



PHARMA DEVILS

QUALITY CONTROL DEPARTMENT

GENERAL TESTING PROCEDURE

Title: Iodine 0.05 M

SOP No.:		Department:	QC
Effective Date:		Review Date:	
Revision No.:	00	Page No.:	1 of 3
Supersede SOP No.:	Nil		

1.0 OBJECTIVE:

1.1 To lay down a procedure for the preparation and standardisation of 0.05 M Iodine.

2.0 SCOPE:

2.1 It is applicable for the estimation of Raw material, bulk product, intermediate product and finish products.

3.0 RESPONSIBILITY:

3.1 Analyst / Officer / Executive follow the procedure.

3.2 Head-QC are responsible for effective implementation of this SOP.

4.0 REFERENCE:

4.1 BP

5.0 DEFINITION:

5.1 Molarity is the number of mole of substance that are present in the given Volume of the solution.

6.0 PROCEDURE:

6.1 Material and Equipment:

6.1.1 Volumetric flask 1000 ml, Iodine, Potassium iodide, conical flask, record book etc.

6.2 Preparation:

6.2.1 Dissolved 12.7 g of Iodine and 20 g of Potassium Iodide in water and dilute to 1000 ml with the same solvent.

6.3 Standardisation:

6.3.1 To 10 ml of the iodine solution add 1 ml of dilute acetic acid and 50 ml of water. Titrate with 0.1 M sodium thiosulfate, determining the end-point potentiometrically or using starch solution as indicator.



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Calculation: $N_1V_1=N_2V_2$

7.0 Annexures:

7.1 Annexure-I: Molarity Calculation format of Volumetric Solution 0.05 M Iodine.

8.0 Distribution:

8.1 Display copy 1: Instrument Lab

9.0 Abbreviation:

GTP : General Test Procedure

QC : Quality Control laboratories

10.0 Revision History:

10.1 Revision history table:

Document Number	CC Number/Date	Brief Description of Change



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ANNEXURE-I

Molarity Calculation format of Volumetric Solution 0.05 M Iodine

Sr. No.	Date	Qty. Prep.	Batch no.	Primary Std. ID. No.	Primary Std. Weight	Calculation	RSD NMT 0.2%	Mean Molarity	Date of Standardization.
1.									
2.									
3.									

Prepared By (Sign/Date):

Checked By (Sign/Date):