

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

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