which Authorization, Annual report has been uploaded.
8. Photograph of the bio-medical waste management register to be maintained and updated on day to day basis.
9. Photograph of established Bar- Code System for bags or containers containing bio-medical waste to be sent out of the premises.
10. Photograph of the area for a safe, ventilated and secured location for storage of segregated biomedical waste in colored bags or containers in the manner as specified.
11. Upload details of global positioning system for transportation of bio- medical waste (for CBWTF).
12. List of the occupiers which are not handing over the segregated bio-medical waste in accordance with these rules (for CBWTF).
13. Photo copy of maintained log book for each of its treatment equipment according to weight of batch; categories of waste treated; time, date and duration of treatment cycle and total hours of operation (for CBWTF).
14. Previous authorization copy with its compliance report.
15. Details of Bio Medical Waste generation with category and quantity.

Note :-

i. Copies of all documents submitted by the applicant shall be duly signed and attested by the authorized signatory:

"I.................................. do hereby certify and declare that all the information and documents uploaded by me are true to the best of my knowledge and personally authorized for the same."

ii. The proponent shall submit all documents in serial order as above index only, otherwise application will not be considered.
Check-list of documents for submission of applications for authorization under Hazardous and other Wastes (Management and Transboundary Movement) Rules, 2016

Documents to be uploaded:

1. Authorization to file application.
2. Duly filled up application form.
3. Copy of valid Consent to Operate (CTO) and its compliance.
4. Copy of valid Consent to Establish (CTE).
5. Registration by District Industry Center, if applicable.
6. Copy of annual return containing the details specified in Form 4.
7. Previous authorization copy with its compliance report.
10. List of facilities for collection, storage, packaging, transportation, treatment, processing, use, destruction, recycling, recovery, pre-processing, co-processing, utilization, offering for sale, transfer or disposal of the hazardous waste.
11. Self-certificate regarding “no storage of Hazardous waste beyond 90 days”.
12. Self-certificate regarding “The labelling is being done as per Form 8 for safe handling, storage and transport of hazardous waste”.
13. Photo copy of Manifest (White) (for general).
15. Details of technical capabilities and equipment complying with the standard operating procedure or other guidelines specified by the Central Pollution Control Board (for TSDF only).
16. Photo copy of Form 3 in which the unit maintains the record of hazardous and other Wastes. (for TSDF only).
17. Photocopy of Passbook (for re-user, recycler, re-processor only).
18. Photo copy of Manifest (Green) (for re-user, recycler, re-processor).

Note:-

i. Copies of all documents submitted by the applicant shall be duly signed and attested by the authorized signatory:

"I .................., do hereby certify and declare that all the information and documents uploaded by me are true to the best of my knowledge and personally authorized for the same."

ii. The proponent shall submit all documents in serial order as above index only, otherwise application will not be considered.
Check-list of documents for submission of applications for
Consent to Operate (CTO)

Documents to be uploaded:

1. Authorization to file application.
2. EC copy with compliance report, if applicable.
3. Copy of CTE and CTO with compliance report (with required documents).
4. Copy of authorization under Hazardous and other Wastes (Management, and Transboundary Movement) Rules, 2016, if these Rules apply to the industry.
5. Details of the pollution control devices preferably accompanied with analysis of treated effluent/emission/solid waste, as the case may be.
6. Detail of effluent treatment plant with design if applicable.
7. Details of air pollution control measures implemented.
8. Self-declaration/Affidavit for procuring valid source of raw material.
10. Affidavit certifying that no expansion/modernization in the plant has not been made after construction of the plant after obtaining CTE.
11. Detail of all sources of emission and fugitive emission and pollution control arrangement at its points.
19. Estimated Project Cost including cost of land, building and plant & machinery (without depreciation) duly certified by CA.
12. Land document such as sale deed/registered land lease deed/rent deed.
13. Latitude and longitude of the unit at four points.

Note:-

i. Copies of all documents submitted by the applicant shall be duly signed and attested by the authorized signatory:

"I, ................................, do hereby certify and declare that all the information and documents uploaded by me are true to the best of my knowledge and personally authorized for the same."

ii. The proponent shall submit all documents in serial order as above index only, otherwise application will not be considered.
15. An affidavit declaring that terms and conditions of the Board shall be abided by the PP, pollution control devices mentioned in item number 12 & 13 shall be installed before commissioning of the unit.

16. Self-declaration/Affidavit for procuring valid source of raw material.

17. Latitude and longitude of the proposed site at four points of battery point.

18. AAQ analysis reports of the proposed area at three points.

Expansion project in existing premises:

(a) Old CTE copy and current CTO copy with its compliance report.

(b) Expansion EC if required.

(c) AAQ report.

(d) Proposed site is out of existing premises then, upload documents mentioned from sl. No. 1 to 18.

Note:-

1. Copies of all documents submitted by the applicant shall be duly signed and attested by the authorized signatory.

   "I, ------------------ do hereby certify and declare that all the information and documents uploaded by me are true to the best of my knowledge and personally authorized for the same."

ii. The proponent shall submit all documents in serial order as above index only, otherwise application will not be considered.
Process for obtaining Consent to Establish (CTE)/Consent to Operate (CTO)/Authorization:

The industry should visit Online Consent Management and Monitoring System (OCMMS) by clicking from Home Page of Single Window portal advantage,jharkhand.gov.in” (New user) or existing user can reach the site through “www.jspcb.nic.in”.

For getting CTE/CTO:

a) Read the industry user manual.
b) Register your industry and create user ID and Password.

Then again login and apply for CTE/CTO.

For getting CTE/CTO:

a) Read the industry user manual.
b) Register your industry and create user ID and Password.

Then again login and apply for CTE/CTO.

Check-list of documents for submission of applications for CONSENT TO ESTABLISH (CTE)

Documents to be uploaded

1. Authorization to file application.
2. Environmental Clearance from Competent Authority i.e. MOEF/SEIAA, if applicable.
3. Mining lease copy, if mining work would be involved.
4. Whether proposed site is part of Reserve forest/protected forest/deemed forest, if yes, then status of clearance under Forest Conservation Act, 1980.
5. Whether proposed site is part of Wildlife sanctuary/ National Park, if yes, then status of permission from NBWL.
6. Whether proposed site is part of ESZ, if yes, then status of approval from NBWL.
7. Land document such as sale deed/registered land lease deed/rent deed.
8. Proposed Industry to be shown on village map to the scale with marked Battery Point.
9. Certificate from concerned DFO regarding distance of forest, National Park and Wild Life Sanctuary from the proposed site in respect of CTE guidelines.
10. Estimated Project Cost including cost of land, building and plant & machinery (without depreciation) duly certified by CA.
13. Process flow diagram and disposal for effluent treatment plant with design details, if applicable.
14. Details of Air pollution source and proposed air pollution control measures.
Ref No: B-592
Ranchi, Dated: 10/3/15

From,
(I/c) Member Secretary
Jharkhand State Pollution Control Board, Ranchi

To,
All Section Head(HO),
All JEE, SA (HO), R.A(HO)

Sub:- Work schedule and file keeping for quick disposal of applications submitted through (Online Consent Management and Monitoring System) OCMMS-Regarding.
Ref.No-2378 dated 11.07.2014

As you know Online Consent Management and Monitoring System has been started in the Board. For quick disposal of the application the maximum time schedule for the disposal of application had been fixed and made available to you vide above referred letter. Now from 1st April, 2015 online submission of CTE and CTO application has been made mandatory and hard copy of application will not be processed after 31.03.2015. The maximum time schedule prescribed is below for your reference:-

<table>
<thead>
<tr>
<th>Receiving clerk</th>
<th>JEE/ASO/RA/SA</th>
<th>Section head</th>
<th>Competent Authority</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>After proper scrutiny he will forward the application to SH within 07 working days with his comment</td>
<td>1(A) If the application is incomplete he will raise the clarification to the unit within 07 working days. 2(B) If the application is complete in all respect he will forward the application and draft to the Member Secretary within 07 days with their clear recommendation.</td>
<td>10 days for disposal of application</td>
</tr>
<tr>
<td></td>
<td>After receipt of the applications he will forward the application to JEE/ASO/RA/SA within 02 working day</td>
<td></td>
<td></td>
</tr>
<tr>
<td>2</td>
<td>2. He will prepare the draft for CTE/CTO grant with condition if the application fulfills all the criteria and CTE/CTO rejection if it does not fulfill the criteria and send it to the concerned within 10 days.</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

You are requested to abide by the above timeline.

I/c Member Secretary

[Signature]
JHARKHAND STATE POLLUTION CONTROL BOARD
T.A Division Building (Ground Floor) HEC Campus, Dhanbad, Ranchi- 834004,
Phone No:- 0651- 2400852, 2400851, Fax No:- 2400850.

Notification
Ref.No: - 862
Ranchi, Dated: 30/11/15

Auto-renewal of consent to establish (CTE) under the Water (Prevention & Control of Pollution) Act, 1974 and the Air (Prevention & Control of Pollution) Act, 1981.

The auto-renewal of Consent to Establish for the industries based on self-certification shall be applicable only in those cases where there is no increase in production capacity and/or load. The auto-renewal can be made for a maximum of half of the period for which the original Consent to Establish was granted.
Self-certification shall be submitted by the authorized official of the industry duly authorized by owner/board of directors:

The industry shall, however, apply online for renewal of Consent to Establish with requisite fee in case of the following:

(i) If there is increase in the production capacity and/or pollution load; and

(ii) If there is increase in capital investment.

This will come into force with immediate effect.

(Sanjay K. Suman)
Member secretary
GOVERNMENT OF INDIA
MINISTRY OF ENVIRONMENT, FOREST AND CLIMATE CHANGE

NOTIFICATION

New Delhi, the 28th March, 2016

G.S.R. 343(E).-Whereas the Bio-Medical Waste (Management and Handling) Rules, 1998 was published vide notification number S.O. 630 (E) dated the 20th July, 1998, by the Government of India in the erstwhile Ministry of Environment and Forests, provided a regulatory frame work for management of bio-medical waste generated in the country;

And whereas, to implement these rules more effectively and to improve the collection, segregation, processing, treatment and disposal of these bio-medical wastes in an environmentally sound management thereby, reducing the bio-medical waste generation and its impact on the environment, the Central Government reviewed the existing rules;

And whereas, in exercise of the powers conferred by sections 6, 8 and 25 of the Environment (Protection) Act, 1986 (29 of 1986), the Central Government published the draft rules in the Gazette vide number G.S.R. 450 (E), dated the 3rd June, 2015 inviting objections or suggestions from the public within sixty days from the date on which copies of the Gazette containing the said notification were made available to the public;

And whereas, the copies of the Gazette containing the said draft rules were made available to the public on the 3rd June, 2015;

And whereas, the objections or comments received within the specified period from the public in respect of the said draft rules have been duly considered by the Central Government;

Now, therefore, in exercise of the powers conferred by section 6, 8 and 25 of the Environment (Protection) Act, 1986 (29 of 1986), and in supersession of the Bio-Medical Waste (Management and Handling) Rules, 1998, except as respects things done or omitted to be done before such suppression, the Central Government hereby makes the following rules, namely:-

1. Short title and commencement.- (1) these rules may be called the Bio-Medical Waste Management Rules, 2016.

(2) They shall come into force on the date of their publication in the Official Gazette.

2. Application.-

(1) These rules shall apply to all persons who generate, collect, receive, store, transport, treat, dispose, or handle bio medical waste in any form including hospitals, nursing homes, clinics, dispensaries, veterinary institutions, animal houses, pathological laboratories, blood banks, ayush
hospitals, clinical establishments, research or educational institutions, health camps, medical or surgical camps, vaccination camps, blood donation camps, first aid rooms of schools, forensic laboratories and research labs.

(2). These rules shall not apply to,-

(a) radioactive wastes as covered under the provisions of the Atomic Energy Act, 1962(33 of 1962) and the rules made there under;

(b) hazardous chemicals covered under the Manufacture, Storage and Import of Hazardous Chemicals Rules, 1989 made under the Act;

(c) solid wastes covered under the Municipal Solid Waste (Management and Handling) Rules, 2000 made under the Act;

(d) the lead acid batteries covered under the Batteries (Management and Handling) Rules, 2001 made under the Act;

(e) hazardous wastes covered under the Hazardous Wastes (Management, Handling and Transboundary Movement) Rules, 2008 made under the Act;

(f) waste covered under the e-Waste (Management and Handling) Rules, 2011 made under the Act; and

(g) hazardous micro organisms, genetically engineered micro organisms and cells covered under the Manufacture, Use, Import, Export and Storage of Hazardous Microorganisms, Genetically Engineered Micro organisms or Cells Rules, 1989 made under the Act.

3. **Definitions.**- In these rules, unless the context otherwise requires, -

(a) "Act" means the Environment (Protection) Act, 1986 (29 of 1986);

(b) "animal house" means a place where animals are reared or kept for the purpose of experiments or testing;

(c) "authorisation" means permission granted by the prescribed authority for the generation, collection, reception, storage, transportation, treatment, processing, disposal or any other form of handling of bio-medical waste in accordance with these rules and guidelines issued by the Central Government or Central Pollution Control Board as the case may be;

(d) "authorised person" means an occupier or operator authorised by the prescribed authority to generate, collect, receive, store, transport, treat, process, dispose or handle bio-medical waste in accordance with these rules and the guidelines issued by the Central Government or the Central Pollution Control Board, as the case may be;
(e) "biological" means any preparation made from organisms or micro-organisms or product of metabolism and biochemical reactions intended for use in the diagnosis, immunisation or the treatment of human beings or animals or in research activities pertaining thereto;

(f) "bio-medical waste" means any waste, which is generated during the diagnosis, treatment or immunisation of human beings or animals or research activities pertaining thereto or in the production or testing of biological or in health camps, including the categories mentioned in Schedule I appended to these rules;

(g) "bio-medical waste treatment and disposal facility” means any facility wherein treatment, disposal of bio-medical waste or processes incidental to such treatment and disposal is carried out, and includes common bio-medical waste treatment facilities;

(h) “Form” means the Form appended to these rules;

(i) “handling” in relation to bio-medical waste includes the generation, sorting, segregation, collection, use, storage, packaging, loading, transportation, unloading, processing, treatment, destruction, conversion, or offering for sale, transfer, disposal of such waste;

(j) “health care facility” means a place where diagnosis, treatment or immunisation of human beings or animals is provided irrespective of type and size of health treatment system, and research activity pertaining thereto;

(k) “major accident” means accident occurring while handling of bio-medical waste having potential to affect large masses of public and includes toppling of the truck carrying bio-medical waste, accidental release of bio-medical waste in any water body but exclude accidents like needle prick injuries, mercury spills;

(l) “management” includes all steps required to ensure that bio-medical waste is managed in such a manner as to protect health and environment against any adverse effects due to handling of such waste;

(m) "occupier” means a person having administrative control over the institution and the premises generating bio-medical waste, which includes a hospital, nursing home, clinic, dispensary, veterinary institution, animal house, pathological laboratory, blood bank, health care facility and clinical establishment, irrespective of their system of medicine and by whatever name they are called;

(n) "operator of a common bio-medical waste treatment facility” means a person who owns or controls a Common Bio-medical Waste Treatment Facility (CBMWTF) for the collection, reception, storage, transport, treatment, disposal or any other form of handling of bio-medical waste;

(o) “prescribed authority” means the State Pollution Control Board in respect of a State and Pollution Control Committees in respect of an Union territory;

(p) "Schedule” means the Schedule appended to these rules.
4. **Duties of the Occupier.**- It shall be the duty of every occupier to-

(a) take all necessary steps to ensure that bio-medical waste is handled without any adverse effect to human health and the environment and in accordance with these rules;

(b) make a provision within the premises for a safe, ventilated and secured location for storage of segregated biomedical waste in colored bags or containers in the manner as specified in Schedule I, to ensure that there shall be no secondary handling, pilferage of recyclables or inadvertent scattering or spillage by animals and the bio-medical waste from such place or premises shall be directly transported in the manner as prescribed in these rules to the common bio-medical waste treatment facility or for the appropriate treatment and disposal, as the case may be, in the manner as prescribed in Schedule I;

(c) pre-treat the laboratory waste, microbiological waste, blood samples and blood bags through disinfection or sterilisation on-site in the manner as prescribed by the World Health Organisation (WHO) or National AIDS Control Organisation (NACO) guidelines and then sent to the common bio-medical waste treatment facility for final disposal;

(d) phase out use of chlorinated plastic bags, gloves and blood bags within two years from the date of notification of these rules;

(e) dispose of solid waste other than bio-medical waste in accordance with the provisions of respective waste management rules made under the relevant laws and amended from time to time;

(f) not to give treated bio-medical waste with municipal solid waste;

(g) provide training to all its health care workers and others, involved in handling of bio medical waste at the time of induction and thereafter at least once every year and the details of training programmes conducted, number of personnel trained and number of personnel not undergone any training shall be provided in the Annual Report;

(h) immunise all its health care workers and others, involved in handling of bio-medical waste for protection against diseases including Hepatitis B and Tetanus that are likely to be transmitted by handling of bio-medical waste, in the manner as prescribed in the National Immunisation Policy or the guidelines of the Ministry of Health and Family Welfare issued from time to time;

(i) establish a Bar-Code System for bags or containers containing bio-medical waste to be sent out of the premises or place for any purpose within one year from the date of the notification of these rules;

(j) ensure segregation of liquid chemical waste at source and ensure pre-treatment or neutralisation prior to mixing with other effluent generated from health care facilities;

(k) ensure treatment and disposal of liquid waste in accordance with the Water (Prevention and Control of Pollution) Act, 1974 (6 of 1974);
(l) ensure occupational safety of all its health care workers and others involved in handling of bio-
medical waste by providing appropriate and adequate personal protective equipments;

(m) conduct health check up at the time of induction and at least once in a year for all its health care
workers and others involved in handling of bio- medical waste and maintain the records for the
same;

(n) maintain and update on day to day basis the bio-medical waste management register and display
the monthly record on its website according to the bio-medical waste generated in terms of
category and colour coding as specified in Schedule I;

(o) report major accidents including accidents caused by fire hazards, blasts during handling of bio-
medical waste and the remedial action taken and the records relevant thereto, (including nil report)
in Form I to the prescribed authority and also along with the annual report;

(p) make available the annual report on its web-site and all the health care facilities shall make own
website within two years from the date of notification of these rules;

(q) inform the prescribed authority immediately in case the operator of a facility does not collect the
bio-medical waste within the intended time or as per the agreed time;

(r) establish a system to review and monitor the activities related to bio-medical waste management,
either through an existing committee or by forming a new committee and the Committee shall
meet once in every six months and the record of the minutes of the meetings of this committee
shall be submitted along with the annual report to the prescribed authority and the healthcare
establishments having less than thirty beds shall designate a qualified person to review and monitor
the activities relating to bio-medical waste management within that establishment and submit the
annual report;

(s) maintain all record for operation of incineration, hydro or autoclaving etc., for a period of five
years;

(t) existing incinerators to achieve the standards for treatment and disposal of bio-medical waste as
specified in Schedule II for retention time in secondary chamber and Dioxin and Furans within two
years from the date of this notification.

5. Duties of the operator of a common bio-medical waste treatment and disposal facility.-It shall be
the duty of every operator to -

(a) take all necessary steps to ensure that the bio-medical waste collected from the occupier is
transported, handled, stored, treated and disposed of, without any adverse effect to the human
health and the environment, in accordance with these rules and guidelines issued by the Central
Government or, as the case may be, the central pollution control board from time to time;

(b) ensure timely collection of bio-medical waste from the occupier as prescribed under these rules;

(c) establish bar coding and global positioning system for handling of bio-medical waste within one
year;
(d) inform the prescribed authority immediately regarding the occupiers which are not handing over the segregated bio-medical waste in accordance with these rules;

(e) provide training for all its workers involved in handling of bio-medical waste at the time of induction and at least once a year thereafter;

(f) assist the occupier in training conducted by them for bio-medical waste management;

(g) undertake appropriate medical examination at the time of induction and at least once in a year and immunise all its workers involved in handling of bio-medical waste for protection against diseases, including Hepatitis B and Tetanus, that are likely to be transmitted while handling bio-medical waste and maintain the records for the same;

(h) ensure occupational safety of all its workers involved in handling of bio-medical waste by providing appropriate and adequate personal protective equipment;

(i) report major accidents including accidents caused by fire hazards, blasts during handling of bio-medical waste and the remedial action taken and the records relevant thereto, (including nil report) in Form I to the prescribed authority and also along with the annual report;

(j) maintain a log book for each of its treatment equipment according to weight of batch; categories of waste treated; time, date and duration of treatment cycle and total hours of operation;

(k) allow occupier, who are giving waste for treatment to the operator, to see whether the treatment is carried out as per the rules;

(l) shall display details of authorisation, treatment, annual report etc on its web-site;

(m) after ensuring treatment by autoclaving or microwaving followed by mutilation or shredding, whichever is applicable, the recyclables from the treated bio-medical wastes such as plastics and glass, shall be given to recyclers having valid consent or authorisation or registration from the respective State Pollution Control Board or Pollution Control Committee;

(n) supply non-chlorinated plastic coloured bags to the occupier on chargeable basis, if required;

(o) common bio-medical waste treatment facility shall ensure collection of biomedical waste on holidays also;

(p) maintain all record for operation of incineration, hydro or autoclaving for a period of five years; and

(q) upgrade existing incinerators to achieve the standards for retention time in secondary chamber and Dioxin and Furans within two years from the date of this notification.

6. **Duties of authorities.**- The Authority specified in column (2) of Schedule-III shall perform the duties as specified in column (3) thereof in accordance with the provisions of these rules.
7. **Treatment and disposal.** (1) Bio-medical waste shall be treated and disposed of in accordance with Schedule I, and in compliance with the standards provided in Schedule-II by the health care facilities and common bio-medical waste treatment facility.

(2) Occupier shall hand over segregated waste as per the Schedule-I to common bio-medical waste treatment facility for treatment, processing and final disposal:

Provided that the lab and highly infectious bio-medical waste generated shall be pre-treated by equipment like autoclave or microwave.

(3) No occupier shall establish on-site treatment and disposal facility, if a service of `common bio-medical waste treatment facility is available at a distance of seventy-five kilometer.

(4) In cases where service of the common bio-medical waste treatment facility is not available, the Occupiers shall set up requisite biomedical waste treatment equipment like incinerator, autoclave or microwave, shredder prior to commencement of its operation, as per the authorisation given by the prescribed authority.

(5) Any person including an occupier or operator of a common bio medical waste treatment facility, intending to use new technologies for treatment of bio medical waste other than those listed in Schedule I shall request the Central Government for laying down the standards or operating parameters.

(6) On receipt of a request referred to in sub-rule (5), the Central Government may determine the standards and operating parameters for new technology which may be published in Gazette by the Central Government.

(7) Every operator of common bio-medical waste treatment facility shall set up requisite biomedical waste treatment equipments like incinerator, autoclave or microwave, shredder and effluent treatment plant as a part of treatment, prior to commencement of its operation.

(8) Every occupier shall phase out use of non-chlorinated plastic bags within two years from the date of publication of these rules and after two years from such publication of these rules, the chlorinated plastic bags shall not be used for storing and transporting of bio-medical waste and the occupier or operator of a common bio-medical waste treatment facility shall not dispose of such plastics by incineration and the bags used for storing and transporting biomedical waste shall be in compliance with the Bureau of Indian Standards. Till the Standards are published, the carry bags shall be as per the Plastic Waste Management Rules, 2011.

(9) After ensuring treatment by autoclaving or microwaving followed by mutilation or shredding, whichever is applicable, the recyclables from the treated bio-medical wastes such as plastics and glass shall be given to such recyclers having valid authorisation or registration from the respective prescribed authority.

(10) The Occupier or Operator of a common bio-medical waste treatment facility shall maintain a record of recyclable wastes referred to in sub-rule (9) which are auctioned or sold and the same shall be submitted to the prescribed authority as part of its annual report. The record shall be open for inspection by the prescribed authorities.
(11) The handling and disposal of all the mercury waste and lead waste shall be in accordance with the respective rules and regulations.

8. Segregation, packaging, transportation and storage.-(1) No untreated bio-medical waste shall be mixed with other wastes.

(2) The bio-medical waste shall be segregated into containers or bags at the point of generation in accordance with Schedule I prior to its storage, transportation, treatment and disposal.

(3) The containers or bags referred to in sub-rule (2) shall be labeled as specified in Schedule IV.

(4) Bar code and global positioning system shall be added by the Occupier and common bio-medical waste treatment facility in one year time.

(5) The operator of common bio-medical waste treatment facility shall transport the bio-medical waste from the premises of an occupier to any off-site bio-medical waste treatment facility only in the vehicles having label as provided in part ‘A’ of the Schedule IV along with necessary information as specified in part ‘B’ of the Schedule IV.

(6) The vehicles used for transportation of bio-medical waste shall comply with the conditions if any stipulated by the State Pollution Control Board or Pollution Control Committee in addition to the requirement contained in the Motor Vehicles Act, 1988 (59 of 1988), if any or the rules made there under for transportation of such infectious waste.

(7) Untreated human anatomical waste, animal anatomical waste, soiled waste and, biotechnology waste shall not be stored beyond a period of forty–eight hours:

Provided that in case for any reason it becomes necessary to store such waste beyond such a period, the occupier shall take appropriate measures to ensure that the waste does not adversely affect human health and the environment and inform the prescribed authority along with the reasons for doing so.

(8) Microbiology waste and all other clinical laboratory waste shall be pre-treated by sterilisation to Log 6 or disinfection to Log 4, as per the World Health Organisation guidelines before packing and sending to the common bio-medical waste treatment facility.

9. Prescribed authority.-(1) The prescribed authority for implementation of the provisions of these rules shall be the State Pollution Control Boards in respect of States and Pollution Control Committees in respect of Union territories.

(2) The prescribed authority for enforcement of the provisions of these rules in respect of all health care establishments including hospitals, nursing homes, clinics, dispensaries, veterinary institutions, animal houses, pathological laboratories and blood banks of the Armed Forces under the Ministry of Defence shall be the Director General, Armed Forces Medical Services, who shall function under the supervision and control of the Ministry of Defence.
(3) The prescribed authorities shall comply with the responsibilities as stipulated in Schedule III of these rules.

10. **Procedure for authorisation.** Every occupier or operator handling bio-medical waste, irrespective of the quantity shall make an application in Form II to the prescribed authority i.e. State Pollution Control Board and Pollution Control Committee, as the case may be, for grant of authorisation and the prescribed authority shall grant the provisional authorisation in Form III and the validity of such authorisation for bedded health care facility and operator of a common facility shall be synchronised with the validity of the consents.

(1) The authorisation shall be one time for non-bedded occupiers and the authorisation in such cases shall be deemed to have been granted, if not objected by the prescribed authority within a period of ninety days from the date of receipt of duly completed application along with such necessary documents.

(2) In case of refusal of renewal, cancellation or suspension of the authorisation by the prescribed authority, the reasons shall be recorded in writing:

Provided that the prescribed authority shall give an opportunity of being heard to the applicant before such refusal of the authorisation.

(3) Every application for authorisation shall be disposed of by the prescribed authority within a period of ninety days from the date of receipt of duly completed application along with such necessary documents, failing which it shall be deemed that the authorisation is granted under these rules.

(4) In case of any change in the bio-medical waste generation, handling, treatment and disposal for which authorisation was earlier granted, the occupier or operator shall intimate to the prescribed authority about the change or variation in the activity and shall submit a fresh application in Form II for modification of the conditions of authorisation.

11. **Advisory Committee.** (1) Every State Government or Union territory Administration shall constitute an Advisory Committee for the respective State or Union territory under the chairmanship of the respective health secretary to oversee the implementation of the rules in the respective state and to advice any improvements and the Advisory Committee shall include representatives from the Departments of Health, Environment, Urban Development, Animal Husbandry and Veterinary Sciences of that State Government or Union territory Administration, State Pollution Control Board or Pollution Control Committee, urban local bodies or local bodies or Municipal Corporation, representatives from Indian Medical Association, common bio-medical waste treatment facility and non-governmental organisation.

(2) Notwithstanding anything contained in sub-rule (1), the Ministry of Defence shall constitute the Advisory Committee (Defence) under the chairmanship of Director General of Health Services of Armed Forces consisting of representatives from the Ministry of Defence, Ministry of Environment, Forest and Climate Change, Central Pollution Control Board, Ministry of Health and Family Welfare, Armed Forces Medical College or Command Hospital.
(3) The Advisory Committee constituted under sub-rule (1) and (2) shall meet at least once in six months and review all matters related to implementation of the provisions of these rules in the State and Armed Forces Health Care Facilities, as the case may be.

(4) The Ministry of Health and Defence may co-opt representatives from the other Governmental and non-governmental organisations having expertise in the field of bio-medical waste management.

12. Monitoring of implementation of the rules in health care facilities.- (1) The Ministry of Environment, Forest and Climate Change shall review the implementation of the rules in the country once in a year through the State Health Secretaries and Chairmen or Member Secretary of State Pollution Control Boards and Central Pollution Control Board and the Ministry may also invite experts in the field of bio-medical waste management, if required.

(2) The Central Pollution Control Board shall monitor the implementation of these rules in respect of all the Armed Forces health care establishments under the Ministry of Defence.

(3) The Central Pollution Control Board along with one or more representatives of the Advisory Committee constituted under sub-rule (2) of rule 11, may inspect any Armed Forces health care establishments after prior intimation to the Director General Armed Forces Medical Services.

(4) Every State Government or Union territory Administration shall constitute District Level Monitoring Committee in the districts under the chairmanship of District Collector or District Magistrate or Deputy Commissioner or Additional District Magistrate to monitor the compliance of the provisions of these rules in the health care facilities generating bio-medical waste and in the common bio-medical waste treatment and disposal facilities, where the bio-medical waste is treated and disposed of.

(5) The District Level Monitoring Committee constituted under sub-rule (4) shall submit its report once in six months to the State Advisory Committee and a copy thereof shall also be forwarded to State Pollution Control Board or Pollution Control Committee concerned for taking further necessary action.

(6) The District Level Monitoring Committee shall comprise of District Medical Officer or District Health Officer, representatives from State Pollution Control Board or Pollution Control Committee, Public Health Engineering Department, local bodies or municipal corporation, Indian Medical Association, common bio-medical waste treatment facility and registered non-governmental organisations working in the field of bio-medical waste management and the Committee may co-opt other members and experts, if necessary and the District Medical Officer shall be the Member Secretary of this Committee.

13. Annual report.- (1) Every occupier or operator of common bio-medical waste treatment facility shall submit an annual report to the prescribed authority in Form-IV, on or before the 30th June of every year.

(2) The prescribed authority shall compile, review and analyse the information received and send this information to the Central Pollution Control Board on or before the 31st July of every year.
(3) The Central Pollution Control Board shall compile, review and analyse the information received and send this information, along with its comments or suggestions or observations to the Ministry of Environment, Forest and Climate Change on or before 31st August every year.

(4) The Annual Reports shall also be available online on the websites of Occupiers, State Pollution Control Boards and Central Pollution Control Board.

14. **Maintenance of records.**-(1) Every authorised person shall maintain records related to the generation, collection, reception, storage, transportation, treatment, disposal or any other form of handling of bio-medical waste, for a period of five years, in accordance with these rules and guidelines issued by the Central Government or the Central Pollution Control Board or the prescribed authority as the case may be.

(2) All records shall be subject to inspection and verification by the prescribed authority or the Ministry of Environment, Forest and Climate Change at any time.

15. **Accident reporting.**-(1) In case of any major accident at any institution or facility or any other site while handling bio-medical waste, the authorised person shall intimate immediately to the prescribed authority about such accident and forward a report within twenty-four hours in writing regarding the remedial steps taken in Form I.

(2) Information regarding all other accidents and remedial steps taken shall be provided in the annual report in accordance with rule 13 by the occupier.

16. **Appeal.**-(1) Any person aggrieved by an order made by the prescribed authority under these rules may, within a period of thirty days from the date on which the order is communicated to him, prefer an appeal in Form V to the Secretary (Environment) of the State Government or Union territory administration.

(2) Any person aggrieved by an order of the Director General Armed Forces Medical Services under these rules may, within thirty days from the date on which the order is communicated to him, prefer an appeal in Form V to the Secretary, Ministry of Environment, Forest and Climate Change.

(3) The authority referred to in sub-para (1) and (2) as the case may be, may entertain the appeal after the expiry of the said period of thirty days, if it is satisfied that the appellant was prevented by sufficient cause from filing the appeal in time.

(4) The appeal shall be disposed of within a period of ninety days from the date of its filing.

17. **Site for common bio-medical waste treatment and disposal facility.**-(1) Without prejudice to rule 5 of these rules, the department in the business allocation of land assignment shall be responsible for providing suitable site for setting up of common biomedical waste treatment and disposal facility in the State Government or Union territory Administration.
(2) The selection of site for setting up of such facility shall be made in consultation with the prescribed authority, other stakeholders and in accordance with guidelines published by the Ministry of Environment, Forest and Climate Change or Central Pollution Control Board.

18. **Liability of the occupier, operator of a facility**.- (1) The occupier or an operator of a common bio-medical waste treatment facility shall be liable for all the damages caused to the environment or the public due to improper handling of bio-medical wastes.

(2) The occupier or operator of common bio-medical waste treatment facility shall be liable for action under section 5 and section 15 of the Act, in case of any violation.

**SCHEDULE I**

[See rules 3 (e), 4(b), 7(1), 7(2), 7(5), 7 (6) and 8(2)]

_Part-1_

**Biomedical wastes categories and their segregation, collection, treatment, processing and disposal options**

<table>
<thead>
<tr>
<th>Category</th>
<th>Type of Waste</th>
<th>Type of Bag or Container to be used</th>
<th>Treatment and Disposal options</th>
</tr>
</thead>
<tbody>
<tr>
<td>Yellow</td>
<td>(a) Human Anatomical Waste: Human tissues, organs, body parts and fetus below the viability period (as per the Medical Termination of Pregnancy Act 1971, amended from time to time).</td>
<td>Yellow coloured non-chlorinated plastic bags</td>
<td>Incineration or Plasma Pyrolysis or deep burial*</td>
</tr>
<tr>
<td></td>
<td>(b) Animal 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.</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>(c) Soiled Waste: Items contaminated with blood, body fluids like dressings, plaster casts, cotton swabs and</td>
<td></td>
<td>Incineration or Plasma Pyrolysis or deep burial*</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>In absence of above facilities, autoclaving or micro-waving/</td>
</tr>
<tr>
<td>Bags containing residual or discarded blood and blood components.</td>
<td>Hydroclaving followed by shredding or mutilation or combination of sterilization and shredding. Treated waste to be sent for energy recovery.</td>
<td>(d) Expired or Discarded Medicines: Pharmaceutical waste like antibiotics, cytotoxic drugs including all items contaminated with cytotoxic drugs along with glass or plastic ampoules, vials etc. Yellow coloured non-chlorinated plastic bags or containers</td>
<td>Expired cytotoxic drugs and items contaminated with cytotoxic drugs to be returned back to the manufacturer or supplier for incineration at temperature &gt;1200 °C or to common bio-medical waste treatment facility or hazardous waste treatment, storage and disposal facility for incineration at &gt;1200°C Or Encapsulation or Plasma Pyrolysis at &gt;1200°C. All other discarded medicines shall be either sent back to manufacturer or disposed by incineration.</td>
</tr>
<tr>
<td>---</td>
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<td>---</td>
<td>---</td>
</tr>
<tr>
<td>(e) Chemical Waste: Chemicals used in production of biological and used or discarded disinfectants. Yellow coloured containers or non-chlorinated plastic bags</td>
<td>Disposed of by incineration or Plasma Pyrolysis or Encapsulation in hazardous waste treatment, storage and disposal facility.</td>
<td>(f) Chemical Liquid Waste: Liquid waste generated due to use of chemicals in production of biological and used or discarded disinfectants, Silver X-ray film developing liquid, discarded Formalin, infected secretions, aspirated body fluids, liquid from laboratories and floor washings, cleaning, house-keeping and disinfecting activities etc. Separate collection system leading to effluent treatment system</td>
<td>After resource recovery, the chemical liquid waste shall be pre-treated before mixing with other wastewater. The combined discharge shall conform to the discharge norms given in Schedule-III.</td>
</tr>
<tr>
<td>(g) Discarded linen, mattresses, beddings contaminated with blood or body fluid. Non-chlorinated yellow plastic bags or suitable packing material</td>
<td>Non- chlorinated chemical disinfection followed by incineration or Plazma Pyrolysis or for energy recovery. In absence of above facilities, shredding or mutilation or combination of sterilization and shredding. Treated waste to be sent for energy recovery or incineration or Plazma Pyrolysis.</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
### Microbiology, Biotechnology and other clinical laboratory waste:

Blood bags, Laboratory cultures, stocks or specimens of microorganisms, live or attenuated vaccines, human and animal cell cultures used in research, industrial laboratories, production of biological, residual toxins, dishes and devices used for cultures.

<table>
<thead>
<tr>
<th>Color</th>
<th>Description</th>
<th>Treatment Options</th>
</tr>
</thead>
<tbody>
<tr>
<td>Red</td>
<td><strong>Contaminated Waste (Recyclable)</strong>&lt;br&gt;Wastes generated from disposable items such as tubing, bottles, intravenous tubes and sets, catheters, urine bags, syringes (without needles and fixed needle syringes) and vaccutainers with their needles cut) and gloves.</td>
<td>Autoclaving or micro-waving/hydroclaving followed by shredding or mutilation or combination of sterilization and shredding. Treated waste to be sent to registered or authorized recyclers or for energy recovery or plastics to diesel or fuel oil or for road making, whichever is possible. Plastic waste should not be sent to landfill sites.</td>
</tr>
<tr>
<td>White</td>
<td><strong>Waste sharps including Metals:</strong>&lt;br&gt;Needles, syringes with fixed needles, needles from needle tip cutter or burner, scalpels, blades, or any other contaminated sharp object that may cause puncture and cuts. This includes both used, discarded and contaminated metal sharps</td>
<td>Autoclaving or Dry Heat Sterilization followed by shredding or mutilation or encapsulation in metal container or cement concrete; combination of shredding cum autoclaving; and sent for final disposal to iron foundries (having consent to operate from the State Pollution Control Boards or Pollution Control Committees) or sanitary landfill or designated concrete waste sharp pit.</td>
</tr>
<tr>
<td>Blue</td>
<td><strong>Glassware:</strong>&lt;br&gt;Broken or discarded and contaminated glass including medicine vials and ampoules except those contaminated with cytotoxic wastes.</td>
<td>Disinfection (by soaking the washed glass waste after cleaning with detergent and Sodium Hypochlorite treatment) or through autoclaving or microwaving or hydroclaving and then sent for recycling.</td>
</tr>
</tbody>
</table>

Pre-treat to sterilize with non-chlorinated chemicals on-site as per National AIDS Control Organisation or World Health Organisation guidelines thereafter for Incineration.
Disposal by deep burial is permitted only in rural or remote areas where there is no access to common bio-medical waste treatment facility. This will be carried out with prior approval from the prescribed authority and as per the Standards specified in Schedule-III. The deep burial facility shall be located as per the provisions and guidelines issued by Central Pollution Control Board from time to time.

Part -2

(1) All plastic bags shall be as per BIS standards as and when published, till then the prevailing Plastic Waste Management Rules shall be applicable.

(2) Chemical treatment using at least 10% Sodium Hypochlorite having 30% residual chlorine for twenty minutes or any other equivalent chemical reagent that should demonstrate $\log_{10} 4$ reduction efficiency for microorganisms as given in Schedule-III.

(3) Mutilation or shredding must be to an extent to prevent unauthorized reuse.

(4) There will be no chemical pretreatment before incineration, except for microbiological, lab and highly infectious waste.

(5) Incineration ash (ash from incineration of any bio-medical waste) shall be disposed through hazardous waste treatment, storage and disposal facility, if toxic or hazardous constituents are present beyond the prescribed limits as given in the Hazardous Waste (Management, Handling and Transboundary Movement) Rules, 2008 or as revised from time to time.

(6) Dead Fetus below the viability period (as per the Medical Termination of Pregnancy Act 1971, amended from time to time) can be considered as human anatomical waste. Such waste should be handed over to the operator of common bio-medical waste treatment and disposal facility in yellow bag with a copy of the official Medical Termination of Pregnancy certificate from the Obstetrician or the Medical Superintendent of hospital or healthcare establishment.

(7) Cytotoxic drug vials shall not be handed over to unauthorised person under any circumstances. These shall be sent back to the manufactures for necessary disposal at a single point. As a second option, these may be sent for incineration at common bio-medical waste treatment and disposal facility or TSDFs or plasma pyrolys at temperature $>1200^{\circ}C$.

(8) Residual or discarded chemical wastes, used or discarded disinfectants and chemical sludge can be disposed at hazardous waste treatment, storage and disposal facility. In such case, the waste should be sent to hazardous waste treatment, storage and disposal facility through operator of common bio-medical waste treatment and disposal facility only.
(9) On-site pre-treatment of laboratory waste, microbiological waste, blood samples, blood bags should be disinfected or sterilized as per the Guidelines of World Health Organisation or National AIDS Control Organisation and then given to the common bio-medical waste treatment and disposal facility.

(10) Installation of in-house incinerator is not allowed. However in case there is no common biomedical facility nearby, the same may be installed by the occupier after taking authorisation from the State Pollution Control Board.

(11) Syringes should be either mutilated or needles should be cut and or stored in tamper proof, leak proof and puncture proof containers for sharps storage. Wherever the occupier is not linked to a disposal facility it shall be the responsibility of the occupier to sterilize and dispose in the manner prescribed.

(12) Bio-medical waste generated in households during healthcare activities shall be segregated as per these rules and handed over in separate bags or containers to municipal waste collectors. Urban Local Bodies shall have tie up with the common bio-medical waste treatment and disposal facility to pickup this waste from the Material Recovery Facility (MRF) or from the household directly, for final disposal in the manner as prescribed in this Schedule.

SCHEDULE II
[See rule 4(t), 7(1) and 7(6)]

STANDARDS FOR TREATMENT AND DISPOSAL OF BIO-MEDICAL WASTES

1. STANDARDS FOR INCINERATION.-

All incinerators shall meet the following operating and emission standards-

A. Operating Standards

1). Combustion efficiency (CE) shall be at least 99.00%.

2). The Combustion efficiency is computed as follows:

\[
C.E. = \frac{\%CO_2}{\%CO_2 + \%CO} \times 100
\]

3). The temperature of the primary chamber shall be a minimum of 800 °C and the secondary chamber shall be minimum of 1050°C + or - 50°C.

4). The secondary chamber gas residence time shall be at least two seconds.

B. Emission Standards
<table>
<thead>
<tr>
<th>Sl. No.</th>
<th>Parameter</th>
<th>Limiting concentration in mg Nm$^3$ unless stated</th>
<th>Sampling Duration in minutes, unless stated</th>
</tr>
</thead>
<tbody>
<tr>
<td>1.</td>
<td>Particulate matter</td>
<td>50</td>
<td>30 or 1NM$^3$ of sample volume, whichever is more</td>
</tr>
<tr>
<td>2.</td>
<td>Nitrogen Oxides NO and NO$_2$ expressed as NO$_2$</td>
<td>400</td>
<td>30 for online sampling or grab sample</td>
</tr>
<tr>
<td>3.</td>
<td>HCl</td>
<td>50</td>
<td>30 or 1NM$^3$ of sample volume, whichever is more</td>
</tr>
<tr>
<td>4.</td>
<td>Total Dioxins and Furans</td>
<td>0.1ngTEQ/Nm$^3$ (at 11% O$_2$)</td>
<td>8 hours or 5NM$^3$ of sample volume, whichever is more</td>
</tr>
<tr>
<td>5.</td>
<td>Hg and its compounds</td>
<td>0.05</td>
<td>2 hours or 1NM$^3$ of sample volume, whichever is more</td>
</tr>
</tbody>
</table>

**C. Stack Height:** Minimum stack height shall be 30 meters above the ground and shall be attached with the necessary monitoring facilities as per requirement of monitoring of ‘general parameters’ as notified under the Environment (Protection) Act, 1986 and in accordance with the Central Pollution Control Board Guidelines of Emission Regulation Part-III.

**Note:**
(a) The existing incinerators shall comply with the above within a period of two years from the date of the notification.

(b) The existing incinerators shall comply with the standards for Dioxins and Furans of 0.1ngTEQ/Nm$^3$, as given below within two years from the date of commencement of these rules.

(c) All upcoming common bio-medical waste treatment facilities having incineration facility or captive incinerator shall comply with standards for Dioxins and Furans.

(d) The existing secondary combustion chambers of the incinerator and the pollution control devices shall be suitably retrofitted, if necessary, to achieve the emission limits.

(e) Wastes to be incinerated shall not be chemically treated with any chlorinated disinfectants.

(f) Ash from incineration of biomedical waste shall be disposed of at common hazardous waste treatment and disposal facility. However, it may be disposed of in municipal landfill, if the toxic metals in incineration ash are within the regulatory quantities as defined under the Hazardous Waste (Management and Handling and Transboundary Movement) Rules, 2008 as amended from time to time.

(g) Only low Sulphur fuel like Light Diesel Oil or Low Sulphur Heavy Stock or Diesel, Compressed Natural Gas, Liquefied Natural Gas or Liquefied Petroleum Gas shall be used as fuel in the incinerator.
(h) The occupier or operator of a common bio-medical waste treatment facility shall monitor the stack gaseous emissions (under optimum capacity of the incinerator) once in three months through a laboratory approved under the Environment (Protection) Act, 1986 and record of such analysis results shall be maintained and submitted to the prescribed authority. In case of dioxins and furans, monitoring should be done once in a year.

(i) The occupier or operator of the common bio-medical waste treatment facility shall install continuous emission monitoring system for the parameters as stipulated by State Pollution Control Board or Pollution Control Committees in authorisation and transmit the data real time to the servers at State Pollution Control Board or Pollution Control Committees and Central Pollution Control Board.

(j) All monitored values shall be corrected to 11% Oxygen on dry basis.

(k) Incinerators (combustion chambers) shall be operated with such temperature, retention time and turbulence, as to achieve Total Organic Carbon content in the slag and bottom ashes less than 3% or their loss on ignition shall be less than 5% of the dry weight.

(l) The occupier or operator of a common bio-medical waste incinerator shall use combustion gas analyzer to measure CO₂, CO and O₂.

2. Operating and Emission Standards for Disposal by Plasma Pyrolysis or Gasification:

A. Operating Standards:

All the operators of the Plasma Pyrolysis or Gasification shall meet the following operating and emission standards:

1) Combustion Efficiency (CE) shall be at least 99.99%.

2) The Combustion Efficiency is computed as follows.

\[
\text{C.E} = \frac{\% \text{ CO}_2}{\% \text{ CO}_2 + \% \text{ CO}} \times 100
\]

3) The temperature of the combustion chamber after plasma gasification shall be 1050 ± 50 °C with gas residence time of at least 2(two) second, with minimum 3 % Oxygen in the stack gas.

4) The Stack height should be minimum of 30 m above ground level and shall be attached with the necessary monitoring facilities as per requirement of monitoring of ‘general parameters’ as notified under the Environment (Protection) Act, 1986 and in accordance with the CPCB Guidelines of Emission Regulation Part-III.

B. Air Emission Standards and Air Pollution Control Measures

(i) Emission standards for incinerator, notified at Sl No.1 above in this Schedule, and revised from time to time, shall be applicable for the Plasma Pyrolysis or Gasification also.
(ii) Suitably designed air pollution control devices shall be installed or retrofitted with the ‘Plasma Pyrolysis or Gasification to achieve the above emission limits, if necessary.

(iii) Wastes to be treated using Plasma Pyrolysis or Gasification shall not be chemically treated with any chlorinated disinfectants and chlorinated plastics shall not be treated in the system.

C. Disposal of Ash Vitrified Material: The ash or vitrified material generated from the ‘Plasma Pyrolysis or Gasification shall be disposed off in accordance with the Hazardous Waste (Management, Handling and Transboundary Movement) Rules 2008 and revisions made thereafter in case the constituents exceed the limits prescribed under Schedule II of the said Rules or else in accordance with the provisions of the Environment (Protection) Act, 1986, whichever is applicable.

3. STANDARDS FOR AUTOCLAVING OF BIO-MEDICAL WASTE.-

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

(1) When operating a gravity flow autoclave, medical waste shall be subjected to:

(1a) 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

(1b) 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

(1c) 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.

(2) When operating a vacuum autoclave, medical waste shall be subjected to a minimum of three pre-vacuum pulse to purge the autoclave of all air. The air removed during the pre-vacuum, cycle should be decontaminated by means of HEPA and activated carbon filtration, steam treatment, or any other method to prevent release of pathogen. The waste shall be subjected to the following:

(i) a temperature of not less than 121°C and pressure of 15 psi per 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;

(3) Medical waste shall not be considered as 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 indicator 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.
(4) **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.

(5) **Validation test for autoclave:** The validation test shall use four biological indicator strips, one shall be used as a control and left at room temperature, and three shall be placed in the approximate center of three containers with the waste. Personal protective equipment (gloves, face mask and coveralls) shall be used when opening containers for the purpose of placing the biological indicators. At least one of the containers with a biological indicator should be placed in the most difficult location for steam to penetrate, generally the bottom center of the waste pile. The occupier or operator shall conduct this test three consecutive times to define the minimum operating conditions. The temperature, pressure and residence time at which all biological indicator vials or strips for three consecutive tests show complete inactivation of the spores shall define the minimum operating conditions for the autoclave. After determining the minimum temperature, pressure and residence time, the occupier or operator of a common biomedical waste treatment facility shall conduct this test once in three months and records in this regard shall be maintained.

(6) **Routine Test:** A chemical indicator strip or tape that changes colour 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 locations to ensure that the inner content of the package has been adequately autoclaved. The occupier or operator of a common biomedical waste treatment facility shall conduct this test during autoclaving of each batch and records in this regard shall be maintained.

(7) **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 Geobacillus stearothermophilus spores using vials or spore Strips; with at least 1X10⁵ spores. Under no circumstances will an autoclave have minimum operating parameters less than a residence time of 30 minutes, a temperature less than 121°C or a pressure less than 15 psi. The occupier or operator of a common biomedical waste treatment and disposal facility shall conduct this test at least once in every week and records in this regard shall be maintained.

4. **STANDARDS OF MICROWAVING.-**

(1) Microwave treatment shall not be used for cytotoxic, hazardous or radioactive wastes, contaminated animal carcasses, body parts and large metal items.

(2) The microwave system shall comply with the efficacy test or routine tests and a performance guarantee may be provided by the supplier before operation of the limit.

(3) The microwave should completely and consistently kill the bacteria and other pathogenic organisms that are ensured by approved biological indicator at the maximum design capacity of each microwave unit. Biological indicators for microwave shall be Bacillus atrophaeus spores using vials or spore strips with at least 1 x 10⁵ spores per detachable strip. The biological indicator shall be placed with waste and exposed to same conditions as the waste during a normal treatment cycle.
5. **STANDARDS FOR DEEP BURIAL.**

   (1) A pit or trench should be dug about two meters deep. It should be half filled with waste, then covered with lime within 50 cm of the surface, before filling the rest of the pit with soil.

   (2) It must be ensured that animals do not have any access to burial sites. Covers of galvanised iron or wire meshes may be used.

   (3) On each occasion, when wastes are added to the pit, a layer of 10 cm of soil shall be added to cover the wastes.

   (4) Burial must be performed under close and dedicated supervision.

   (5) The deep burial site should be relatively impermeable and no shallow well should be close to the site.

   (6) The pits should be distant from habitation, and located so as to ensure that no contamination occurs to surface water or ground water. The area should not be prone to flooding or erosion.

   (7) The location of the deep burial site shall be authorised by the prescribed authority.

   (8) The institution shall maintain a record of all pits used for deep burial.

   (9) The ground water table level should be a minimum of six meters below the lower level of deep burial pit.

6. **STANDARDS FOR EFFICACY OF CHEMICAL DISINFECTION**

   Microbial inactivation efficacy is equated to “Log10 kill” which is defined as the difference between the logarithms of number of test microorganisms before and after chemical treatment. Chemical disinfection methods shall demonstrate a 4 Log10 reduction or greater for Bacillus Subtilis (ATCC 19659) in chemical treatment systems.

7. **STANDARDS FOR DRY HEAT STERILIZATION**

   Waste sharps can be treated by dry heat sterilization at a temperature not less than 185°C, at least for a residence period of 150 minutes in each cycle, which sterilization period of 90 minutes. There should be automatic recording system to monitor operating parameters.

   (i) **Validation test for Sharps sterilization unit**

   Waste sharps sterilization unit should completely and consistently kill the biological indicator GeobacillusStearothermophilus or Bacillus Atropheausspoers using vials with at least log₁₀ 6 spores per ml. The test shall be carried out once in three months

   (ii) **Routine test**
A chemical indicator strip or tape that changes colour 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 to ensure that the inner content of the sharps has been adequately disinfected. This test shall be performed once in a week and records in this regard shall be maintained.

8. STANDARDS FOR LIQUID WASTE.

(1) The effluent generated or treated from the premises of occupier or operator of a common bio medical waste treatment and disposal facility, before discharge into the sewer should conform to the following limits-

<table>
<thead>
<tr>
<th>PARAMETERS</th>
<th>PERMISSIBLE LIMITS</th>
</tr>
</thead>
<tbody>
<tr>
<td>pH</td>
<td>6.5-9.0</td>
</tr>
<tr>
<td>Suspended solids</td>
<td>100 mg/l</td>
</tr>
<tr>
<td>Oil and grease</td>
<td>10 mg/l</td>
</tr>
<tr>
<td>BOD</td>
<td>30 mg/l</td>
</tr>
<tr>
<td>COD</td>
<td>250 mg/l</td>
</tr>
<tr>
<td>Bio-assay test</td>
<td>90% survival of fish after 96 hours in 100% effluent.</td>
</tr>
</tbody>
</table>

(2) Sludge from Effluent Treatment Plant shall be given to common bio-medical waste treatment facility for incineration or to hazardous waste treatment, storage and disposal facility for disposal.

Schedule III
[See rule 6 and 9(3)]

List of Prescribed Authorities and the Corresponding Duties

<table>
<thead>
<tr>
<th>Sl. No. (1)</th>
<th>Authority (2)</th>
<th>Corresponding Duties (3)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Ministry of Environment, Forest and Climate Change, Government of India</td>
<td>(i) Making Policies concerning bio-medical waste Management in the Country including notification of Rules and amendments to the Rules as and when required.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>(ii) Providing financial assistance for training and awareness programmes on bio-medical waste management related activities to for the State Pollution Control Boards or Pollution Control Committees.</td>
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<td></td>
<td></td>
<td>(iii) Facilitating financial assistance for setting up or up-gradation of common bio-medical waste treatment facilities.</td>
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<td></td>
<td></td>
<td>(iv) Undertake or support operational research and assessment with reference to risks to environment and health due to bio-medical waste and</td>
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<td>2</td>
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<td></td>
<td>Central or State Ministry of Health and Family Welfare, Central Ministry for Animal Husbandry and Veterinary or State Department of Animal Husbandry and Veterinary.</td>
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<td></td>
<td>previously unknown disposables and wastes from new types of equipment.</td>
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<td></td>
<td>(v) Constitution of Monitoring Committee for implementation of the rules.</td>
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<td></td>
<td>(vi) Hearing Appeals and give decision made in Form-V against order passed by the prescribed authorities.</td>
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<td></td>
<td>(viii) Notify the standards or operating parameters for new technologies for treatment of bio medical waste other than those listed in Schedule- I.</td>
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<td></td>
<td>(i) Grant of license to health care facilities or nursing homes or veterinary establishments with a condition to obtain authorisation from the prescribed authority for bio-medical waste management.</td>
<td></td>
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<tr>
<td></td>
<td>(ii) Monitoring, Refusal or Cancellation of license for health care facilities or nursing homes or veterinary establishments for violations of conditions of authorisation or provisions under these Rules.</td>
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<td></td>
<td>(iii) Publication of list of registered health care facilities with regard to bio-medical waste generation, treatment and disposal.</td>
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<td></td>
<td>(iv) Undertake or support operational research and assessment with reference to risks to environment and health due to bio-medical waste and previously unknown disposables and wastes from new types of equipment.</td>
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<td></td>
<td>(v) Coordinate with State Pollution Control Boards for organizing training programmes to staff of health care facilities and municipal workers on bio-medical waste.</td>
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<td></td>
<td>(vi) Constitution of Expert Committees at National or State level for overall review and promotion of clean or new technologies for bio-medical waste management.</td>
<td></td>
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</tbody>
</table>
(vii) Organizing or Sponsoring of trainings for the regulatory authorities and health care facilities on bio-medical waste management related activities.

(viii) Sponsoring of mass awareness campaigns in electronic media and print media.

<table>
<thead>
<tr>
<th></th>
<th>Ministry of Defence</th>
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<td>3</td>
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<td></td>
<td>(i) Grant and renewal of authorisation to Armed Forces health care facilities or common bio-medical waste treatment facilities (Rule 9).</td>
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<td></td>
<td>(ii) Conduct training courses for authorities dealing with management of bio-medical wastes in Armed Forces health care facilities or treatment facilities in association with State Pollution Control Boards or Pollution Control Committees or Central Pollution Control Board or Ministry of Environment, Forest and Climate Change.</td>
<td></td>
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<td></td>
<td>(iii) Publication of inventory of occupiers and bio-medical waste generation from Armed Forces health care facilities or occupiers</td>
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<td></td>
<td>(iv) Constitution of Advisory Committee for implementation of the rules.</td>
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<td></td>
<td>(v) Review of management of bio-medical waste generation in the Armed Forces health care facilities through its Advisory Committee (Rule 11).</td>
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<td></td>
<td>(vi) Submission of annual report to Central Pollution Control Board within the stipulated time period (Rule 13).</td>
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<tr>
<th></th>
<th>Central Pollution Control Board</th>
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<tr>
<td>4</td>
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<td></td>
<td>(i) Prepare Guidelines on bio-medical waste Management and submit to the Ministry of Environment, Forest and Climate Change.</td>
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<tr>
<td></td>
<td>(ii) Co-ordination of activities of State Pollution Control Boards or Pollution Control Committees on bio-medical waste.</td>
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<td></td>
<td>(iii) Conduct training courses for authorities dealing with management of bio-medical waste.</td>
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<td>(iv) Lay down standards for new technologies for</td>
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<tr>
<td><strong>(vi)</strong> Random inspection or monitoring of health care facilities and common bio-medical waste treatment facilities.</td>
<td><strong>(vi)</strong> Random inspection or monitoring of health care facilities and common bio-medical waste treatment facilities.</td>
<td></td>
</tr>
<tr>
<td><strong>(vii)</strong> Review and analysis of data submitted by the State Pollution Control Boards on bio-medical waste and submission of compiled information in the form of annual report along with its observations to Ministry of Environment, Forest and Climate Change.</td>
<td><strong>(vii)</strong> Review and analysis of data submitted by the State Pollution Control Boards on bio-medical waste and submission of compiled information in the form of annual report along with its observations to Ministry of Environment, Forest and Climate Change.</td>
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</tr>
<tr>
<td><strong>(viii)</strong> Inspection and monitoring of health care facilities operated by the Director General, Armed Forces Medical Services (Rule 9).</td>
<td><strong>(viii)</strong> Inspection and monitoring of health care facilities operated by the Director General, Armed Forces Medical Services (Rule 9).</td>
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</tr>
<tr>
<td><strong>(ix)</strong> Undertake or support research or operational research regarding bio-medical waste.</td>
<td><strong>(ix)</strong> Undertake or support research or operational research regarding bio-medical waste.</td>
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<tr>
<td><strong>5. State Government of Health or Union Territory Government or Administration</strong></td>
<td><strong>(i)</strong> To ensure implementation of the rule in all health care facilities or occupiers.</td>
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<td><strong>(ii)</strong> Allocation of adequate funds to Government health care facilities for bio-medical waste management.</td>
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<td></td>
<td><strong>(iii)</strong> Procurement and allocation of treatment equipments and make provision for consumables for bio-medical waste management in Government health care facilities.</td>
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<td></td>
<td><strong>(iv)</strong> Constitute State or District Level Advisory Committees under the District Magistrate or Additional District Magistrate to oversee the bio-medical waste management in the Districts.</td>
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<td></td>
<td><strong>(v)</strong> Advise State Pollution Control Boards or Pollution Control Committees on implementation of these Rules.</td>
<td></td>
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<tr>
<td></td>
<td><strong>(vi)</strong> Implementation of recommendations of the Advisory Committee in all the health care facilities.</td>
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</tr>
</tbody>
</table>
| 6. | State Pollution Control Boards or Pollution Control Committees | (i) Inventorisation of Occupiers and data on bio-medical waste generation, treatment & disposal.  
(ii) Compilation of data and submission of the same in annual report to Central Pollution Control Board within the stipulated time period.  
(iii) Grant and renewal, suspension or refusal cancellation or of authorisation under these rules (Rule 7, 8 and 10).  
(iv) Monitoring of compliance of various provisions and conditions of authorisation.  
(v) Action against health care facilities or common bio-medical waste treatment facilities for violation of these rules (Rule 18).  
(vi) Organizing training programmes to staff of health care facilities and common bio-medical waste treatment facilities and State Pollution Control Boards or Pollution Control Committees Staff on segregation, collection, storage, transportation, treatment and disposal of bio-medical wastes.  
(vii) Undertake or support research or operational research regarding bio-medical waste management.  
(viii) Any other function under these rules assigned by Ministry of Environment, Forest and Climate Change or Central Pollution Control Board from time to time.  
(ix) Implementation of recommendations of the Advisory Committee.  
(x) Publish the list of Registered or Authorised (or give consent) Recyclers.  
(xi) Undertake and support third party audits of the common bio-medical waste treatment facilities in their State. |
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<tbody>
<tr>
<td>7</td>
<td>Municipalities or Corporations, Urban Local Bodies and Gram Panchayats</td>
<td>(i) Provide or allocate suitable land for development of common bio-medical waste treatment facilities in their respective jurisdictions as per the guidelines of</td>
</tr>
</tbody>
</table>
Central Pollution Control Board.

(ii) Collect other solid waste (other than the biomedical waste) from the health care facilities as per the Municipal Solid Waste (Management and handling) Rules, 2000 or as amended time to time.

(iii) Any other function stipulated under these Rules.
SCHEDULE IV
[See rule 8(3) and (5)]

Part A

LABEL FOR BIO-MEDICAL WASTE CONTAINERS or BAGS

[Image of biohazard symbol]

HANDLE WITH CARE

[Image of cytotoxic hazard symbol]

HANDLE WITH CARE

Part B

LABEL FOR TRANSPORTING BIO-MEDICAL WASTE BAGS OR CONTAINERS

- Day ...........Month ............
- Year ..........
- Date of generation ................

Waste category Number ........
Waste quantity ...............
Sender's Name and Address Receiver's Name and Address:
Phone Number .......... Phone Number .............
Fax Number ............... Fax Number ............... 
Contact Person .......... Contact Person ........

In case of emergency please contact :
Name and Address :
Phone No.

Note : Label shall be non-washable and prominently visible.

FORM – I
[ (See rule 4(o), 5(i) and 15 (2)]

ACCIDENT REPORTING

1. Date and time of accident :

2. Type of Accident :
3. Sequence of events leading to accident:

4. Has the Authority been informed immediately:

5. The type of waste involved in accident:

6. Assessment of the effects of the accidents on human health and the environment:

7. Emergency measures taken:

8. Steps taken to alleviate the effects of accidents:

9. Steps taken to prevent the recurrence of such an accident:

10. Does your facility have an Emergency Control Policy? If yes give details:

Date: ……………………
Signature ……………………
Place: ……………………
Designation ……………………

FORM - II
(See rule10)

APPLICATION FOR AUTHORISATION OR RENEWAL OF AUTHORITY
(To be submitted by occupier of health care facility or common bio-medical waste treatment facility)

To
The Prescribed Authority
(Name of the State or UT Administration)
Address.

1. Particulars of Applicant:

   (i) Name of the Applicant:
   (In block letters & in full)

   (ii) Name of the health care facility (HCF) or common bio-medical waste treatment facility (CBWTF):

   (iii) Address for correspondence:

   (iv) Tele No., Fax No.:
2. Activity for which authorisation is sought:

<table>
<thead>
<tr>
<th>Activity</th>
<th>Please tick</th>
</tr>
</thead>
<tbody>
<tr>
<td>Generation, segregation</td>
<td></td>
</tr>
<tr>
<td>Collection,</td>
<td></td>
</tr>
<tr>
<td>Storage</td>
<td></td>
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<tr>
<td>packaging</td>
<td></td>
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<tr>
<td>Reception</td>
<td></td>
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<tr>
<td>Transportation</td>
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<tr>
<td>Treatment or processing or conversion</td>
<td></td>
</tr>
<tr>
<td>Recycling</td>
<td></td>
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<tr>
<td>Disposal or destruction</td>
<td></td>
</tr>
<tr>
<td>use</td>
<td></td>
</tr>
<tr>
<td>offering for sale, transfer</td>
<td></td>
</tr>
<tr>
<td>Any other form of handling</td>
<td></td>
</tr>
</tbody>
</table>

3. Application for □ fresh or □ renewal of authorisation (please tick whatever is applicable):

(i) Applied for CTO/CTE Yes/No

(ii) In case of renewal previous authorisation number and date:

(iii) Status of Consents:

(a) under the Water (Prevention and Control of Pollution) Act, 1974

(b) under the Air (Prevention and Control of Pollution) Act, 1981:

4. (i) Address of the health care facility (HCF) or common bio-medical waste treatment facility (CBWTF):

(ii) GPS coordinates of health care facility (HCF) or common bio-medical waste treatment facility (CBWTF):

5. Details of health care facility (HCF) or common bio-medical waste treatment facility (CBWTF):

(i) Number of beds of HCF:

(ii) Number of patients treated per month by HCF:

(iii) Number healthcare facilities covered by CBMWTF: ______

(iv) No of beds covered by CBMWTF: ______

(v) Installed treatment and disposal capacity of CBMWTF:_______ Kg per day

(vi) Quantity of biomedical waste treated or disposed by CBMWTF:_____ Kg/ day

(vii) Area or distance covered by CBMWTF:_____________
(pl. attach map a map with GPS locations of CBMWTF and area of coverage)
(viii) Quantity of Biomedical waste handled, treated or disposed:

<table>
<thead>
<tr>
<th>Category</th>
<th>Type of Waste</th>
</tr>
</thead>
<tbody>
<tr>
<td>Yellow</td>
<td>(a) Human Anatomical Waste:</td>
</tr>
<tr>
<td></td>
<td>(b) Animal Anatomical Waste:</td>
</tr>
<tr>
<td></td>
<td>(c) Soiled Waste:</td>
</tr>
<tr>
<td></td>
<td>(d) Expired or Discarded Medicines:</td>
</tr>
<tr>
<td></td>
<td>(e) Chemical Solid Waste:</td>
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<tr>
<td></td>
<td>(f) Chemical Liquid Waste:</td>
</tr>
<tr>
<td></td>
<td>(g) Discarded linen, mattresses, beddings contaminated with blood or body fluid.</td>
</tr>
<tr>
<td></td>
<td>(h) Microbiology, Biotechnology and other clinical laboratory waste:</td>
</tr>
</tbody>
</table>

| Red       | Contaminated Waste (Recyclable) |

<table>
<thead>
<tr>
<th>White</th>
<th>Waste sharps including Metals:</th>
</tr>
</thead>
</table>

| Blue      | Glassware: |
|           | Metallic Body Implants |

6. Brief description of arrangements for handling of biomedical waste (attach details):
   (i) Mode of transportation (if any) of bio-medical waste:
   (ii) Details of treatment equipment (please give details such as the number, type & capacity of each unit)

| Incinerators : |
| Plasma Pyrolysis: |
| Autoclaves: |
| Microwave: |
| Hydroclave: |
| Shredder: |
| Needle tip cutter or destroyer |
| Sharps encapsulation or concrete pit: |
| Deep burial pits: |
| Chemical disinfection: |
| Any other treatment equipment: |
| No of units | Capacity of each unit |

31
7. Contingency plan of common bio-medical waste treatment facility (CBWTF) (attach documents):
8. Details of directions or notices or legal actions if any during the period of earlier authorisation
9. Declaration

I do hereby declare that the statements made and information given above are true to the best of my knowledge and belief and that I have not concealed any information.

I do also hereby undertake to provide any further information sought by the prescribed authority in relation to these rules and to fulfill any conditions stipulated by the prescribed authority.

Date: 
Place:

Signature of the Applicant
Designation of the Applicant

FORM – III
(See rule 10)

AUTHORISATION

(Authorisation for operating a facility for generation, collection, reception, treatment, storage, transport and disposal of biomedical wastes)

1. File number of authorisation and date of issue......................................................

2. M/s _______________________ an occupier or operator of the facility located at ________________________ is hereby granted an authorisation for;

   Activity
   Please tick
   Generation, segregation
   Collection,
   Storage
   packaging
   Reception
   Transportation
   Treatment or processing or conversion
   Recycling
   Disposal or destruction
   use
   offering for sale, transfer
   Any other form of handling

3. M/s ___________________________ is hereby authorized for handling of biomedical waste as per the capacity given below;
   (i) Number of beds of HCF:
   (ii) Number healthcare facilities covered by CBMWTF: ______
(iii) Installed treatment and disposal capacity:_______ Kg per day
(iv) Area or distance covered by CBMWTF:____________

(v) Quantity of Biomedical waste handled, treated or disposed:

<table>
<thead>
<tr>
<th>Type of Waste</th>
<th>Category</th>
<th>Quantity permitted for Handling</th>
</tr>
</thead>
<tbody>
<tr>
<td>Yellow</td>
<td></td>
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<tr>
<td>Red</td>
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<tr>
<td>White (Translucent)</td>
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<tr>
<td>Blue</td>
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</tbody>
</table>

3. This authorisation shall be in force for a period of ………… Years from the date of issue.

4. This authorisation is subject to 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.

Date …………….  Signature……………………..
Place: ……………..  Designation …………………..

Terms and conditions of authorisation *

1. The authorisation shall comply with the provisions of the Environment (Protection) Act, 1986 and the rules made there under.
2. The authorisation or its renewal shall be produced for inspection at the request of an officer authorised by the prescribed authority.
3. The person authorized shall not rent, lend, sell, transfer or otherwise transport the biomedical wastes without obtaining prior permission of the prescribed authority.
4. Any unauthorised change in personnel, equipment or working conditions as mentioned in the application by the person authorised shall constitute a breach of his authorisation.
5. It is the duty of the authorised person to take prior permission of the prescribed authority to close down the facility and such other terms and conditions may be stipulated by the prescribed authority.

Form - IV
(See rule 13)

ANNUAL REPORT

[To be submitted to the prescribed authority on or before 30th June every year for the period from January to December of the preceding year, by the occupier of health care facility (HCF) or common bio-medical waste treatment facility (CBWTF)]

<table>
<thead>
<tr>
<th>Sl. No.</th>
<th>Particulars</th>
</tr>
</thead>
<tbody>
<tr>
<td>1.</td>
<td>Particulars of the Occupier</td>
</tr>
<tr>
<td></td>
<td>(i) Name of the authorised person (occupier or operator of facility)</td>
</tr>
</tbody>
</table>


<table>
<thead>
<tr>
<th>(ii) Name of HCF or CBMWT</th>
<th>:</th>
</tr>
</thead>
<tbody>
<tr>
<td>(iii) Address for Correspondence</td>
<td>:</td>
</tr>
<tr>
<td>(iv) Address of Facility</td>
<td>:</td>
</tr>
<tr>
<td>(v) Tel. No, Fax. No</td>
<td>:</td>
</tr>
<tr>
<td>(vi) E-mail ID</td>
<td>:</td>
</tr>
<tr>
<td>(vii) URL of Website</td>
<td>:</td>
</tr>
<tr>
<td>(viii) GPS coordinates of HCF or CBMWT</td>
<td>:</td>
</tr>
<tr>
<td>(ix) Ownership of HCF or CBMWT</td>
<td>: (State Government or Private or Semi Govt. or any other)</td>
</tr>
<tr>
<td>(x). Status of Authorisation under the Bio-Medical Waste (Management and Handling) Rules</td>
<td>: Authorisation No.: ……………………………….. valid up to …………</td>
</tr>
<tr>
<td>(xi). Status of Consents under Water Act and Air Act</td>
<td>: Valid up to:</td>
</tr>
</tbody>
</table>

2. Type of Health Care Facility

(i) Bedded Hospital

No. of Beds:…..

(ii) Non-bedded hospital

(Clinic or Blood Bank or Clinical Laboratory or Research Institute or Veterinary Hospital or any other)

(iii) License number and its date of expiry

3. Details of CBMWT

(i) Number healthcare facilities covered by CBMWT

(ii) No of beds covered by CBMWT

(iii) Installed treatment and disposal capacity of CBMWT:

(iv) Quantity of biomedical waste treated or disposed by CBMWT:

4. Quantity of waste generated or disposed in Kg per annum (on monthly average basis)

   Yellow Category : 

   Red Category : 

   White:

   Blue Category :

   General Solid waste:

5. Details of the Storage, treatment, transportation, processing and Disposal Facility

(i) Details of the on-site storage facility

<table>
<thead>
<tr>
<th>Size</th>
</tr>
</thead>
<tbody>
<tr>
<td>Capacity</td>
</tr>
</tbody>
</table>

Provision of on-site storage : (cold storage or any other provision)
<table>
<thead>
<tr>
<th>disposal facilities</th>
<th>Type of treatment equipment</th>
<th>No of units</th>
<th>Cap acty Kg/day</th>
<th>Quantity treated or disposed in kg per annum</th>
</tr>
</thead>
<tbody>
<tr>
<td>Incinerators</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Plasma Pyrolysis</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Autoclaves</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Microwave</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Hydroclave</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Shredder</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Needle tip cutter or destroyer</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sharps</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>encapsulation or concrete pit</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Deep burial pits:</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Chemical disinfection:</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Any other treatment equipment:</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

(iii) Quantity of recyclable wastes sold to authorized recyclers after treatment in kg per annum: Red Category (like plastic, glass etc.)

(iv) No of vehicles used for collection and transportation of biomedical waste:

(v) Details of incineration ash and ETP sludge generated and disposed during the treatment of wastes in Kg per annum:

<table>
<thead>
<tr>
<th></th>
<th>Quantity generated</th>
<th>Where disposed</th>
</tr>
</thead>
<tbody>
<tr>
<td>Incineration Ash</td>
<td></td>
<td></td>
</tr>
<tr>
<td>ETP Sludge</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

(vi) Name of the Common Bio-Medical Waste Treatment Facility Operator through which wastes are disposed of:

(vii) List of member HCF not handed over bio-medical waste:

6 Do you have bio-medical waste management committee? If yes, attach minutes of the meetings held during the reporting period

7 Details trainings conducted on BMW

(i) Number of trainings conducted on BMW Management.
(ii) number of personnel trained
(iii) number of personnel trained at the time of induction
(iv) number of personnel not undergone any training so far
(v) whether standard manual for training is available?
(vi) any other information)

8 Details of the accident occurred during the year
(i) Number of Accidents occurred
(ii) Number of the persons affected
(iii) Remedial Action taken (Please attach details if any)
(iv) Any Fatality occurred, details.

9. Are you meeting the standards of air Pollution from the incinerator? How many times in last year could not met the standards?

Details of Continuous online emission monitoring systems installed

10 Liquid waste generated and treatment methods in place. How many times you have not met the standards in a year?

11 Is the disinfection method or sterilization meeting the log 4 standards? How many times you have not met the standards in a year?

12 Any other relevant information : (Air Pollution Control Devices attached with the Incinerator)

Certified that the above report is for the period from
................................................................................................................................................
................................................................................................................................................
................................................................................................................................................
................................................................................................................................................
........................................................................

Name and Signature of the Head of the Institution

Date:
Place

FORM –V
(See rule 16)

Application for filing appeal against order passed by the prescribed authority

1. Name and address of the person applying for appeal:

2. Number, date of order and address of the authority which passed the order, against which appeal is being made (certified copy of order to be attached):

3. Ground on which the appeal is being made:

4. List of enclosures other than the order referred in para 2 against which appeal is being filed:

Signature ………………………
Date :
Name and Address……………………

[F. No. 3-1/2000-HSMD]

(Bishwanath Sinha)
Joint secretary to the Government of India

******
In supersession of the previous notification No.- JSPCB/EODB/01 dated 09.06.2017 this is notified that the officer/ Agency of the Jharkhand State Pollution Control Board shall submit/upload his/their inspection report(s) of the industry including analysis report(s) of air/water, if any, under the provision of Water(Prevention & Control of Pollution) Act,1974 and Air(Prevention & Control of Pollution) Act,1981 within 24 hours (one working day) after the completion of the inspection and receipt of the analysis report(s) of the samples as indicated above.
INSPECTION PROCEDURE

[Under Water (Prevention and Control of Pollution) Act, 1974 and Air (Prevention and Control of Pollution) Act, 1981]

Regular inspection of Red, Orange-high risk and Orange-low risk industries shall be conducted at the end of 2 years, 3 years and 5 years respectively. For Green category, inspection shall be done once in 15 years. Date of inspection shall be calculated from the last inspection done.

List of establishments to be inspected is generated based on computerized risk based assessment on following parameters for regular inspection:

- Categorization of industries into Red, Orange (low and high) and Green
- Previous compliance (Based on Checklist)
- Time period of inspection lapsed

Weightage of these parameters is as follows:

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Weightage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Categorization : Red</td>
<td>0.40</td>
</tr>
<tr>
<td>Categorization : Orange-High Risk</td>
<td>0.30</td>
</tr>
<tr>
<td>Categorization : Orange-Low Risk</td>
<td>0.20</td>
</tr>
<tr>
<td>Categorization : Green</td>
<td>0.10</td>
</tr>
<tr>
<td>Previous compliance (Based on Checklist) : Non Compliance</td>
<td>0.40</td>
</tr>
<tr>
<td>Previous compliance (Based on Checklist) : Partial Compliance</td>
<td>0.20</td>
</tr>
<tr>
<td>Previous compliance (Based on Checklist) : Full Compliance</td>
<td>0.0</td>
</tr>
<tr>
<td>Time lapsed (0-15 days)</td>
<td>0.05</td>
</tr>
<tr>
<td>Time lapsed (16-30 days)</td>
<td>0.10</td>
</tr>
<tr>
<td>Time lapsed (31-45 days)</td>
<td>0.15</td>
</tr>
<tr>
<td>Time lapsed (46 days and above)</td>
<td>0.20</td>
</tr>
</tbody>
</table>
For new industry registration, following shall be the risk parameters

- Categorization of industries into Red, Orange (low and high) and Green
- Time period of inspection lapsed

Weightage of these parameters is as follows:

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Weightage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Categorization : Red</td>
<td>0.40</td>
</tr>
<tr>
<td>Categorization : Orange-High Risk</td>
<td>0.30</td>
</tr>
<tr>
<td>Categorization : Orange-Low Risk</td>
<td>0.20</td>
</tr>
<tr>
<td>Categorization : Green</td>
<td>0.10</td>
</tr>
<tr>
<td>Time lapsed (1-15 days)</td>
<td>0.15</td>
</tr>
<tr>
<td>Time lapsed (16-25 days)</td>
<td>0.30</td>
</tr>
<tr>
<td>Time lapsed (26-35 days)</td>
<td>0.45</td>
</tr>
<tr>
<td>Time lapsed (36 days and above)</td>
<td>0.60</td>
</tr>
</tbody>
</table>

The establishments are categorized into High, Medium and Low risk establishment as per the following definition:

<table>
<thead>
<tr>
<th>Risk Type</th>
<th>Range</th>
</tr>
</thead>
<tbody>
<tr>
<td>High Risk</td>
<td>0.70 to 1.0</td>
</tr>
<tr>
<td>Medium Risk</td>
<td>0.4 to 0.65</td>
</tr>
<tr>
<td>Low Risk</td>
<td>0.0 to 0.35</td>
</tr>
</tbody>
</table>

a) Computerized allocation of Inspectors shall take place based on the following criteria:

- In case of Consent to Establish (CTE) with total investment more than 20 Lakhs, two officers from Regional office including Regional Officer along with one officer from Head office shall do inspection. Allocation of inspectors shall be done at Head office
- In case of Consent to Establish (CTE) with total investment less than 20 Lakhs, two officers from Regional office including Regional Officer shall do inspection. Allocation of inspectors shall be done at Head office
Incase of Consent to Operate (CTO) and Authorizations:

- If **RED** industry falls in “CAT 17 GPI” (refer to annexure I), one officer from Head office and two officers from Regional Office shall do inspection. Allocation of inspectors shall be done at Head office
- For others, two officers from Regional Office shall do inspection. Allocation of inspectors shall be done at Regional office

**During Inspection**

- Inspection shall be made during regular working hours of the establishment / industry except when special circumstances indicate otherwise
- Before entering into the industry, the Inspector shall inspect surrounding areas along the boundary of the industry to verify if there is any discharge of effluents or dumping of solid or hazardous waste
- Thereafter, the inspector shall inspect the industry. At the beginning of the inspection, the inspector shall locate the occupier / representative of the occupier / person in charge of the industry and inform about the purpose of the proposed inspection
- The inspector shall inspect following areas in the industry premises, in addition to other areas as required, to verify compliance status:
  - Manufacturing area
  - Raw materials and finished products storage area
  - Water storage area
  - Effluent generating sources
  - Effluent collection, storage, treatment and disposal areas
  - Air pollution sources and control systems
  - Solid and hazardous waste generation sources, storage and disposal areas
  - Environment management cell to verify record maintenance
- The inspector shall verify all the records pertaining to production, water pollution & control, air pollution & control, management of solid and hazardous waste.

- The inspector shall verify and collect specific information on the following points:
  - Operational status of the industry (whether in operation / seasonal / sick / closed, etc)
  - Production records for a period of last six months (RG-I copies)
  - Consumption quantities of major raw materials during a period of last six months
  - Fresh water source and storage capacity along with details of tanks.
  - Status of flow meters provided for recording water consumption for various purposes along with meter readings at the time of inspection.
  - Details of Water Cess assessment and payment
  - Effluent generation sources and quantities (as per records)
  - Effluent collection, treatment and disposal details.
  - Operation and maintenance of effluent treatment systems.
  - Status of flow meters provided for recording effluent generation, treatment and disposal along with meter readings at the time of inspection.
  - Air pollution sources and control measures.
  - Operation and maintenance of air pollution control equipment / systems.
  - Status of energy meters provided for effluent treatment and air pollution control systems along with meter readings at the time of inspection
  - Details of solid and hazardous waste generation, storage and disposal

- The Inspector shall collect photographic evidence for observations of noncompliance

- The inspector may seek information / clarifications or documentary evidence from the industry to ascertain compliance status

- If any representative / occupier of the industry willfully delays or obstructs the inspector in carrying out inspection, the same shall be
recorded in the inspection report for taking further necessary action in the matter

- During inspection, the Inspector shall take all safety precautions required such as PPE, helmets, masks, goggles, etc

- At the end of the inspection, the Inspector may inform the representative of the industry about apparent non-compliance observed during inspection so that industry may initiate necessary corrective action wherever required

- Signed copy of the report is given to the establishment

1. Post-inspection activities:

- The inspector shall carefully record all the information collected and observation made during the inspection

- Any additional information may be recorded under remarks

- The inspection report shall be carefully checked and then submitted online within 48 hours of the inspection

- The supporting documents and photographs obtained during inspection, if any shall be uploaded along with the inspection report

- An intimation is sent to the establishment with a copy of the inspection report so that it may log onto the portal and view the submitted inspection report
कार्यालय आदेश

झारखंड राज्य प्रदूषण नियंत्रण पर्षद के अधीन विभिन्न प्रकार के इकाइयों के विरुद्ध प्राप्त शिकायतों के आलोक में Surprise Inspection पर्षद मुख्यालय के आदेश पर ही किया जाएगा।

हृ/-
(शंजय कुमार शुमन)
सभस्त सचिव

झापा, 15-1970

प्रतिलिपि : सभी संबंधित पदाधिकारी/ सभी शाखा प्रमुख मुख्यालय/ सभी क्षेत्रीय पदाधिकारी/ सभी कर्मचारियों को सूचनार्थ।

(शंजय कुमार शुमन)
सभस्त सचिव
Office Order

WHEREAS, as per the guideline of CPCB revised classification of industries in the category of Red, Orange, Green and White was prepared and notified by the Board vide order No B 2436, dated 31.03.2016. The order and the list of industrial units classified under Red (60 nos.), Orange (91 nos.), Green (63 nos.) and White (36 nos.) is also available in Board’s website www.jspcb.org:

AND WHEREAS, further, some additional units were identified by the Board, which need to be classified/exempted under Red/Orange/Green/White categories. The matter was placed before the internal committee for evaluation and as per the recommendation of the committee following units have been placed under Red/Orange/Green categories/exempted and was approved by competent authority in anticipation of Board as per the following list:

<table>
<thead>
<tr>
<th>S.N.</th>
<th>Unit Name</th>
<th>Industries type</th>
<th>Implementing date</th>
</tr>
</thead>
<tbody>
<tr>
<td>1.</td>
<td>Wood Plastic Composite unit</td>
<td>Green</td>
<td>Date of issue of concern letter i.e. 28.11.2017</td>
</tr>
<tr>
<td>2.</td>
<td>Assembling of automobile brakes</td>
<td>Green</td>
<td>Date of issue of concern letter i.e. 28.11.2017</td>
</tr>
<tr>
<td>3.</td>
<td>Water Park</td>
<td>Green</td>
<td>Date of issue of concern letter i.e. 13.12.2017</td>
</tr>
<tr>
<td>4.</td>
<td>Oil Depot</td>
<td>Green</td>
<td>Date of issue of concern letter i.e. 13.12.2017</td>
</tr>
<tr>
<td>5.</td>
<td>Washing of Petroleum empty drum</td>
<td>Green</td>
<td>Date of issue of concern letter i.e. 13.12.2017</td>
</tr>
<tr>
<td>6.</td>
<td>Tyre pyrolysis process</td>
<td>Red</td>
<td>Date of issue of this order</td>
</tr>
<tr>
<td>7.</td>
<td>Acetylene gas</td>
<td>Green</td>
<td>Date of issue of this order</td>
</tr>
<tr>
<td>8.</td>
<td>Sodium Silicate plant</td>
<td>Orange low risk</td>
<td>Date of issue of this order</td>
</tr>
</tbody>
</table>

This is for your information.

Memo No: P-558

Ranchi, Dated: 15/5/18

(Rajeev Lochan Bakshi)
Member Secretary

Copy forwarded to Additional Chief Secretary, Deptt. Of Forest, Environment and Climate Change, Jharkhand, Ranchi/ Director, Industry, Jharkhand, Ranchi for information.

(Rajeev Lochan Bakshi)
Member Secretary

Copy forwarded to All Regional Officer/Section Head for information & necessary action.
Notice

It is brought to the notice of all the industries of Jharkhand that it is mandatory on the part of the industry to obtain the consent to operate from the Jharkhand State Pollution Control Board u/s 25 of Water (Prevention & Control of Pollution) Act, 1974 and u/s 21 of the Air (Prevention & Control of Pollution) Act, 1981. So all such Industries which are not having valid Consent to Operate (CTO), are directed not to operate their plants/units without obtaining consent to operate (CTO) from JSPCB.

Industries running without valid CTO are liable to penal action u/s section 41 to 44 of Water (Prevention & Control of Pollution), Act 1974 and 37 of Air (Prevention & Control of Pollution), Act 1981 which includes imprisonment with fine.

Application for obtaining CTO can be submitted online through "https://www.advantage.jharkhand.gov.in/SingleWindow/".

(Sanjay Kumar Suman)
Member Secretary

Memo No.- β - 2081
Ranchi, Dated- 12/12/11

Copy to all Regional Officers, Jharkhand State Pollution Control Board for information and necessary action.

(Sanjay Kumar Suman)
Member Secretary
झारखण्ड राज्य प्रदूषण नियंत्रण पर्षद

नगर प्रशासन भवन, एच0 ई0 सी0, धौरा, रैंची
दरभाष : (0651) 2400851, 2400852, फॅक्स : 0651-2400850

पत्रांक…………693378
रैंची, डिनांक…………17.5.13

सूचना

सर्वसाधारण को सूचित किया जाता है कि हरी श्रेणी की वैसी इकाइय़ों जो माननीय उच्चतम न्यायालय के वाद संख्या-19628-19629/09-दीपक कुमार बनाम हरियाणा राज्य एवं अन्य में आदेश दिनांक 27.02.2012 से आच्छादित नहीं हैं, में पर्यावरणीय स्वीकृति की आवश्यकता झारखण्ड राज्य में नहीं है। हरी श्रेणी के इकाइय़ों की विस्तृत सूची पर्षद के वेब साइट www.jspcb.org पर उपलब्ध है।

यह भी स्पष्ट किया जाता है कि झारखण्ड राज्य में किसी भी इकाई के विद्युत संयोजन (connection) हेतु झारखण्ड राज्य प्रदूषण नियंत्रण पर्षद से प्रदूषण नियंत्रण संबंधी प्रमाण पत्र की आवश्यकता नहीं है। वर्तमान कि पर्षद के द्वारा विद्युत विच्छेद आदेश सम्बंधित इकाई के विरूद्ध पूर्व में नहीं दिया गया है।

सम्पूर्णात्मक

प्रभारी सदस्य सचिव

प्रतिलिपि: सभी शाखा मुख्य/प्रशासी पदाधिकारी/सहायक वैज्ञानिक पदाधिकारी/कर्मी पर्यावरण अभियंता/वैज्ञानिक सहायक/प्रशाखा पदाधिकारी, मुख्यालय/सभी क्षेत्रीय पदाधिकारी को सूचनार्थ एवं आवश्यक कार्रवाई हेतु प्रस्तुत है।
पर्यावरण नियन्त्रण आयोग
Jharkhand State Pollution Control Board
नगर प्रशासन भवन, एचओसी, धौरण, रांची
दूरभाष: 2400851, 2400852, 2400979 फैक्स: 0651-2400850,
Web: www.jspcb.nic.in (new) www.jspcb.org(old)
E-mail: ranchijspxb@gmail.com

अधिसूचना

परियोजना की 37वीं बैठक (विशेष बैठक) की कार्यवाली संख्या-4 में लिये गये निर्णयानुसार पर्यावरण नियन्त्रण आयोग द्वारा पूर्व में छोटी अवधि के संदर्भ में निर्माण आदेश में निम्नलिखित बिन्दुओं को छोड़कर अन्य मामलों में न्यूनतम सी०टी०० अवधि पौंच वर्ष रखने का निर्णय लिया जाता है:-

क) खनन पट्टों के मामलों में खनन पट्टे की कैदीत अथवा पौंच वर्ष में जो भी कम होगा, लागू होगी।
ख) जमीन संबंधी लीज़ की कैदीत अथवा पौंच वर्ष में जो भी कम होगा, लागू होगी।
ग) 50 लाख से कम पूंजी निवेश के उद्देश्यों द्वारा अधिक का पौंच वर्ष से कम अवधि का सी०टी०० आवेदन कर सकते हैं।
घ) अति प्रभावी इकाईयों के मामलों में यदि परियोजना द्वारा बैंक गारंटी लेना आवश्यक हो, तो बैंक गारंटी कैदीत अथवा पौंच वर्ष में जो भी कम होगा, लागू होगी।

(राजीव लोचन बकरी)
सदस्य सचिव

डांपंक...05-02-2019.

प्रतिलिपि: - सभी संबंधित पदाधिकारी/कर्मचारी को सूचनार्थ एवं आवश्यक कार्यरात्रि हेतु प्रेरित।

(राजीव लोचन बकरी)
सदस्य सचिव