

CERTIFICATE OF ANALYSIS

Product: **CANIVERM comprimate á 0,7 g**

Batch No.: **A037-01** 0.7 g

Expiry: **06/2026**

Date of production: **10.7.2024**

Results of analysis:

Tests	Release limits	Shelf-life limits	Results
Appearance	Oval, yellowish, biconvex tablets with small spots and a dividing line on both sides.		complies
Content of original package	2 x 1 tbl., 6 x 1 tbl., 100 x 1 tbl.		complies
Microbiological quality			
- TAMC	10 ⁴ CFU/g		150 CFU/g
- TYMC	10 ² CFU/g		54 CFU/g
- Bile tolerant gram-negative bacteria	Max. 10 ² CFU v 1g		<10 CFU v 1g
- Absence of Salmonella	Absence in 10 g		absent
- Absence of Escherichia coli	Absence in 1 g		absent
Absence of Staphylococcus aureus	Absence in 1 g		absent
Content of fenbendazole	142.5 – 157.5 mg/0.7 g		153.01 mg/0.7 g
Fenbendazole identity	Must comply.		complies
Content of pyrantel embonate	136.8 – 151.2 mg/0.7 g		142.76 mg/0.7 g
Pyrantel identity	Must comply.		complies
Content of praziquantel	47.5 – 52.5 mg/0.7 g		51.50 mg/0.7 g
Praziquantel identity	Must comply.		complies
Weight uniformity of tablets	0.665 – 0.735 g		0.697 g
Impurities of fenbendazole			
- content of oxfendazole	Max. 1.0 %		0.1 %
- any individual unidentified impurity	Max. 1.0 %		0.1 %
- sum of impurities	Max. 3.0 %		0.1 %
Impurities of pyrantel embonate			
- any individual unidentified impurity	Max. 1.0 %		0.0 %
- sum of impurities	Max. 3.0 %		0.0 %
Impurities of praziquantel			
- any individual unidentified impurity	Max. 1.0 %		0.0 %
- sum of impurities	Max. 3.0 %		0.0 %
Determination of tablet strength	100-200 N		138 N
Tablet disintegration	Max. 15 min.		6 min.
Determination of abrasion of uncoated tablets	Max.1.0%		0.10 %

I certify that this product has been tested in the laboratories of Bioveta a.s. and has been found to meet specification requirements. Preparation is suitable with control regulations of the valid PN No. 316,V003.



09/09/2024
Mgr. Hana Nezvalová
HEAD of QC, QP