



Edward Food Research & Analysis Centre Limited



QA.15.0.0.3

**FORM 39**

**[RULE 150 - E(F)] APPROVAL NUMBER: T-22/DC**

**TEST REPORT**

ISSUED TO : Meghalayan Medical Drugs And Services Limited New colony shillong, DHS, Iaitumkhrah, Office of the mission director, NHM, Shillong, East Khasi Hills Meghalaya, 793003 Shillong - 793003 MFG. LICENCE No :	REPORT NO : EFRAC/2023/DRG/RG/03248 ISSUE DATE : 15/10/2023 CUSTOMER REFERENCE : Mail DATE : 28/09/2023 PAGE NO : 1 of 2
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**SAMPLE DETAILS**

<b>SAMPLE REGISTRATION DETAILS</b>	
Sample Name : Tab Aceclofenac 200mg	Sample Quantity Received : 80.00 Tablets
	Sample Registration Date : 09/10/2023
	Sample Submitted/Drawn by : Client
	Batch Size : N/A
	Date of Expiry : Jun-2026
Sample Receipt Date : 28/09/2023	
Registration No : EFRAC/2023/DRG/RG/03248	
Name of Manufacturer : N/A	
Batch No. : MMDSL/QC-0007	
Date of Mfg. : Jul-2023	
<b>SAMPLE ANALYSIS DETAILS</b>	
Analysis Starting Date : 12/10/2023	Analysis Completion Date : 14/10/2023

**TEST RESULT**

SL No.	TEST PARAMETER	TESTING / REF. PROCEDURE	LABEL CLAIM	UOM	ACCEPTANCE LIMIT	RESULTS
1	Identification	Indian Pharmacopoeia 2022	-	—	In the assay , the principal peak in the chromatogram obtained with the test solution should correspond to the peak in the chromatogram obtained with the reference solution.	In the assay , the principal peak in the chromatogram obtained with the test solution corresponds to the peak in the chromatogram obtained with the reference solution.
2	Disintegration test	Indian Pharmacopoeia 2022	-	Min./Sec.	NMT 15.0	Min : 04 Min 07 Sec , Max : 04 Min 32 Sec
3	Dissolution Test	Indian Pharmacopoeia 2022	200 mg/Tab	% of L.C.	NLT 75.0	Avg : 98.35 , Min : 97.98 , Max : 98.59





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Meghalayan Medical Drugs And Services Limited	ISSUE DATE :	15/10/2023
New colony shillong, DHS, Iaitumkhrah, Office of the mission	CUSTOMER REFERENCE :	Mail
director, NHM, Shillong, East Khasi Hills Meghalaya, 793003	DATE :	28/09/2023
Shillong - 793003	PAGE NO :	2 of 2
MFG. LICENCE No :		

TEST RESULT

SL No.	TEST PARAMETER	TESTING / REF. PROCEDURE	LABEL CLAIM	UOM	ACCEPTANCE LIMIT	RESULTS
4	Uniformity of the weight	Indian Pharmacopoeia 2022	-	%	(±)5.0	Avg wt : 328.06 mg , (-) Dev : 1.60 , (+) Dev : 1.81
5	Assay	Indian Pharmacopoeia 2022	200 mg/Tab	% of L.C.	180.0 - 220.0 mg/Tab i.e. 90.0 - 110.0	200.19 mg/Tab i.e. 100.10

In the opinion of the undersigned, the Sample referred to above is of Standard Quality / is not of Standard Quality as defined in the Act or the Rules made there under for the reasons given below:

Date : 15/10/2023

UOM : Unit of Measurement


REMARKS : The sample is tested as per test procedure shared by client and Opinion on Quality is drawn against the results obtained for tested parameters only.

Note : *This Report is valid only for use in Indian (Domestic) Market & is Not valid for use in any International Market.*

**-END OF THE TEST REPORT-**



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