

Certificate of Analysis

Certificate No.: 39965

Product: Boflox Flavour 80mg 72er RO
Article number: 86090 Dosage form: Tablet
Batch: 12304203 Split: 0 Package size: 72 tablets
Quantity: 312 Units Marketing authorisation holder: Livisto Int'l
Manufacturing date: 23.02.2021 Registration number: ~~15999/2~~ 170033
Expiry date: 31.01.2026 Distributor / Importer: RO - BistriVet Rumänien

6/280521

Parameter	Specification	Result
Appearance	Beige round tablets, brown speckled, with breaking notch on both sides	Beige round tablets, brown speckled, with breaking notch on both sides
Length	20,2 ... 21,1 mm	20,7 mm
Width	8,9 ... 9,3 mm	9,1 mm
Height	6,6 ... 7,4 mm	7,0 mm
Average mass	1064 ... 1176 mg	1109 mg
Uniformity of dosage by mass variation	Must correspond to Ph. Eur. 2.9.40	Corresponds to
Acceptance value (AV)	Max. 15,0	2,0
Hardness	100 ... 200 N	152 N
Disintegration with disc	Max. 15,0 min	2,0 min
Acceptance level dissolution	Must correspond to Ph. Eur. 2.9.3	Corresponds to
Dissolution	After 30 min.: 80% Q	102,6 %
Identity: marbofloxacin	Retention time and UV spectrum of standard and sample must correspond	Corresponds to
Assay: marbofloxacin	76,0 ... 84,0	77,7 mg/Tabl.
Impurity D	Max. 1,0 %	< 0,1 %
Impurity F	Max. 1,0 %	< 0,1 %
Unknown impurity, single	Max. 1,0 %	< 0,1 %
Total impurities	Max. 1,5 %	< 0,1 %
Microbiological purity: TAMC	Max. 1000	< 1000 CFU/g
Microbiological purity: TYMC	Max. 100	< 100 CFU/g
Microbiological purity: Escherichia coli	Absent/g	Absent/g

Template: 1340

The finished product has been tested in accordance with the testing instruction and is of the required quality.

26. MAI 2021



Date, Signature Head of Quality Control



Batch certificate

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aniMedica GmbH
a LIVISTO company

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I certify that the above information is authentic and correct. This batch has been manufactured, including packaging and quality control, in full compliance with the GMP requirements of the local regulatory authority and with the information in the marketing authorisation as laid down in the manufacturing and quality control instructions. The batch processing, packaging and analytical documentation has been reviewed and is in full compliance with GMP guidelines. All deviations were evaluated before batch release.

This batch has the required quality and

- ☐ is released for delivery
- ☒ is released for the market

Remarks:

/ / / /

280521 
Date, Signature Qualified Person

- ☒ Dr. Hans-Jörg Lange ☐ Gudrun Prußeit ☐ Dr. Jürgen Schlüter ☐ Dr. Bernd Manfred Müller ☐