

**DAFFOHILS LABORATORIES PVT. LTD.**

(A WHO-GMP, GLP Certified Company)

F-109-110, UPSIDC Industrial Area, Central Hope Town, Setaqui, 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	AMOXYSHEAL-CV DRY SYRUP		
Generic Name	Amoxycillin & Potassium Clavulanate Oral Suspension IP		
Batch No.	DLBS-152	Batch Size	5000 Bottles
Mfg. Date	10/2023	Exp. Date	03/2026
Report No.	DLBS/10/152/2023	Specification No.	DQC/SP/FP/147
Ref. No.	B014	Sampled Qty	5 Bottles
Date of Sampling	28/10/2023	Date of Release	30/10/2023

S. No.	Test / Parameter	Specification	Observation		
1.	Description	White colored powder filled in transparent glass bottle on reconstitution with water, forms orange color suspension.	White colored powder filled in transparent glass bottle on reconstitution with water, forms orange color suspension.		
2.	Identification	Should be positive for Amoxycillin Trihydrate & Potassium Clavulanate	Positive for Amoxycillin Trihydrate & Potassium Clavulanate		
3.	Nominal fill Weight powder	3.3 g	3.319 g		
4.	Average fill Weight	NLT 3.3 g	Complies		
5.	Nominal Fill Volume	30.0 ml	30.07 ml		
6.	pH(after reconstitution)	3.8 to 6.6	6.26		
7.	Water	NMT 8.5%	6.6%		
8.	Weight/ml (after reconstitution)	1.00 g/ml to 1.40 g/ml	1.114g/ml		
9.	Assay:				
	Each 5ml (after reconstitution) suspension contains:	Label Claim	Result	Percentage	Limit
	Amoxycillin Trihydrate IP Eq. to Amoxycillin	200 mg	197.12 mg	98.56%	180.0 mg to 240.0 mg (90.0% to 120.0%)
	Potassium Clavulanate Diluted IP Eq. to Clavulanic Acid	28.5 mg	28.04 mg	98.39%	25.65 mg to 35.62 mg (90.0% to 125.0%)

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:	30/10/23	Date:	30/10/23	Date:	30/10/23

