

## PHARMA DEVILS

QUALITY CONTROL DEPARTMENT

STANDARD OPERATING PROCEDURE				
<b>Department:</b> Quality Control	SOP No.:			
Title: Quality Control Instruments Activity Log	<b>Effective Date:</b>			
Supersedes: Nil	Review Date:			
Issue Date:	Page No.:			

#### 1.0 **OBJECTIVE:**

To lay down a procedure for QC instruments Activity log.

#### 2.0 SCOPE:

This procedure is applicable to QC instruments Activity log in the QC laboratory.

#### 3.0 **RESPONSIBILITY:**

Executive, Officer, Sr. Officer – Quality control

Head – Quality control

#### 4.0 PROCEDURE:

- 4.1 Activity log of the instruments shall be filled by the analyst online during the analysis.
- 4.2 Analyst shall complete the activity log in the logbook of each instrument as per Annexure-I.

After each and every activity like Operation, Calibration and Maintenance must be log in

- 4.3 activity log.
- Each activity logbook shall be checked by the section head or his designee daily at end of the shift.
- 4.5 After completion of logbook, section head or his designee shall preserve the logbook in a designated place.

### 5.0 ANNEXURE (S):

Annexure-I: Instrument Activity log

### 6.0 REFERENCE (S):

SOP: Preparation, approval, distribution, control, revision and destruction of Standard Operating Procedure (SOP).

### 7.0 ABBREVIATION (S) / DEFINITION (S):

QC: Quality Control

QA: Quality Assurance



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## **REVISION CARD**

S.No.	REVISION No.	REVISION DATE	DETAILS OF REVISION	REASON (S) FOR REVISION	REFERENCE CHANGE CONTROL No.
1	00			New SOP	-



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# ANNEXURE I INSTRUMENT ACTIVITY LOG

Instrument ID No.:													
S.No.	Date	Material/ Product	A.R. No/ Test B.No. Performed		Operation		Operation				Done By	Checked By	Remark
					Started on	<b>Completed on</b>							