

(THE DRUGS & COSMETICS ACT 1940 & RULES THEREUNDER)

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<b>Sample Name</b>	DICLOFENAC POT. & PARACETAMOL TABLETS.	<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>	19-Mar-24	
<b>Mfg By</b>	N.S.	<b>Report Date</b>	02-Apr-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-0110	10/2023	09/2025	N.S.	60 TABS.

**RESULT OF ANALYSIS**

Date / Period of Performance of test 19/03/2024 to 02/04/2024.

Ref. to protocol :- In house specification.

Description :- White elongated biconvex uncoated tablet having scored on one side.

Identification :- Positive for Paracetamol & Diclofenac Potassium.

Average weight :- 816.9mg.

Uniformity of weight :- Within limit.

Disintegration time :- 03 to 04 minutes. (Limit NMT 15 minutes)

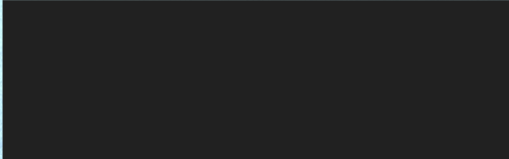
Assay :- Each uncoated tablet contains:-

LIMIT

Content of	Obtd./Av.wt.	Claim	Lower	Upper	Method
Paracetamol	:- 498.61mg	500.0mg.	450.0mg.	550.0mg.	SP-04.
Diclofenac Potassium	:- 49.83mg	50.0mg.	45.0mg.	55.0mg.	HPLC.

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 with respect above test only.



- NOTE :
1. The result listed refer only to the tested samples and applicable parameter. Endorsement of products is neither inferred nor implied.
  2. Sample not drawn by us. Total liability of this laboratory limited to the invoice amount.
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