[redacted]

TEST RESULTS

Date / Period of Performance of test 06/09/2024 to 17/09/2024.

Reference to protocol :- I.P-2022.

Description :- White elongated biconvex film coated tablet.

Identification (by HPLC) :- Complies.

Average weight :- 1512.3mg.

Uniformity of weight :- Within limit.

Dissolution for Amoxycillin (by HPLC) :- Complies(90.64% to 95.23%) Limit NLT 85%+5%

Dissolution for Clavulanic Acid (by HPLC) :- Complies(90.06% to 95.31%) Limit: NLT 80%+5%

Uniformity of Content by(HPLC) (For Clavulanic Acid) :- Within limit.

Water(by KF) :- 6.74%w/w Limit:NMT 11.0%w/w

Assay (by HPLC) :- Each film coated tablet contains:-

	Obtd./Avg.wt.	Claim	Limit Lower	Limit Upper	Method
Content of Amoxycillin Trihydrate eq. to Amoxycillin	:- 870.63mg	875.0mg	787.5mg	1050.0mg	IP.
Potassium Clavulanate diluted eq. to Clavulanic Acid	:- 124.82mg	125.0mg	112.5mg	150.0mg	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.

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