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Benestermycin, intramammaarsuspensioon veistele

Authorised

- Framycetin sulfate
- Benethamine penicillin
- Penethamate hydriodide

Product identification

Medicine name:
Benestermycin, intramammaarsuspensioon veistele

Active substance:
Framycetin sulfate

Benethamine penicillin

Penethamate hydriodide

Target species:
Cattle (dry cow)

Route of administration:
Intramammary use

Product details

Active substance and strength:

Framycetin sulfate
100.00 milligram(s) / 1.00 Syringe

Benethamine penicillin
280.00 milligram(s) / 1.00 Syringe

Penethamate hydriodide
100.00 milligram(s) / 1.00 Syringe

Pharmaceutical form:
Intramammary suspension

Withdrawal period by route of administration:

Intramammary use:

- **Cattle (dry cow)**
  - Meat and offal. 8 day
  - Milk. 36 hour

Piimale: 36 tundi pärast poegimist, kui ravimit manustati 35 päeva või rohkem enne poegimist. 37 päeva pärast ravi, kui ravimit manustati vähem kui 35 päeva enne poegimist.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51RC25

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

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Authorisation status:
Valid

Authorised in:
Estonia

Available in:
Estonia

Package description:
Available only in Estonian

Available only in Estonian

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:
Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:
27/04/2008

Manufacturing sites for batch release:
Haupt Pharma Latina S.r.l.
Lohmann Pharma Herstellung GmbH

Responsible authority:
State Agency Of Medicines

Authorisation number:
1513

Date of authorisation status change:
27/04/2008

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

Documents

Summary of Product Characteristics

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