

CERTIFICATE OF ANALYSIS

Report issued to	NEOLPHARMA Grupo Farmaceutico	Report No.	CMC/LAB/NPCL06/2026
Sample Name	NeolPharma CLOMIA 50 MG/TAB 30 TABLET	ULR No.	NA
Manufacturer	NEOLPHARMA Grupo Farmaceutico (Mexico)	Reporting Date	14-04-2026
Active Ingredient	Clomiphene Citrate	Sample Submitted By	Ing. Ricardo Alvarez Mendez, QA Manager (NEOLPHARMA)
Package	Bottle of 30 tablets	Exp. Date	04/2029
Batch / Lot No.	NPCL06	Sample Qty.	30 tablets
Received Date	10-04-2026	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

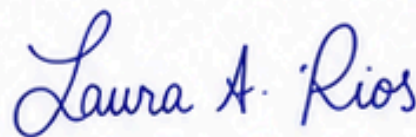
Reference to Protocol / Method	USP 2024 · CMC-LAB/SOP/NPCL06	Analysis Started on:	11-04-2026
Analysis Completed on:	13-04-2026	Review / Release Date:	14-04-2026

RESULT OF ANALYSIS

S.No	Test Parameter	Claim	Units	Results	Specifications	Test Method
1	Description	-	-	Complies	Yellow to light yellow, round, biconvex, film coated tablets	Visual Evaluation.
2	Average Weight	-	mg	198.6	190.0 - 210.0 mg	USP <2091>
3	Identification	-	-	Complies	Positive for Clomiphene Citrate	HPLC
4	Assay of Clomiphene Citrate	50	mg/tab	50.3	45.0 - 55.0 mg/tab	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	-	%	0.24	NMT 1.0 %	USP <731>
9	Bacterial Endotoxin	-	EU/tab	< 0.25	NMT 1.0 EU/tab	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.