

CERTIFICATE OF ANALYSIS

Report issued to	NEOLPHARMA Grupo Farmaceutico	Report No.	CMC/LAB/NPGLT2/2026
Sample Name	NeolPharma GLUTAMAX 500 MG/CAP 30 Capsules	ULR No.	NA
Manufacturer	NEOLPHARMA Grupo Farmaceutico (Mexico)	Reporting Date	25-03-2026
Active Ingredient	L-Glutamine	Sample Submitted By	Ing. Ricardo Alvarez Mendez, QA Manager (NEOLPHARMA)
Package	Bottle of 30 capsules	Exp. Date	03/2029
Batch / Lot No.	NPGLT2	Sample Qty.	30 capsules
Received Date	21-03-2026	Mfg. Lic. No.	COFEPRIS 153300201X0092
Sample Condition	Received in sealed intact condition	Ref. No.	NIL
Storage Condition	Store below 30C away from direct sunlight	Country of Origin	Mexico

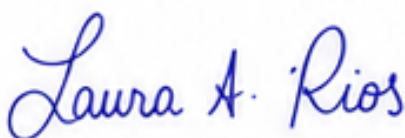
Reference to Protocol / Method	USP 2024 · CMC-LAB/SOP/NPGLT2	Analysis Started on:	22-03-2026
Analysis Completed on:	24-03-2026	Review / Release Date:	25-03-2026

RESULT OF ANALYSIS

S.No	Test Parameter	Claim	Units	Results	Specifications	Test Method
1	Description	-	-	Complies	White to off-white powder filled in capsules	Visual Evaluation.
2	Average Weight	-	mg	503.1	485.0 - 515.0 mg	USP <2091>
3	Identification	-	-	Complies	Positive for L-Glutamine	HPLC
4	Assay of L-Glutamine	500	mg/cap	508.4	475.0 - 525.0 mg/cap	HPLC
5	Uniformity of Dosage Units	-	-	Complies	AV ≤ 15.0	USP <905>
6	Dissolution	-	% / 30 min	96	Not less than 80 %	USP <711>
7	Related Substances	-	%	Complies	NMT 0.5% any individual impurity NMT 1.0% total impurities	HPLC
8	Loss on Drying	-	%	0.28	NMT 1.0 %	USP <731>
9	Bacterial Endotoxin	-	EU/cap	< 0.25	NMT 1.0 EU/cap	USP <85>

*The sample submitted by NEOLPHARMA Grupo Farmaceutico **COMPLIES** with the declared specifications and is of standard pharmaceutical quality with respect to the parameters tested.*

USP = United States Pharmacopeia · HPLC = High Performance Liquid Chromatography · KF = Karl Fischer · NMT = Not More Than
EU = Endotoxin Units · AV = Acceptance Value · COFEPRIS = Comision Federal para la Proteccion contra Riesgos Sanitarios



QFB. Laura A. Martinez Rios
Chief Analyst, Costamed Laboratory



Dr. Arturo Gonzalez Pena
Technical Director, Costamed Laboratory

**** End of Report ****
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Terms and Conditions

- This report is issued by Costamed Grupo Medico and refers only to the sample tested.
- Sample will be destroyed one year from the date of issue unless otherwise specified.
- Total liability of the laboratory is limited to the invoiced amount.
- All disputes subject to Cozumel, Quintana Roo jurisdiction, Mexico.