Nafpenzal dc, suspensão intramamária para bovinos e ovinos

Not authorised

- Dihydrostreptomycin
- Benzylpenicillin procaine
- Nafcillin

Product identification

Medicine name:
Nafpenzal dc, suspensão intramamária para bovinos e ovinos

Active substance:
Dihydrostreptomycin

Benzylpenicillin procaine

Nafcillin

Target species:
Cattle (dairy cow at drying-off)
Sheep (dry ewe)

Route of administration:
Intramammary use

Product details

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

Benzylpenicillin procaine
300.00 milligram(s) / 1.00 Syringe

Nafcillin
100.00 milligram(s) / 1.00 Syringe

Pharmaceutical form:
Intramammary suspension

Withdrawal period by route of administration:
Intramammary use:

- **Cattle (dairy cow at drying-off)**
  - Meat and offal. 16 day

Os animais não devem ser abatidos para consumo humano durante o tratamento

  - Milk. 42 day

Intervalo no tratamento-parto > 42 dias = 48 horas após o parto. Intervalo no tratamento-parto ≤ 42 dias = 44 dias após o tratamento.

- **Sheep (dry ewe)**
  - Meat and offal. 28 day

Os animais não devem ser abatidos para consumo humano durante o tratamento

  - Milk. 3 month

Intervalo no tratamento-parto ≥ 3 meses = 6 dias após o parto. Intervalo no tratamento-parto < 3 meses = 14 dias após o parto.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51RC23

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

Authorisation status:
Revoked

Authorised in:
Portugal

Package description:
Available only in Portuguese
Available only in Portuguese

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:
MSD Animal Health Lda.

Marketing authorisation date:
10/09/1990

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

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
1248/01/19NFVPT

Date of authorisation status change:
9/02/2024

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

Documents

Combined File of all Documents

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