

MASTIJET FORT, 200 mg + 250 mg + 2000 i.j. + 10 mg, intramamarna suspenzija za krave u laktaciji

Authorised

- Prednisolone
- Bacitracin
- Neomycin
- Tetracycline hydrochloride

Product identification

Medicine name:

MASTIJET FORT, 200 mg + 250 mg + 2000 i.j. + 10 mg, intramamarna suspenzija za krave u laktaciji

Active substance:

Prednisolone

Bacitracin

Neomycin

Tetracycline hydrochloride

Target species:

Cattle (lactating cow)

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:

-

Cattle (lactating cow)

- Meat and offal. 14 day

- Milk. 5 day

Mlijeko nije prikladno za hranu tijekom liječenja i još 10 mužnji (120 sati) nakon posljednje aplikacije pripravka tj. može se koristiti prilikom 11. mužnje, uz uvjet da se krave muzu 2× na dan.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ51RV01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Croatia

Available in:

Croatia

Package description:

Available only in [Croatian](#)

Available only in [Croatian](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Intervet International B.V. Subsidiary In The Republic Of Croatia

Marketing authorisation date:

26/09/2014

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:

UP/I-322-05/22-01/701

Date of authorisation status change:

27/12/2023

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

Documents

Summary of Product Characteristics

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