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-AM	**A.R. No.**	FG/G/26A0592
Generic Name	Telmisartan and Amlodipine Tablets IP	**Sample Quantity**	60 Tablets
Batch No.:	AGT60276A	**Sample Received on**	23/03/2026
Batch Size:	1.0 Lac	**Analysis Date**	23/03/2026
Mfg. Date.	02/2026	**Release Date**	28/03/2026
Exp. Date	01/2028	**Page No.:**	Page 1 of 2

Sr. No.	Test Parameter	Acceptance Criteria	Result
1.	**Description**	White and sky blue coloured, round shaped, biconvex, both side plain, uncoated bilayered tablet.	White and sky blue 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 (c).	Complies
3.	**Average weight**	300.00 mg ± 3%	297.85 mg
4.	**Uniformity of weight**	± 5.0% of its average weight.	Min: 293.58 mg ; Max: 304.56 mg -2.42% : +2.83%
5.	**Disintegration**	Not more than 15 minutes.	03 minutes 40 seconds.
6.	**Hardness**	Not less than 4.0 Kg/cm^2	5.6 Kg/cm^2
7.	**Friability**	Not more than 1.0%w/w	0.24 %w/w
8.	**Uniformity of content (For Amlodipine Besylate)**	The acceptance value (AV) calculated using 10 dosage units must be less than or equal to 15 (L1 =15, L2 =25)	8.6
9.	**Dissolution:**		
	For Telmisartan	Not less than 80.0% (Q)	Minimum = 89.54% Maximum = 89.74% Average = 89.65%
	For Amlodipine Besylate	Not less than 80.0% (Q)	Minimum = 88.92% Maximum = 92.26% Average = 91.02%
10.	**Related Substances:**		
	Impurity D	Not more than 1.0%	Not Detected
	Any Secondary impurity	Not more than 0.5%	Not Detected
	Total impurities	Not more than 2.0%	Not Detected

Particulars	Prepared By	Checked By	Approved By
Name	Praveen Kumar	Kamal Rana	Sukh Ram
Designation	Executive	Executive	Head QC
Signature			
Date	28/03/2026	28/03/2026	28/03/2026

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11.	**Assay:**				
	Each uncoated bilayered tablet contains:	**Claim**	**Limit**	**mg**	**%**
	Telmisartan IP	**40 mg**	Between 90.0% to 110.0% (Between 36.0 mg to 44.0 mg)	40.79 mg	101.98%
	Amlodipine Besylate IP eq. to Amlodipine	**5 mg**	Between 90.0% to 110.0% (Between 4.5 mg to 5.5 mg)	5.27 mg	105.37%
12.	**Microbial Limit Tests:**				
i.	**Total aerobic microbial count**	NMT 1000 cfu/g		35 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	Bavesh Kumar	Komal Kant	Subash Rajd
Designation	Executive	Executive	HOD
Signature			
Date	28/03/2026	28/03/2026	28/03/2026