

Standard Operating Procedure For Bio Medical Management	
Document Title: Bio Medical Management	Page 1 of 7
Document Number/Version No.: SOP-eMRDS-03/1.0	Proprietary & Confidential

# **Standard Operating Procedure for Bio Medical Management**

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Standard Operation Procedure, SOP-eMRDS-03/1.0

Standard Operating Procedure For Bio Medical Management	
Document Title: Bio Medical Management	Page 2 of 7
Document Number/Version No.: SOP-eMRDS-03/1.0	Proprietary & Confidential

### APPROVALS

By signing the below section digitally, the individuals listed below have reviewed and approved this document:

#### Prepared By:

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Standard Operating Procedure For Bio Medical Management	
Document Title: Bio Medical Management	Page 3 of 7
Document Number/Version No.: SOP-eMRDS-03/1.0	Proprietary & Confidential

## REVISION HISTORY

This document has been revised as follows:

Document Version	Effective Date (Date of final Approval)	Revised By	Description/Reason for Revision

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Standard Operation Procedure, SOP-eMRDS-03/1.0

Standard Operating Procedure For Bio Medical Management	
Document Title: Bio Medical Management	Page 4 of 7
Document Number/Version No.: SOP-eMRDS-03/1.0	Proprietary & Confidential

## Contents

<b>1.0</b>	<b>OBJECTIVE .....</b>	<b>5</b>
<b>2.0</b>	<b>SCOPE.....</b>	<b>5</b>
<b>3.0</b>	<b>RESPONSIBILITY .....</b>	<b>5</b>
<b>4.0</b>	<b>FREQUENCY .....</b>	<b>5</b>
<b>5.0</b>	<b>GENERAL INFORMATION/DEFINITION .....</b>	<b>5</b>
<b>6.0</b>	<b>PROCEDURE: .....</b>	<b>6</b>

Standard Operating Procedure For Bio Medical Management	
Document Title: Bio Medical Management	Page 5 of 7
Document Number/Version No.: SOP-eMRDS-03/1.0	Proprietary & Confidential

## Standard Operating Procedure (SOP) for Bio-Medical Waste Management (BMWM)

### 1.0 Objective

The objective of this SOP is to establish a systematic process for the management of bio-medical waste (BMW) within our organization, ensuring safe disposal, record-keeping, and compliance with all relevant regulations.

### 2.0 Scope

This SOP applies to all personnel involved in the generation, handling, and disposal of bio-medical waste within the organization. It encompasses the entire lifecycle of BMW, from generation to final disposal.

### 3.0 Responsibility

#### 1. BMW Handler:

- The BMW handler is responsible for segregating and packaging bio-medical waste into Yellow bags, Red bags, White boxes, or Blue boxes, as appropriate for incineration or autoclave treatment.
- The handler must update the system with the date and time of each BMW handover.

#### 2. End User (Microbiology Lab/Safety Dept):

- End users shall log in to the system to acknowledge BMW handling requests and initiate further bio-medical waste management processes.
- End users are responsible for referencing the bio-medical waste handover record register.

#### 3. System Administrator:

- The system administrator is responsible for maintaining the electronic system for bio-medical waste management.
- The administrator ensures the proper functioning of the system, user access, data security, and report generation.

### 4.0 Frequency

- BMW handling and system updates shall occur as needed, with a daily record maintained for each activity.
- Monthly reports shall be generated summarizing BMW handling activities.

### 5.0 General Information/Definition

- Bio-medical waste (BMW) includes waste materials generated during healthcare processes, such as discarded medical equipment, contaminated materials, or biological waste.

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Standard Operating Procedure For Bio Medical Management	
Document Title: Bio Medical Management	Page 6 of 7
Document Number/Version No.: SOP-eMRDS-03/1.0	Proprietary & Confidential

- Yellow bags, Red bags, White boxes, and Blue boxes are used for the segregation and disposal of different types of BMW

**Local Version Control System:**

- A local version control system shall be implemented to manage changes to this SOP within the organization.

**Centralized Version Control Systems**

- A centralized version control system shall be maintained for this SOP to ensure uniformity and consistency in procedures across different departments or locations.

**Distributed Version Control System:**

- In case of multiple facilities or locations, a distributed version control system shall be used to manage local variations while maintaining a central standard.

**Bug fixes** - - Any issues or discrepancies in the BMW management system shall be reported to the system administrator for resolution.

## **6.0 Procedure:**

**1. BMW Handler:**

- Segregate BMW into appropriate containers (Yellow bags, Red bags, White boxes, or Blue boxes) based on type.
- Update the system with the date and time of each BMW handover.

**2. End User (Microbiology Lab/Safety Dept):**

- Log in to the system to acknowledge BMW handling requests.
- Refer to the bio-medical waste handover record register for reference.

**3. System Administrator:**

- Maintain the BMW management system.
- Ensure data security and access control.
- Generate monthly reports as required.
- Address any system issues or bug fixes.

**Abbreviations:**

- BMW: Bio-Medical Waste
- SOP: Standard Operating Procedure
- OHC: Occupational Health and Safety

**REFERENCES:**

- Refer to local and national regulations governing bio-medical waste management.
- Internal documents and guidelines related to bio-medical waste management.

**Confidential and Proprietary**

Standard Operating Procedure For Bio Medical Management	
Document Title: Bio Medical Management	Page 7 of 7
Document Number/Version No.: SOP-eMRDS-03/1.0	Proprietary & Confidential

This SOP ensures the safe and efficient management of bio-medical waste within the organization, promoting compliance with regulatory standards and maintaining comprehensive records for transparency and accountability.

**Confidential and Proprietary**

Standard Operation Procedure, SOP-eMRDS-03/1.0