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NAFPENZAL DC

Not authorised

- Nafcillin
- Dihydrostreptomycin
- Benzylpenicillin procaine

Product identification

Medicine name:
NAFPENZAL DC

Active substance:
Nafcillin
Dihydrostreptomycin
Benzylpenicillin procaine

Target species:
Cattle (cow)
Sheep
Goat (adult female)

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 (cow)**
  - Meat and offal. 14 day
  - Milk. 2 day
- **Sheep**
  - Meat and offal. 21 day
  - Milk. 6 day
- **Goat (adult female)**
  - Meat and offal. 21 day
  - Milk. 6 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51RC23

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

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Authorisation status:
Surrendered

Authorised in:
Romania

Package description:
Available only in Romanian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Directive No 2001/82/EC

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
19/08/2010

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

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:
160341

Date of authorisation status change:
28/03/2024

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

Documents

Summary of Product Characteristics

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