



CERTIFICATE OF ANALYSIS

Product: **SELIVERM 30 mg, soluție spot-on pentru câini S**

Batch No.: **0324A01**

Expiry: **10/2026**

Date of production: **25.11.2024**

Results of analysis:

Tests	Release limits	Shelf-life limits	Results
Appearance	A clear, colourless to yellow or yellow-green solution. The solution is clear like water R. The yellow colour of the solution is not more intense than the colour of the reference solution Y1. The green-yellow colour of the solution is not more intense than the colour of the reference solution GY1.		complies
Microbiological quality:			
TAMC	< 10 ² (CFU/g or CFU/ml)		<10 CFU/ml
TYMC	< 10 ¹ (CFU/g or CFU/ml)		<10 CFU/ml
<i>Staphylococcus aureus</i>	absent (in 1 g or 1 ml)		absent
<i>Pseudomonas aeruginosa</i>	absent (in 1 g or 1 ml)		absent
Water content	Max. 2.0 %		0.37 %
Density	0.8370 – 0.8600 g/cm ³		0.8500 g/cm ³
Refractive index	1.37 – 1.43		1.404
Selamectin identity test	RT and UV spectrum match with standard		complies
Selamectin assay	28.50 – 31.50 mg/tube	27.90 – 32.10 mg/tube	29.565 mg/tube
Butylhydroxytoluene identity test	Rt and UV spectrum match with standard		complies
BHT assay	0.72 – 0.88 mg/ml	Max. 0.88 mg/ml	0.801 mg/ml
Purity test	Impurity A – max. 2.0 %		0.2196 %
	Impurity B – max. 2.0 %		0.2856 %
	Impurity C – max. 1.5 %		0.1751 %
	Impurity D – max. 1.5 %		0.2853 %
	Any individual undefined impurity - max. 1.0 %		<1.0 %
	Sum of impurities – max. 4.0 %		2.86 %
Uniformity of dosage units (Weight variability)	Must comply.		complies
Average extractable volume	0.2250 - 0.2750 ml		0.2430 ml

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. 556a,V001.



Nezvalová
.....10/03/2025
Mgr. Hana Nezvalová
HEAD of QC, QP