

## Certificate of Analysis

**Product name:** **Amiloridi hydrochloridum dihyd**

**Batch number / Weight:** **23E30-B10-233822 / 5 G**

**Analysed according to:** **Ph.Eur.11.2**

**Number of analysis / Inspection Code** **23E30-B10 / INS-23-4744**

**Reference Code / No.:** **V01535 / 2001012303002**

Tests	Requirement	Result	Unit	Standard remark
Appearance	pale yellow or greenish-yellow powder	Conform		
Identification A	Conform	Conform		
Identification C	Conform	Conform		
Identification D	11,0 - 13,0 %	Conform		
Free acid	Conform	0,064	ml	
Related substances	acc. to Ph. Eur.	Conform		
Impurity C	<= 0,2	<0,05	%	
Unspecified impurities	<=0,10	0,063	%	
Total impurities	<= 0,4	0,06	%	
Water (Karl Fischer)	11,0 - 13,0	11,72	%	
Sulphated ash	<=0,1	0,03	%	
Metallic residues	CHMP/ICH/353369/2013	Conform		
Residual solvents	CPMP/ICH/82 260/06	Conform		
Assay	98,0 - 101,0	99,58	%m/m	
TSE/BSE-statement:	No contamination with TSE/BSE-risk materials	Conform		

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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

21-11-2023

Expiration: 29-02-2028

Conclusion: APPROVED

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