



ALVENTA PHARMA LIMITED

VILL. KISHANPURA, TEHSIL BADDI- NALAGARH ROAD, DISTT.- SOLAN (H.P) 174101

Quality Control Department

CERTIFICATE OF ANALYSIS (FINISHED PRODUCT)

Product Name	Telmafox-H	A.R. No.	FG/G/25A1463
Generic Name	Telmisartan and Hydrochlorothiazide Tablets IP	Sample Quantity	60 Tablets
Batch No.:	AGT50891B	Sample Received on	04/09/2025
Batch Size:	1.0 Lac	Analysis Date	04/09/2025
Mfg. Date.	08/2025	Release Date	11/09/2025
Exp. Date	07/2027	Page No.:	Page 1 of 2

Sr. No.	Test Parameter	Acceptance Criteria	Result
1.	Description	White and orange coloured, round shaped, biconvex, both side plain, uncoated bilayered tablet.	White and orange coloured, round shaped, biconvex, both side plain, uncoated bilayered tablet.
2.	Identification	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.	Complies
3.	Average weight	300.00 mg \pm 3%	302.39 mg
4.	Uniformity of weight	\pm 5.0% of its average weight.	Min: 295.38 mg ; Max: 309.43 mg -2.32% ; + 2.33%
5.	Disintegration	Not more than 15 minutes.	03 minutes 02 seconds.
6.	Hardness	Not less than 4.0 Kg/cm ²	7.38 Kg/cm ²
7.	Friability	Not more than 1.0%w/w	0.27 %w/w
8.	Dissolution:		
	For Telmisartan	Not less than 80.00% (Q)	Minimum = 97.14 % Maximum = 105.79 % Average = 101.43 %
	For Hydrochlorothiazide	Not less than 80.00% (Q)	Minimum = 94.26 % Maximum = 103.79 % Average = 97.78 %
9.	Uniformity of content (Hydrochlorothiazide)	The acceptance value (AV) calculated using 10 dosage units must be less than or equal to 15 (L1 =15, L2 =25)	6.16
10.	Assay :		
	Each uncoated bilayered tablet contains:	Claim	Limit
	Telmisartan IP	40 mg	Between 95.00% to 105.00% Between 38.00 mg to 42.00 mg
	Hydrochlorothiazide IP	12.5 mg	Between 90.00% to 107.50% Between 11.25 mg to 13.44 mg
			mg
			%
			39.14 mg
			97.85%
			12.421 mg
			99.37%

Particulars	Prepared By	Checked By	Approved By
Name	Praveen Kumar	Kamal Khat	Tuljeet Kaur
Designation	Executive	Executive	Head QC
Signature			
Date	11/09/2025	11/09/2025	11/09/2025



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11. Microbial Limit Tests:			
i.	Total aerobic microbial count	NMT 1000 cfu/g	40 cfu/g
ii.	Total yeast and mould count	NMT 100 cfu/g	Less than 10 cfu/g
iii.	Pathogens: Escherichia coli	Should be absent/g	Absent/g

Remarks: The above test parameters are complies/ not complies as per IP/BP/USP & In-House Specification.

Particulars	Prepared By	Checked By	Approved By
Name	Barveen Kumar	Karraj Kant	Kuljeet Kaur
Designation	Executive	Executive	Head QC
Signature			
Date	11/09/2025	11/09/2025	11/09/2025