

FAGRON a.s.

Inspectional laboratory no. 536 Holická 1098/31m, 779 00 Olomouc tel. 585222590, fax 585226521

email: orj@fagron.cz

CERTIFICATE OF ANALYSIS

Product name:

LANOLINA

Inspection No:

H05835/0620/536

Customer:

Fagron PL

Batch No:

077351

Batch size:

200 x 1 kg

Expiry date:

31.03.2022

Analysed according to: Ph. Eur. 04/2012:0134

Tests	Requirements:	Results:
Appearance	Yellow, unctuous substance. When melted, it is a clear or	conform
	almost clear, yellow liquid. A solution in light petroleum	
	in opalescent.	
Solubility	Practically insoluble in water, slightly soluble in boiling	conform
	anhydrous ethanol.	
Odour	Characteristic odour.	conform
Identification		
Identification A	green	conform
Identification B	red colour, fluorescence	conform
Tests		
Water-soluble acid or alk. subs.	complies with Ph. Eur.	conform
Water-absorption capacity	≥ 20 [ml]	35 ml
Acid value	≤ 1,0	0,70
Peroxide value	≤ 20	7,4
Saponification value	90 - 105	101
Water-soluble oxidisable subs.	complies with Ph. Eur.	conform
Paraffins	≤1,0 [%]	0,20 %
Pesticides residues	complies with Ph. Eur.	conform
Chlorides	≤150 [ppm]	conform
Loss on drying	≤ 0,5 [%]	0,01 %
Sulfated ash	≤ 0,15 [%]	0,065 %



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Tests	Requirements:	Results:
Microbiology	complies with Ph. Eur.	conform
TAMC	≤ 10³ [CFU/g]	conform
TYMC	≤ 10 ² [CFU/g]	conform
TSE/BSE	No contamination with TSE/BSE-risk materials.	conform
Drop point	38 – 44 [°C]	43 °C

The product conforms to Ph. Eur. 04/2012:0134

Conclusion: I certify that the above-mentioned batch of pharmaceutical raw material has been manufactured and tested in accordance with the requirements of Marketing Authorization (License) and requirements of Good Manufacturing Practice.

Approved by:

Ivana Urbánková, QP

Date and signature:

9.4.2020 lld.