

DAFFOHILS LABORATORIES PVT. LTD.**(A WHO-GMP, GLP Certified Company)**

F-109-110, UPSIDC Industrial Area, Central Hope Town, Selaqui, Dehradun U.K. - 248 011 INDIA.

CERTIFICATE OF ANALYSIS

Quality Control Department

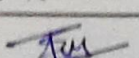
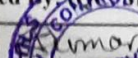
(The Drugs & Cosmetics Act 1940 and rules made there under)

Mfg. Lic. No. 107/UA/2007 & 113/UA/SC/P-2007

Product Name	AMOXHYHEAL-CV DDS DRY SYRUP		
Generic Name	Amoxycillin & Potassium Clavulanate Oral Suspension IP		
Batch No.	DLBS-148	Batch Size	5,000 Bottles
Mfg. Date	10/2023	Exp. Date	03/2025
Report No.	DLBS/10/148/2023	Specification No.	DQC/SP/FP/224
Ref. No.	B010	Sampled Qty	5 Bottles
Date of Sampling	22/11/2023	Date of Release	24/11/2023

S. No.	Test / Parameter	Specification	Observation		
1.	Description	White colored powder filled in transparent glass bottle on reconstitution with water, forms pink color suspension.	White colored powder filled in transparent glass bottle on reconstitution with water, forms pink color suspension.		
2.	Identification	Should be positive for Amoxycillin Trihydrate & Potassium Clavulanate	Positive for Amoxycillin Trihydrate & Potassium Clavulanate		
3.	Nominal fill Weight powder	4.5 g	4.512 g		
4.	Average fill Weight	NLT 4.5 g	Complies		
5.	Nominal Fill Volume	30.0 ml	30.04 ml		
6.	pH(after reconstitution)	3.8 to 6.6	6.22		
7.	Weight/ml (after reconstitution)	1.00 g/ml to 1.40 g/ml	1.113g/ml		
Assay:					
8.	Each 5ml (after reconstitution) suspension contains:	Label Claim	Result	Percentage	Limit
	Amoxycillin Trihydrate IP Eq. to Amoxycillin	400 mg	394.32 mg	98.58%	360.0 mg to 480.0 mg (90% to 120%)
	Potassium Clavulanate Diluted IP Eq. to Clavulanic Acid	57 mg	56.12 mg	98.46%	51.30 mg to 71.25 mg (90% to 125%)

REPORT: The sample complies /Does not comply as per IP/In-house Specification.

Prepared By: (QC.Officer)		Checked By: (QC. Executive)		Approved By: (QC.Manager)	
Signature:		Signature:		Signature:	
Date:	24/11/23	Date:	24/11/23	Date:	24/11/23

