[redacted]

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| | | | |
|---|---|---|---|
| **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-0434 | 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 :- 9.0839gm.
Uniformity of weight :- Within limit.
Related substances (by HPLC) :- Complies.
Weight per ml :- 1.0948gm/ml

| | | LIMIT | |
|---|---|---|---|
| | | Lower | Upper |
| pH(after reconstitution) | :- 5.26 | 4.5 | 6.0 |
| Water(by KF) | :- 0.93%w/w | - | 2.0%w/w |

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

| | | | LIMIT | | |
|---|---|---|---|---|---|
| Contents of | Obtained | Claim | Lower | Upper | Method |
| Cefadroxil (anhydrous) | :- 124.85mg | 125.0mg. | 112.5mg. | 150.0mg. | IP. |
| STABILITY AFTER FOUR DAYS | :- | | | | |
| Cefadroxil (anhydrous) | :- 122.39mg | 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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