



MEDOZ PHARMACEUTICALS PVT LTD
VILL.CHANAL MAJRA, NEAR MANPURA, TEH. NALAGARH ,DIST. SOLAN (H.P.)

QUALITY CONTROL DEPARTMENT
CERTIFICATE OF ANALYSIS FINISHED PRODUCT

Product	LABECANOL-100 Labetalol Hydrochloride Tablets I.P.	A.R.No.	FGT/24/823
		Mfg. Date	DEC.2024
		Exp. Date	NOV.2026
Batch no.	MGT-24823	Date of Receiving	02/01/2025
Batch Size	97942 Tabs	Date of Analysis	02/01/2025
Sample Qty.	60 Tablets	Date of Release	15/01/2025

S. No	Test	Specification	Observation
1.	Description	White coloured, circular, biconvex, break line on one side, uncoated tablets.	White coloured, circular, biconvex, break line on one side, uncoated tablets.
2.	Identification	To Comply	Complies
3.	Avg. wt. of tablet	310.0 mg \pm 5 %	309.57 mg
4.	Uniformity of weight	Not more than 2 of the individual weights of tablets deviate from the average weight by more than \pm 5% & none deviates from the average weight by more than \pm 10 %	Complies
5.	Hardness	Not less than 3.0 kg/cm ²	4.0 kg/cm ²
6.	Friability	Not more than 1.0% w/w	0.45% w/w
7.	Disintegration Time	Not more than 15 minutes	5 Minutes. 48 seconds
8.	Dissolution	Not less than 80%(Q)	Min-96.09%, Max-101.80%, Mean-99.03%
9.	Related Substance	To comply	Complies
10.	Assay: Each Uncoated tablet contains:		
	Labetalol Hydrochloride I.P. 100 mg	Limit:95.0% to 105.0% of the stated amount	99.6190 mg (99.62%)

Remark: In the opinion of the undersigned, the sample complies/~~does not comply~~ with the prescribed standard of quality with respect to the above test as per IP/BP/USP/HP specification.

	Prepared By	Checked By	Approved By
Designation	Officer QC	Executive QC	Manager QC
Name	Elisam Sana	Jai prakash singh	Dr. M. Mishra
Signature			
Date	15/01/2025	15/01/2025	15/01/2025

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