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MASTIJET FORT

Authorised

- Prednisolone
- Bacitracin
- Neomycin
- Tetracycline hydrochloride

Product identification

Medicine name:

MASTIJET FORT

Active substance:

Prednisolone

Bacitracin

Neomycin

Tetracycline hydrochloride

Target species:

Cattle

Route of administration:

Intramammary use

Product details

Active substance and strength:

Prednisolone

10.00 milligram(s) / 1.00 Syringe

Bacitracin

2000.00 international unit(s) / 1.00 Syringe

Neomycin

250.00 milligram(s) / 1.00 Syringe

Tetracycline hydrochloride

200.00 milligram(s) / 1.00 Syringe

Pharmaceutical form:

Intramammary suspension

Withdrawal period by route of administration:

Intramammary use:

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Cattle

- Meat and offal. 14 day

- Milk. no withdrawal period
Мляко: 8 издоявания.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ51RV01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Bulgaria

Available in:

Bulgaria

Package description:

Available only in Bulgarian

Available only in Bulgarian

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Intervet International B.V.

Marketing authorisation date:

10/04/2013

Manufacturing sites for batch release:

Intos B.V.

Responsible authority:

Bulgarian Food Safety Authority

Authorisation number:

0022-2008

Date of authorisation status change:

10/04/2013

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet and Labelling

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