



SWAPNROOP DRUGS & PHARMACEUTICALS

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CERTIFICATE OF ANALYSIS

Product Name	Valsartan USP	CAS No:	137862-53-4
Batch No.	2605001	Manufactured by	-
Date of Mfg.	MAY-2026	Date of Analysis	20/05/2026
Batch Quantity	-	Retest Date	APR-2031

Sr No	Test	Acceptance Criteria	Results
1	Description	A white to almost white, hygroscopic powder.	White powder.
2	Solubility	Freely soluble in anhydrous ethanol; Sparingly soluble in methylene chlorides; practically insoluble in water.	Complies
3	Identification by - IR	Determine by Infrared absorption spectrophotometry compare the spectrum with that obtained with Valsartan WRS or with the reference spectrum of Valsartan.	Complies
4	Identification by - HPLC	The retention time of the major peak of the Sample solution corresponds to that of the standard solution, as obtained in the Assay.	Complies
5	Absorbance (By UV)	The absorbance divided by the path length is NMT 0.02	0.00755
6	Procedure 1 (By HPLC) Limit of valsartan related compound A	NMT 1.0%	Not detected
7	Procedure 2 (By HPLC) Limit of valsartan Related compound B	NMT 0.2%	Not detected
8	Related compound C	NMT 0.1%	Not detected
9	Any other individual impurity	NMT 0.1%	0.050%
10	Total impurities	NMT 0.3%	0.112%
11	Residue on ignition	NMT 0.1%	0.03%
12	Water	NMT 2.0%	1.10%
13	Assay By HPLC (Anhydrous basis)	NLT 98.0% and NMT 102.0%	99.60%

Storage: Store in air tight container below room Temperature

Conclusion: Product Complies as per above Specifications

Prepared By  Sonali Nirmal Date: 23/05/2026	Checked by  Dadarao Kakde Date: 23/05/2026	Approved By  Nayna Nehete Date: 23/05/2026 
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