



aniMedica Herstellungs GmbH
a LIVISTO company

Certificate of Analysis

Certificate No.: 48258

Product:	Dexafast 2mg/ml inj 100ml EP RO	Dosage form:	Solution for injection
Article number:	00100357	Package size:	100ml
Batch:	22020003 Split: 0	Marketing authorisation holder:	Livisto Int'l
Quantity:	3.808 Units	Registration number:	190002
Manufacturing date:	01.03.2022	Distributor / Importer:	RO - BistriVet Rumänien
Expiry date:	28.02.2026		

Parameter	Specification	Result
Appearance	Clear, colourless solution essentially free of visible particles	Clear, colourless solution essentially free of visible particles
Batch packaging record	batch packaging record complies	Corresponds to
pH	7,0 ... 8,0	7,4
Relative density	1,003 ... 1,013	1,010
Extractable volume	Min. 100,0 %	102,0 %
Identity: dexamethasone sodium phosphate	Retention times and UV spectrum of standard and sample must correspond	Corresponds to
Identity: benzyl alcohol	Retention times and UV spectrum of standard and sample must correspond	Corresponds to
Assay: dexamethasone sodium phosphate	2,50 ... 2,76 mg/ml	2,71 mg/ml
Assay: benzyl alcohol	14,0 ... 17,2 mg/ml	15,7 mg/ml
Impurity: dexamethasone	Max. 1,0 %	< 0,1 %
Any unspecified impurity	Max. 1,0 %	0,1 %
Total impurities	Max. 5,0 %	0,5 %
Sterility	Sterile	Corresponds to

Template: 2116

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

29. MARZ. 2022

Date, Signature Head of Quality Control

Amtsgericht Coesfeld
HRB-Nr. 10339
Geschäftsführer:
Thomas Höltker

Im Südfeld 9
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www.livisto.com

Along with you


LIVISTOaniMedica Herstellungs GmbH
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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:

28.2.22
Date, Signature Qualified Person

- ☒ Dr. Hans-Jörg Lange ☐ Gudrun Prüßert ☐ Dr. Jürgen Schlüter ☐ Dr. Bernd Manfred Müller
☐ Ulrich Kontny

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