TACROLIMUS MONOHIDRATADO (EUR. PH.)

PRODUCT CODE: 0316	CAS Nº: 109581-93-3	ANALYSIS Nº: 311/25	
MANUFACTURER BATCH:	HNTN2508501E	CERTIFICATE ID:	48.541
SUPPLIER BATCH:	----	MANUFACTURING DATE:	03/08/2025
METAPH BATCH:	0031025	RETEST DATE:	02/08/2028

ATTRIBUTES	SHOULD BE	IS
Appearance	White or almost white, crystalline powder	White crystalline powder
Identification	Complies	Complies
Appearance of solution	Clear and not more intensely coloured than ref. sol. GY7	Complies
Related substances		
Impurity A	=< 0.5 %	< 0.05 %
Unspecified impurities	=< 0.15 %	< 0.15 % (#)
Total impurities	=< 1.0 %	0.2 %
Water	1.5 - 4.0 %	2.2 %
Sulfated ash	=< 0.1 %	0.06 %
Assay	97.0 - 102.0 %	100.8 %

COMPLIES WITH

European Pharmacopoeia 11.0

REMARKS

(#) Unspecified impurities detailed underneath:
RT (13.78 min) = 0.12 %
RT (14.35 min) = 0.08 %

Product fully analyzed within the EU, in compliance with the currents regulations "EudraLex - Volume 4 - Good Manufacturing Practice (GMP) guidelines" Part-II.

Tacrolimus Monohydrate is subjected to the requirements of the ICH Q3D "Elemental Impurities" guideline and the requirements of guides EMA/CHMP/ICH/82260/2006.

All data controlled by Metapharmaceutical Industrial SL.

Absence of N-nitrosamines impurities has been ensured after a risk evaluation according to ICH Q9, ICH M7 and in accordance with guidelines EMA/428592/2019 Rev 2 and EMA/189634/2019.

Certificates of residual solvents, allergens, non-GMO and BSE-TSE, among others, are available upon request.

All methods of analysis are validated by official pharmacopoeias or are validated by internal methods of the manufacturer, which can be obtained at specific request. The above information does not exempt from the obligation to identify the product before use.

STORAGE

Keep the continers tightly closed in a fresh and dry place.

Relase date: 12/12/2025
Signature: Albert Sánchez López (QP)
Conclusion: Complies

Manufacturer: 40000447

NORTH CHINA PHARMACEUTICAL HUASHENG Co., Ltd.
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Original certificate available upon request

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