

Pure & Cure Healthcare Pvt. Ltd.Plot No. 26A-30, Sector-8A, I.I.E., SIDCUL, Ranipur
Haridwar-249403, Uttarakhand, INDIA.**QUALITY CONTROL****CERTIFICATE OF ANALYSIS****(FINISHED PRODUCT)**

Product Name :	VERTIHIST-8		
Generic Name :	Betahistine Tablets IP		
Mfg. Lic. No. :	31/UA/2013	Market:	DOMESTIC
Batch No. :	PA2BD09	A. R. No.:	F20240416070
Mfg. Date :	Apr. 2024	Pack Size:	1x15 Tabs
Expiry date:	Mar. 2026	Pack Type:	Alu-Alu Blister (Sale)
Batch Size :	300000 Tabs	Sampled On:	16/04/24
Product Code :	40050068	Sample Quantity:	150 Tabs
Specification No, Ver No.:	STS/FP/40050068-03	Sampled By:	BHUPENDRA SINGH
Ref. STP No. , Ver No.:	STP/FP/0043	Analyzed By:	PANKAJ KUMAR
Manufactured For :	Healing Pharma India Pvt. Ltd.	Date of Analysis:	16/04/24
Manufactured By :	Pure & Cure Healthcare Pvt. Ltd.	Analysis Completion Date:	26/04/24

S.No.	TEST	ACCEPTANCE CRITERIA	RESULTS
1	Description	White to off white coloured, round, biconvex, plain on both sides uncoated tablets. 15 tablets packed in an alu alu blister of printed aluminium foil and printed base aluminium foil.	White coloured, round, biconvex, plain on both sides uncoated tablets. 15 tablets packed in an alu alu blister of printed aluminium foil and printed base aluminium foil.

	Prepared By QC	Reviewed By QC	Approved By QC
Date	26/04/24	26/04/24	26/04/24
Name	NITIN BHATT	SACHIN CHAUHAN	YOGESH CHAUHAN
Designation	ASST. OPERATOR	DY. MANAGER	SR. MANAGER

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2	Identification	(A) By IR: The IR spectrum of the test preparation should concordant to that of the reference preparation of Betahistine Hydrochloride reference standard or working standard. (B) By HPLC: In the Assay, the principal peak in the chromatogram obtained with the test solution corresponds to the peak in the chromatogram obtained with the reference solution of Betahistine Hydrochloride.	Complies
3	Dimension	As below	As below

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S.No.	TEST	ACCEPTANCE CRITERIA	RESULTS
a.	Diameter	7.1 mm $\pm$ 0.2 mm. (6.9 mm to 7.3 mm)	Min. 7.12 mm Max. 7.15 mm
b.	Thickness	3.5 mm $\pm$ 0.3 mm. (3.2 mm to 3.8 mm)	Min. 3.49 mm Max. 3.53 mm
4	Average weight	138.0 mg $\pm$ 7.5%	136.65 mg
5	Uniformity of weight	Not more than two of individual weight deviate from the average weight by more than 7.5% and none deviates by more than 15%.	-2.5% to +1.6%
6	Disintegration Time	Not more than 15 minutes	Passes (01 Minutes 21 Seconds)
7	Hardness	Not less than 30 Newton	Min. 61 Newton Max. 75 Newton
8	Friability	Not more than 1.0% w/w	0.26 %w/w

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9	Dissolution	Level S1(No. of tablets Tested 6 units): Each unit is not less than 80% (Q) which is equivalent to 85% of the labelled amount is released. Level S2 (No. of tablets tested 6 units) : Average of 12 units (S1+S2) is equal to or greater than 80% and no unit is less than 65% (Q - 15%). Level S3 (No. of tablets tested 12 units): Average of 24 units (S1+S2+S3) is equal to or greater than 80% Not more than 2 units are less than 65% (Q - 15%) and No unit is	94.70%, 91.52%, 91.52%, 94.70%, 88.34%, 95.33%
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		less than 55% (Q - 25%).	

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10	Uniformity of content	The preparation complies with the test if each individual content is 85 to 115 per cent of the average content. The preparation fails to comply with the test if more than one individual content is outside these limits or if one individual content is outside the limits of 75 to 125 per cent of the average content. If one individual content is outside the limits of 85 to 115 per cent of the average content but within the limits of 75 to 125 per cent, repeat the determination using another 20	98.13% to 101.05%

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		dosage units. The preparation complies with the test if not more than one of the individual contents of the total sample of 30 dosage units is outside 85 to 115 per cent of the average content and none is outside the limits of 75 to 125 per cent of the average content.	
11	Related Substance	As below	As below
a.	Betahistine Impurity C	Not more than 2.0%	Below disregard limit
b.	Betahistine Impurity A	Not more than 0.2%	Not detected
c.	Any other Secondary impurity	Not more than 0.2%	Below disregard limit

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d.	Sum of all other secondary impurities	Not more than 2.0%		Below disregard limit
12	Assay - Each uncoated tablet contains:	Shelf Life Limit	Release Limit	
	Betahistine Hydrochloride IP - 8 mg/Tab	NLT 7.60mg/Tab to NMT 8.40mg/Tab (NLT 95.00% to NMT 105.00% of label claimed)	NLT 7.76mg/Tab to NMT 8.40mg/Tab (NLT 97.00% to NMT 105.00% of label claimed)	7.81mg 97.63%

CONCLUSION : The Finished Product complies as per IP Specifications.

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