Mastitar Retard Vet. intramammær salve 500.000 IE+300 mg

Authorised

- Neomycin
- Benzylpenicillin procaine

Product identification

Medicine name:
Mastitar Retard Vet. 500.000 IE +300 mg intramammær salve
Mastitar Retard Vet. intramammær salve 500.000 IE+300 mg

Active substance:
Neomycin
Benzylpenicillin procaine

Target species:
Cattle

Route of administration:
Intramammary use

Product details

Active substance and strength:
Neomycin
300.00 milligram(s) / 1.00 Syringe

Benzylpenicillin procaine
500.00 milligram(s) / 1.00 Syringe

Pharmaceutical form:
Intramammary ointment

Withdrawal period by route of administration:

Intramammary use:

- **Cattle**

  - Meat and offal. 10 day efter kælvning.

  - Meat and offal. 10 day efter kælvning.

  - Milk. 4 day

Ved behandling mere end 30 døgn før kælvning er tilbageholdelsestiden for mælk: 4 døgn efter kælvning.

  - Milk. 4 day

Ved behandling mere end 30 døgn før kælvning er tilbageholdelsestiden for mælk: 4 døgn efter kælvning.

  - Milk. 30 day

Ved behandling mindre end 30 døgn før kælvning er tilbageholdelsestiden for mælk: 30 døgn efter kælvning.

  - Milk. 30 day

Ved behandling mindre end 30 døgn før kælvning er tilbageholdelsestiden for mælk: 30 døgn efter kælvning.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51RC23

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:

Denmark

Package description:
Available only in Danish
Available only in Danish
Available only in Danish
Available only in Danish
Available only in Danish
Available only in Danish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Virbac

Marketing authorisation date:
26/10/1982

Manufacturing sites for batch release:
VIRBAC
INTERVET INTERNATIONAL B.V.

Responsible authority:
Danish Medicines Agency

Authorisation number:
09168

Date of authorisation status change:
26/10/1982

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

Documents

Summary of Product Characteristics

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