

ZEE LABORATORIES

47, Industrial Area, Gondpur Paonta Sahib Sirmour HP INDIA 173025

CERTIFICATE OF ANALYSIS (FINISHED PRODUCT)

(The Drugs & Cosmetics Act 1940 at the under form 39)

Work Order No. : 252610924

Document No. : 252612754

Ref. STS No.	348	Product Code	FG03348
Product Name	TAB. VALPROLEX CHRONO 500CR		
Generic Name	Sodium Valproate & Valproic Acid Controlled Release Tablets 500mg		
Batch No.	425-704	Date Of Sampling	18/05/2025
Batch Size	170000.000 Tab	Sample Quantity	100.000 Tab
Mfg. Date	04/2025	Exp. Date	03/2028
A. R. No.	ZL25FG02244	Date Of Analysis	18/05/2025
Date Of Release	20/05/2025		

S NO.	Tests	Specification	Results
1	Description	Off white colour, elongated, biconvex, controlled release tablet, scored on one side and plain on another side	Complies
2	Identification	Should be positive for Sodium Valproate	Complies
3	Average Weight of Tablet	730 mg ± 5.0%	736.09 mg
4	Uniformity of Weight	Average Weight of Tablet + 5.0%	Complies
5	Dissolution (1 Hours)	NLT 10% and NMT 50%	34.25 %
6	Dissolution (3 Hours)	NLT 35% and NMT 75%	45.14 %
7	Dissolution (8 Hours)	NLT 70%	88.63 %
8	Assay: Each controlled release tablet Contains Sodium Valproate I.P. 333 mg Valproic Acid I.P. 145 mg (Both together correspond to Sodium Valproate 500 mg)	NLT 90.0% and NMT 110.0% of labeled amount	98.75 %

Remarks The above sample complies/does not comply with prescribed standards of Quality with respect to above tests as per IP/BP/USP/In house specification.

Particulars	Prepared / Analysed by	Reviewed By	Approved / Authorized By
Signature:	<i>Amr</i>	<i>S</i>	
Name:	<i>Amargeet Kumar</i>	<i>Sunil Katal</i>	<i>Amr Quality Control Manager</i>
Designation:	QC Executive	Asst. QCM	QCM
Date	20/05/2025	20/05/2025	20/05/2025