Nafpenzal DC, intramammaarsuspensioon veistele

Authorised

- Nafcillin
- Dihydrostreptomycin
- Benzylpenicillin procaine

Product identification

Medicine name:
Nafpenzal DC, intramammaarsuspensioon veistele

Active substance:
Nafcillin

Dihydrostreptomycin

Benzylpenicillin procaine

Target species:
Cattle (dry cow)

Route of administration:
Intramammary use

Product details

Active substance and strength:

Nafcillin
100.00 milligram(s) / 1.00 Syringe

Dihydrostreptomycin
100.00 milligram(s) / 1.00 Syringe

Benzylpenicillin procaine
300.00 milligram(s) / 1.00 Syringe

Pharmaceutical form:
Intramammary suspension

Withdrawal period by route of administration:

Intramammary use:

- **Cattle (dry cow)**
  - Meat and offal. 14 day
  - Milk. 36 hour

Piimale: 36 tundi, kui kinnisperiood kestab rohkem kui 6 nädalat. 6 nädalat + 36 h, kui kinnisperiood on lühem kui 6 nädalat.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51RC23

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

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Authorisation status:
Valid

Authorised 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:
Intervet International B.V.

Marketing authorisation date:
5/02/2004

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

Responsible authority:
State Agency Of Medicines

Authorisation number:
1189

Date of authorisation status change:
5/02/2004

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

Documents

Summary of Product Characteristics

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