

<b>Sample Name</b>	Cefixime Oral Suspension-IP. 50mg./5ml	<b>Mfg Lic No</b>	[REDACTED]	
<b>Submitted By</b>	Quality Control Advisor. (MMDSL) Shillong.	<b>Report No</b>	[REDACTED]	
<b>Address</b>	DHS LaitumkhrahNHM, EastKhasi Hills Meghalaya.	<b>Receipt Date</b>	21-Aug-24	
<b>Mfg By</b>	N.S.	<b>Report Date</b>	30-Aug-24	
<b>Supplied By</b>	N.S.	<b>Ref No</b>	N.S.	
<b>Batch No</b>	<b>Mfg. Date</b>	<b>Exp Date</b>	<b>Batch Size</b>	<b>Sample Qty</b>
MMDSL/QC-0426	02/2024	01/2026	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 :- Light pink coloured powder filled in plastic bottle which gives pink coloured suspension upon reconstitution with water.

Identification (by HPLC) :- Complies.

		LIMIT	
		Lower	Upper
pH(after reconstitution)	:- 3.91	2.5	4.5

Weight per ml. :- 1.1085gm/ml.

Average net weight :- 13.1012gm.

Uniformity of weight :- Within limit.

Water(by KF) :- 0.961%w/w - 2.0%w/w

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

	Obtained	Claim	LIMIT		Method
			Lower	Upper	
Contents of Cefixime Trihydrate eq. to anhydrous Cefixime	:- 49.58mg	50.0mg.	45.0mg.	60.0mg.	IP.
Stability after 5 days Cefixime Trihydrate eq. to anhydrous Cefixime	:- 48.66mg	50.0mg.	40.0mg.	-	IP.

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 reason given: Complies as per IP.

