



**Swiss Garniers**  
**BIOTECH PVT. LTD.**

21- Industrial Area, Mehatpur,  
Dist. UNA, (H.P.) - 174 315

**CERTIFICATE OF ANALYSIS**  
**QUALITY CONTROL**

**FINISHED PRODUCT**

**A.R. NUMBER : SG2/2425/QCP/QFG/00281**

<b>PRODUCT NAME</b> : ANGEDIL 5 TABS * 1 X 30	<b>BATCH SIZE</b> : 200000 NOS
<b>BATCH NUMBER</b> : 148TAF008	<b>EXP DATE</b> : 03/2026
<b>MFG DATE</b> : 04/2024	<b>SAMPLE DATE</b> : 29/04/2024
<b>SAMPLE QTY.</b> : 4.000 JAR	<b>A. R. DATE</b> : 30/04/2024
<b>SPECIFICATION NO.</b> : A00055-02-02	

Sr.	TEST NAME	OBSERVATION	SPECIFICATION
1	Appearance	White coloured, round shaped, biconvex, uncoated tablets plain on both sides.	White to off-white coloured, round shaped, biconvex, uncoated tablets plain on both sides.
2	Identification (By HPLC)	Complies	In the test of assay the retention time of principal peak of sample preparation must correspond to the Nicorandil working standard solution.
3	Average weight	175.26 mg	176 mg $\pm$ 2.0 % (172.5 mg to 179.5 mg)
4	Uniformity of weight	- 0.81 % to + 1.26 %	Individual weight of tablet does not deviate by more than $\pm$ 7.5 % from the average weight.
5	Friability	0.18 %	Not More Than 1.0 %
6	Disintegration Time	04 minutes 23 seconds	NMT 15 min in water at 37°C $\pm$ 2°C
7	Dissolution (By HPLC)	97.56 % to 102.66 %	Q. NLT 75 % of the labeled amount of Nicorandil.
8	Uniformity of Dosage Units (By HPLC)	L1 = 5.82	L1 = NMT 15.0

**Assay Details :**

Each uncoated tablet contains:

SRL.	ACTIVE INGREDIENT	RESULT	LIMITS
1	Nicorandil IP	4.9859 mg (99.72 %)	Between 4.5 and 5.5 mg

**REMARKS : THE SAMPLE COMPLIES AS PER IP. SPECIFICATION**

Prepared by   
Date : 30/04/24

Checked by   
Date : 30/04/24

Approved by   
Date : 30/04/24