[redacted]

Sample Name	CEFADROXIL ORAL SUSPENSION-IP.	**Mfg Lic No**	[redacted]
Submitted By	Quality Control Advisor. (MMDSL) Shillong.	**Report No**	
Address	DHS LaitumkhrahNHM,EastKhasi Hills Meghalaya.	**Receipt Date**	21-Aug-24
Mfg By	N.S.	**Report Date**	30-Aug-24
Supplied By	N.S.	**Ref No**	N.S.

Batch No	Mfg. Date	Exp Date	Batch Size	Sample Qty
MMDSL/QC-0432	06/2024	11/2025	N.S.	12 BOTTLES.

TEST RESULTS

Date / Period of Performance of test 21/08/2024 to 30/08/2024.

Reference to protocol :- I.P-2022.

Description :- Off white powder filled in plastic bottle which gives orange coloured suspension upon reconstitution with water.

Identification (by T.L.C.) :- Complies.
Average net weight :- 9.0933gm.
Uniformity of weight :- Within limit.
Related substances (by HPLC) :- Complies.
Weight per ml :- 1.1007gm/ml

		LIMIT	
		Lower	Upper
pH(after reconstitution)	:- 5.23	4.5	6.0
Water(by KF)	:- 0.92%w/w	-	2.0%w/w

Assay (H.P.L.C.) :- Each 5ml. after reconstitution contains:-

Contents of	Obtained	Claim	LIMIT Lower	LIMIT Upper	Method
Cefadroxil (anhydrous)	:- 124.27mg	125.0mg.	112.5mg.	150.0mg.	IP.
STABILITY AFTER FOUR DAYS	:-				
Cefadroxil (anhydrous)	:- 122.05mg	125.0mg.	100.0mg.	-	IP.

DOSE-CUP.
Reference to protocol :- In House Specification.
Description :- A sample name "DOSE.CUP" was submitted by the party in standard shape & size.
Nominal size :- 10ml.
Overflow capacity :- Passes the test.
Graduation mark :- 2.5ml,5ml,7.5ml & 10ml.
Construction Quality :- Passes the test.
NOTE:- SAMPLE CONSUMED IN TESTING.

Report:In Opinion of the undersigned,The sample referred to above is of Standard Quality as defined in the Act and the rules made there under for the [redacted] per IP. with respect above test only.

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