



WINDLAS BIOTECH LIMITED

40/1 Mohabewala Industrial Area Dehradun-248110 (Uttarakhand)

Approved by: FDA Uttarakhand, GLP certified Lab

FORM 39 [See rule 150E(f)]

Report of Analysis by Approved Institution

Lic. No.: 1/UA/CTL*/2024

CERTIFICATE OF ANALYSIS FOR FINISHED PRODUCT

Product Name	TELMAFOX 40 TABLET (SALE)		
Generic Name/ Dosage Form Strength (mg)	Telmisartan Tablets IP 40 mg		
Batch No.	TXV26003	Booking ID/A.R. No.	QC/FG/26/11128
Batch Size	100000 STRIP	Pack Size	01 X 10 TABLET
Mfg. Date	08/2026	Exp. Date	07/2028
Reference	I.P	Sample Qty.	1x75 TABLET
Sample Received on	10/08/2026	Manufacturer Lic. No	34/UA/2013,55/UA/SC/P-2013
Protocol/STP No.	WBL/STP/FG/1155	Ref. SPS No.	WBL/FPS/FG/8594
Date of Analysis started on	11/08/2026	Date of Analysis completed on	13/08/2026
Sample Send by and Address	WINDLAS BIOTECH LIMITED (PLANT-2) ,Khasra No. 141 to 143 & 145, Mohabewala Industrial Area, Dehradun-248110 UK		
Manufactured By	WINDLAS BIOTECH LIMITED (PLANT-2)	Page No.	01 of 03

S.No.	Test	Specification	Result
1	Description	Pink coloured, round, biconvex, film coated tablets, plain on both sides.	Pink coloured, round, biconvex, film coated tablets, plain on both sides.
2	Identification	In the Assay, the principal peak in the chromatogram obtained with the test solution correspond to the peak in the chromatogram obtained with the reference solution.	Complies
3	Disintegration Time	Not more than 30 minutes	08 min 35 sec
4	Average weight	175.75 to 194.25 mg	182.47 mg
5	Uniformity of weight	Not more than 2 tablets in 20 deviates from the average weight by more than 7.5%. No tablet deviates from the average weight by more than 15%.	-3.69% to +4.95%

	Prepared By-QC	Checked By-QC	Reviewed By-Lab.-QA	Approved By- Person-in-charge
Name	Ambika	Ravi Kumar	Manoj Gautam	Jogendra Singh
Designation	Trainee	Assistant Manager	Assistant Manager	G. M.
Date	13/08/2026	13/08/2026	13/08/2026	13/08/2026

Printed By: Vanshita Singh

Date & Time : 14/08/2026 10:49:00

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Manufactured By	WINDLAS BIOTECH LIMITED (PLANT-2)	Page No.	02 of 03

S.No.	Test	Specification	Result
6	Dissolution	Not less than 75% (Q) of the stated amount in 30 minutes. Stage: S1 Number tested: 6 Each unit is not less than Q+5 percent of the labelled content. Stage: S2 Number tested: 6 Average of 12 units (S1+S2) is equal to or greater than Q, and no unit is less than Q-15 percent of the labelled content. Stage: S3 Number tested: 12 Average of 24 units (S1+S2+S3) is equal to or greater than Q, not more than 2 units are less than Q-15 percent of the labelled content and no unit is less than Q-25 percent of the labelled content.	Min-91.2% Max-99.6% Avg-96.4%
7	Uniformity of Dosage Unit (By Content uniformity)	Acceptance value should be less than 15	1.96
8	Related substances		
8a.	Any other secondary peak	Not more than 0.2%	0.045%

	Prepared By-QC	Checked By-QC	Reviewed By-Lab.-QA	Approved By- Person-in-charge
Name	Ambika	Ravi Kumar	Manoj Gautam	Jogendra Singh
Designation	Trainee	Assistant Manager	Assistant Manager	G. M.
Date	13/08/2026	13/08/2026	13/08/2026	13/08/2026

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S.No.	Test	Specification	Result
9	Assay Each film coated tablet contains:		
9a.	Telmisartan IP...40mg	90.0 to 110.0% of labeled amount	99.0%

Remarks : The results of analysis relate only to the analysis applied.

In the opinion of the undersigned, the sample referred to above is of standard Quality as defined in the Act and the rules made thereunder for the reason given below.

	Prepared By-QC	Checked By-QC	Reviewed By-Lab.-QA	Approved By- Person-in-charge
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