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Nafpenzal DC Intramammary Suspension

Not authorised

- Benzylpenicillin procaine
- Dihydrostreptomycin sulfate
- Nafcillin sodium

Product identification

Medicine name:
Nafpenzal DC Intramammary Suspension

Active substance:
Benzylpenicillin procaine

Dihydrostreptomycin sulfate

Nafcillin sodium

Target species:
Cattle

Route of administration:
Intramammary use

Product details

Active substance and strength:
Benzylpenicillin procaine

300.00 milligram(s) / 1.00 Syringe

Dihydrostreptomycin sulfate
116.81 milligram(s) / 1.00 Syringe

Nafcillin sodium
105.31 milligram(s) / 1.00 Syringe

Pharmaceutical form:
Intramammary suspension

Withdrawal period by route of administration:
Intramammary use:

- **Cattle**
  - Meat and offal. 14 day
  - Milk. 48 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51RC23

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

Authorisation status:
Surrendered

Authorised in:
Ireland

Package description:
Each 3 g syringe is made of low density polyethylene. The syringes are packed in a sachet. The sachet is packed in a carton. Cleaning towels packed in a sachet are also included in the carton. Packs of 20 syringes and 20 cleaning towels.

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 (Ireland) Limited

Marketing authorisation date:
1/10/1987

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

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10996/074/001

Date of authorisation status change:
1/09/2023

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

Documents

Summary of Product Characteristics