Mastijet Forte suspensija ievadīšanai tesmenī laktējošām govīm

Authorised

- Chlortetracycline hydrochloride
- Neomycin
- Bacitracin
- Prednisolone

Product identification

Medicine name:
Mastijet Forte suspensija ievadīšanai tesmenī laktējošām govīm

Active substance:
Chlortetracycline hydrochloride

Neomycin

Bacitracin

Prednisolone

Target species:
Cattle (lactating cow)

Route of administration:
Intramammary use

Product details

Active substance and strength:

Chlortetracycline hydrochloride
200.00 milligram(s) / 1.00 Syringe

Neomycin
250.00 milligram(s) / 1.00 Syringe

Bacitracin
2000.00 international unit(s) / 1.00 Syringe

Prednisolone
10.00 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. 96 hour Pienam: 96 stundas (8 slaukšanas reizes).

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ51RV01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Latvia

Available in:

Latvia

Package description:
Available only in Latvian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
19/02/1996

Manufacturing sites for batch release:
Intervet International B.V.

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/NRP/96/0344

Date of authorisation status change:
19/02/1996

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

Documents

Combined File of all Documents

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