



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	ATORVAHEAL 40 TABLET (SALE)		
Generic Name/ Dosage Form Strength (mg)	Atorvastatin Tablets IP 40 mg		
Batch No.	AFJ26001	Booking ID/A.R. No.	QC/FG/26/06197
Batch Size	20000 STRIP	Pack Size	01 X 10 TABLET
Mfg. Date	04/2026	Exp. Date	03/2028
Reference	I.P	Sample Qty.	1x75 TABLET
Sample Received on	02/05/2026	Manufacturer Lic. No	34/UA/2013,55/UA/SC/P-2013
Protocol/STP No.	WBL/STP/FG/0732	Ref. SPS No.	WBL/FPS/FG/8704
Date of Analysis started on	11/05/2026	Date of Analysis completed on	27/05/2026
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	Yellow coloured, round, biconvex, film coated tablets, plain on both sides.	Yellow coloured, round, biconvex, film coated tablets, plain on both sides.
2	Identification	The retention time of the principal peak in the chromatogram of test solution corresponds to that the standard solution obtained as directed in the Assay.	Complies
3	Disintegration Time	Not more than 30 minutes	05 Min 46 Sec
4	Average weight	139.65 to 154.35 mg	147.61 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%.	-5.23% to +4.66%

	Prepared By-QC	Checked By-QC	Reviewed By-Lab.-QA	Approved By- Person-in-charge
Name	Sunil Kumar	Ravi Kumar	Lakshay	Jogendra Singh
Designation	TRAINEE CHEMIST	Assistant Manager	Assistant Manager-QA	G. M.
Date	27/05/2026	27/05/2026	27/05/2026	27/05/2026

Printed By: Vanshita Singh

Date & Time : 28/05/2026 09:35:00

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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 70% (Q) of labeled 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-96.4% Avg-94.6%
7	Uniformity of Dosage Unit (By Weight Variation)	Acceptance value should be less than 15.	6.70
8	Related substances		
8a.	Any secondary peak	Not more than 1.0%	0.149%

	Prepared By-QC	Checked By-QC	Reviewed By-Lab.-QA	Approved By- Person-in-charge
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S.No.	Test	Specification	Result
8b.	Sum of all secondary peak	Not more than 4.0%	0.220%
9	Assay Each film coated tablet contains:		
9a.	Atorvastatin Calcium IP eq. to Atorvastatin...40mg	90.0 to 110.0%	101.6%

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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