Nafpenzal DC 100 mg suspension intramammaire

Authorised

- Nafcillin sodium
- Dihydrostreptomycin sulfate
- Benzylpenicillin procaine

Product identification

Medicine name:
Nafpenzal DC 100 mg suspension intramammaire

Active substance:
Nafcillin sodium

Dihydrostreptomycin sulfate

Benzylpenicillin procaine

Target species:
Cattle (dairy cattle)

Route of administration:
Intramammary use

Product details

Active substance and strength:
Nafcillin sodium
100.00 milligram(s) / 1.00 Syringe

Dihydrostreptomycin sulfate

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 (dairy cattle)**
  - Meat and offal. 5 day
  - Milk. no withdrawal period

Treatment to calving interval ≥ 46 days: 48 hours; Treatment to calving interval < 46 days: 46 days after treatment

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51RC23

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

Authorisation status:
Valid

Authorised in:
Luxembourg

Available in:
Luxembourg

Package description:
20 Intramammary syringes with 3g of Intramammary suspension

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:
1/08/1972

Manufacturing sites for batch release:
Intervet International GmbH

Responsible authority:
Ministry Of Health And Social Security

Authorisation number:
V 817/02/11/0746

Date of authorisation status change:
1/01/2000

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

Documents

Summary of Product Characteristics

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