



# DECODING PHARMA

QUALITY ASSURANCE DEPARTMENT

## STANDARD OPERATING PROCEDURE

<b>Department:</b> Quality Assurance	<b>SOP No.:</b>
<b>Title:</b> Document Retention Period	<b>Effective Date:</b>
<b>Supersedes:</b> Nil	<b>Review Date:</b>
<b>Issue Date:</b>	<b>Page No.:</b>

### 1.0 OBJECTIVE:

To lay down a Procedure for Document Retention Period.

### 2.0 SCOPE:

This Procedure applies for Retention of Critical Documents in .....

### 3.0 RESPONSIBILITY:

Officer /Executive-Concern department: To follow SOP Accordingly.

HOD-Concern department: To ensure Implementation of SOP.

### 4.0 ACCOUNTABILITY:

Head-QA.

### 5.0 PROCEDURE:

#### 5.1 MASTER COPY OF BMR/BPR AND STPs:

Obsolete version of master copy of BMR/BPR and STP shall be retained forever.

#### 5.2 FILLED BMR AND BPR:

Filled copy of BMR and BPR shall be retained for expiry of product plus one year or minimum five years from the date of completion which is more.

#### 5.3 MASTER COPY OF SOP:

Obsolete version of SOPs shall be retained forever.

#### 5.4 MACHINE LOG BOOK:

Machine logbook shall be retained for expiry of product plus one year or minimum five years from the date of last entry which is more.

#### 5.5 CHANGE CONTROL, DEVIATIONS, NON COMPLIENCE, MARKET COMPLAINT AND PRODUCT RECALL RECORD:

Record of Change control, Deviation, Non compliance, Market complaint and Product recall shall be retained forever.

#### 5.6 SITE MASTER FILE:



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Site master file shall retained forever.

### 5.7 **VALIDATION DOCUMENT:**

All validation related documents including BMR/BPR of validation batches should be retained forever.

### 5.8 **PRECAUTION FOR DOCUMENTS TO BE RETAINED FOREVER:**

In case the hard copy of any document is not possible to retain for long time due to un-avoidable reasons, the document shall be scanned on computer and record shall be kept forever on CD.

In case scanning of documents in not possible due to any reason then a duplicate copy shall be prepared, approved by Head-QA or his designate and shall be kept along with original copy.

### 5.9 **PRECAUTION FOR DOCUMENTS TO BE DESTROYED:**

Before destruction of any document, QA Head or his designate shall ensure that no issue pending related to regulatory and/or internal/external investigation(s) against the respective document.

In case any investigation is under proceeding or pending, in such case the document shall be destroyed after completion of investigation(s).

### 6.0 **ABBREVIATIONS:**

SOP	-	Standard Operating Procedure
QA	-	Quality Assurance
BMR	-	Batch Manufacturing Record
BPR	-	Batch Packing Record.
STP	-	Standard Test Procedure
CD	-	Compact Disc

### 7.0 **CROSS REFERENCES:**

NA

### 8.0 **REFERENCES:**

In-House



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### 9.0 ATTACHMENTS:

NA

### 10.0 CIRCULATION LIST

Quality Assurance

Production

Engineering

Quality Control

Warehouse

Personnel & Administration

Purchase

Account

### 11.0 REVISION HISTORY:

SOP NUMBER	REASON FOR CHANGE	VERSION NUMBER	SUPERSEDES	CHANGE CONTROL No.
	New SOP	01	NIL	NA