

Stabilimento di Garbagnate
Via Milano, 186
20024 Garbagnate Milanese (MI)

PEC: olon@pec.olonspa.it
Tel. 02 9944221
Fax 0382.812006

Olon S.p.A.
Sede Legale
Strada Rivoltana km 6/7
Rodano (MI)



Certificate of analysis

Product **MINOXIDIL-USP/EP current edition** Code N° **2610290** Batch N° **1610000856**

Manufacturing date **APR 2016** Retest date (where applicable) **MAR 2021**

Analysis Number **Q1PF042702** Bulletin **4P1642246**

Test	References	Result	M.U.	Limits
Description		Off-white crystalline powder		White to off-white crystalline powder
IR		Conforms to standard		Conforms to standard
UV and VISIBLE absorption		Conforms		Conforms
Loss on drying		0,1	%	≤ 0,5
Residue on ignition		0,1	% db	≤ 0,1
Heavy metals		< 20	ppm	≤ 20
Appearance of solution		Not more coloured than Y6		Not more coloured than Y6
Appearance of solution		Clear	—	Clear
Assay (HCLO4)		100,2	%	99,0 + 101,0
HPLC: Assay		99,4	% dried	97,0 + 103,0
		< 0,05	% dried	≤ 0,15
2,4-Diamino-6-ChloroPyrimidine (Impurity B)		< 0,05	%	≤ 0,2
2,4-Diamino-6-Piperidinpyrimidine (Impurity E or deoxyminoxidil)		< 0,05	%	≤ 0,10
Largest unknown impurity		< 0,05	%	≤ 0,3
Total impurities by EP		< 0,05	%	≤ 0,3
Total impurities by USP		< 0,05	%	≤ 0,3
OVI (USP): Not used		Not tested		Not tested
ICH Class 1 solvents : Not used		Not tested	—	Not tested
ICH Class 2 solvents : Not used		Not tested	—	Not tested
ICH Class 3 solvents		Conforms	—	Conforms
2-Propanol		708	ppm	≤ 2000
Acetone		39	ppm	≤ 1000
Packaging and storage		Preserve well-closed container	—	Preserve well-closed container

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Analysis Date

05/11/2016

QC Manager

11 MAR 2016

Reviewed by Qualified Person

Notes:

This batch has been manufactured, packaged and tested in accordance with EU GMP Guideline Volume 4 Part II (ICH Q7A).
Manufacturing site of the Active Pharmaceutical Ingredient: Stabilimento di Garbagnate