



ALVENTA PHARMA LIMITED

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

Quality Control Department

CERTIFICATE OF ANALYSIS (FINISHED PRODUCT)

Product Name	Telmifox-AM	A.R. No.	FG/G/26A1490
Generic Name	Telmisartan and Amlodipine Tablets IP	Sample Quantity	60 Tablets
Batch No.:	AGT60827B	Sample Received on	06/09/2026
Batch Size:	1.0 Lac	Analysis Date	06/09/2026
Mfg. Date.	08/2026	Release Date	11/09/2026
Exp. Date	07/2028	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%	301.97 mg		
4.	Uniformity of weight	± 5.0% of its average weight.	Min: 299.67 mg ; Max: 304.32 mg -0.76% ; +0.78%		
5.	Disintegration	Not more than 15 minutes.	03 minutes 15 seconds.		
6.	Hardness	Not less than 4.0 Kg/cm ²	6.36 Kg/cm ²		
7.	Friability	Not more than 1.0% w/w	0.22 % 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)	7.08		
9.	Dissolution:				
	For Telmisartan	Not less than 80.0% (Q)	Minimum = 86.01% Maximum = 100.17% Average = 93.26%		
	For Amlodipine Besylate	Not less than 80.0% (Q)	Minimum = 90.87% Maximum = 99.18% Average = 93.39%		
10.	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)	40.90 mg	102.24%

Particulars	Prepared By	Checked By	Approved By
Name	Praveen Kumar	Kamalkant	Sukhdeep
Designation	Executive	Executive	Head of QA
Signature			
Date	11/09/2026	11/09/2026	11/09/2026



ALVENTA PHARMA LIMITED

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

Quality Control Department

CERTIFICATE OF ANALYSIS (FINISHED PRODUCT)

Product Name	Telmifox-AM	A.R. No.	FG/G/26A1490
Generic Name	Telmisartan and Amlodipine Tablets IP	Sample Quantity	60 Tablets
Batch No.:	AGT60827B	Sample Received on	06/09/2026
Batch Size:	1.0 Lac	Analysis Date	06/09/2026
Mfg. Date.	08/2026	Release Date	11/09/2026
Exp. Date	07/2028	Page No.:	Page 2 of 2

Sr. No.	Test Parameter	Acceptance Criteria		Result	
	Amlodipine Besylate IP eq. to Amlodipine	5 mg	Between 90.0% to 110.0% (Between 4.5 mg to 5.5 mg)	5.07 mg	101.36%
11.	Microbial Limit Tests:				
i.	Total aerobic microbial count	NMT 1000 cfu/g		30 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	Harveen Kumar	Kamal Kant	Sushant Redi
Designation	Executive	Executive	Head QC
Signature			
Date	11/09/2026	11/09/2026	11/09/2026