

## Certificate of Analysis

**Product name:** **Erythromycinum**  
**Batch number / Weight:** **23D17-B04-236258 / 100 G**  
**Analysed according to:** **Ph.Eur.11.3**  
**Number of analysis / Inspection Code** **23D17-B04 / INS-23-3977**  
**Reference Code / No.:** **V00596 / EBS/23-004**

Tests	Requirement	Result	Unit	Standard remark
Appearance	White or slightly yellow powder, or colourless	Conform		
Appearance	or slightly yellow crystals, slightly hygroscopic			
Identification A	Conform	Conform		
Related substances	Conform	Conform		
Impurity A	$\leq 2,0$	0,37	%	
Impurity B	$\leq 2,0$	0,66	%	
Impurity C	$\leq 3,0$	0,66	%	
Impurity D	$\leq 1,0$	0,42	%	
Impurity E	$\leq 1,0$	$< 0,2$	%	
Impurity F	$\leq 1,0$	$< 0,2$	%	
Impurity H	$\leq 1,0$	0,19	%	
Impurity L	$\leq 0,4$	$< 0,2$	%	
Any other impurity	$\leq 0,4$	$< 0,2$	%	
Total impurities	$\leq 7,0$	2,31	%	
Thiocyanate	$\leq 0,3$	0,02	%	
Water (Karl Fischer)	$\leq 6,5$	1,87	%	
Sulphated ash	$\leq 0,2$	0,03	%	
Metallic residues	CHMP/ICH/353369/2013	Conform		Data producer
Residual solvents	CPMP/ICH/82 260/06	Conform		Data producer
Assay Erythromycine (A+B+C)	93,0 - 102,0	97,21	%m/m	
Erythromycine B	$\leq 5,0$	0,13	%	
Erythromycine C	$\leq 5,0$	0,52	%	
TSE/BSE-statement:	No contamination with TSE/BSE-risk materials	Conform		Data producer

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Fagron Services Northern Europe Sp. z o.o.  
 ul. Armii Krajowej 3  
 32-540 Trzebinia, Poland  
 e-mail: fsne@fagron.pl

fagron.pl

Fagron sp. z o.o.  
 ul. Pasternik 26, 31-354 Kraków, Poland  
 tel.: +48 12 3343 512  
 e-mail: biuro@fagron.pl

All the manufacturing stages of this batch have been carried out in full compliance with the GMP requirements of the EU and with the national legal requirements.

Analysis performed by the authorized lab Fagron PL lab

Release:  
Dominika Soltysik  
Qualified Person

14-01-2024

Expiration: 31-12-2025

Conclusion: APPROVED

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Fagron Services Northern Europe Sp. z o.o.  
ul. Armii Krajowej 3  
32-540 Trzebinia, Poland  
e-mail:fsne@fagron.pl