



**STANDARD OPERATING PROCEDURE**

**Department:** Quality Assurance

**SOP No.:**

**Title:** Assigning Retest/Expiry period of Raw materials, Packing materials, Intermediates and Finished products

**Effective Date:**

**Supersedes:** Nil

**Review Date:**

**Issue Date:**

**Page No.:**

**1.0 OBJECTIVE:**

To lay down a procedure for assigning the retest/expiry date for Raw materials, Packing materials, Intermediates and Finished products.

**2.0 SCOPE:**

This SOP is applicable to all the raw materials, intermediates and finished products used in .....

**3.0 RESPONSIBILITY:**

Head – Quality Control

Head – Quality Assurance

**4.0 DEFINITION(S):**

NA

**5.0 PROCEDURE:**

5.1 Assign retest/Expiry date for the product, raw material and packing materials as given below.

S.No.	Stage/Material	Retest date	Expiry Date
1.	Raw Material (Active)	6 months	As per manufacturer's COA
2.	Raw Material (Inactive/excipient) For biological origin material	12 months 6 months	As per manufacturer's instructions COA
3.	Primary packing material - Foils etc.	12 months	-----
4.	Secondary Packing materials e.g. adhesive rolls, labels etc, which are stored at ambient temperature	12 months	-----
5.	Dispensed Raw materials	7 days	-----
6.	Blend granules	7 days	As per product Specification
7.	Core tablets / Capsules	14 days	As per product specification
8.	Coated tablets	21 days	As per product specification



# PHARMA DEVILS

QUALITY ASSURANCE DEPARTMENT

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### 6.0 ABBREVIATION(S):

COA : Certificate Of Analysis

### 7.0 REFERENCE(S):

NA

### 8.0 ANNEXURE(S):

Nil

### 9.0 REVISION CARD:

S.No.	REVISION No.	REVISION DATE	DETAILS OF REVISION	REASON (S) FOR REVISION