WRS 139-1



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Vetindia Pharmaceuticals Limited

A-6/1, Electronic Complex, Kushaiguda, Hyderabad-62

QUALITY CONTROL DEPARTMENT CERTIFICATE OF ANALYSIS WORKING STANDARD ANALYSIS REPORT Mfg.lic.No:52/HD/AP/95/F/R Name of the Material: LEVAMISOLE HYDROCHLORIDE Material Code: 067 Mfg.Date: 02/2015 Exp.Date: 01/2019 Name of the Manufacturer: Baijingyu Pharmaceutical Co Ltd

Condition: Good Quantity Received: 50 g

Date of Analysis:23/12/2015 Date of Report: 24/12/2015

Inward No:046 A.R.NO: VWSR/046/15

Ref STP No: VIPL/QC/RM/067/A Ref SOP No: VIPL/QC/GEN/017

S.NO	TEST	SPECIFICATION	RESULT
1.	Appearance	White or almost white, crystalline powder	White crystalline powder
2.	Solubility	Freely soluble in water, soluble in ethanol (96%), slightly soluble in methylene chloride.	Complies
3.	Identification	A. Specific optical rotationB. Infrared absorption spectrophotometryC. It gives reaction (a) of chlorides	Complies Complies Complies
4.	Appearance of solution	Solution S is clear and not more intensely coloured than reference solution Y ₇ .	Solution S was clear and not more intensely coloured than reference solution Y ₇
5.	рH	3.0 to 4.5	4.19
6.	Specific optical rotation	-121 to -128	-122.15 ⁰
7.	Related Substances	Individual impurity: NMT 0.1% Total impurity: NMT 0.3%	0.06% 0.08%
8.	Heavy metals	Maximum 20 ppm	< 20 ppm
9.	Loss on drying	Maximum 0.5%	0.13%
10.	Sulphated Ash	Maximum 0.1%	0.039%
11.	Assay	98.5% to 101.0%	98.80%

REMARKS: THE MATERIAL COMPLIES WITH BP2011 SPECIFICATIONS IN THE ABOVE RESPECTS.

ANALYSED BY Latha

24/12/2015

CHECKED BY Prasad 24/12/2015 APPROVED BY
B.Swaroopa Rani 2

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