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NAFPENZAL T POMMADE INTRAMAMMAIRE POUR VACHES BREBIS ET CHEVRES

Authorised

- Nafcillin
- Dihydrostreptomycin
- Benzylpenicillin (procaine) monohydrate

Product identification

Medicine name:
NAFPENZAL T POMMADE INTRAMAMMAIRE POUR VACHES BREBIS ET CHEVRES

Active substance:
Nafcillin
Dihydrostreptomycin
Benzylpenicillin (procaine) monohydrate

Target species:
Cattle (cow)
Sheep (ewe)
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) monohydrate
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. 48 day

47,5 jours après le traitement si la période de tarissement est inférieure à 46 jours.

  - Milk. 36 hour

36 heures après le vêlage si la période de tarissement est égale ou supérieure à 46 jours.

- **Sheep (ewe)**
  - Meat and offal. 28 day
  - Milk. 14 day

14 jours après agnelage pour une période de tarissement inférieure à 3 mois.

  - Milk. 6 day

6 jours après agnelage quand la période de tarissement est supérieure ou égale à 3 mois.

- **Goat (adult female)**
  - Meat and offal. 28 day
  - Milk. 14 day

14 jours après la mise bas quand la période de tarissement est inférieure à 40 jours.

  - Milk. 10 day

10 jours après la mise bas quand la période de tarissement est supérieure à 40 jours.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51RC23

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

Authorisation status:
Valid

Authorised in:
France

Available in:
France

Package description:
Available only in French
Available only in French
Available only in French
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Intervet

Marketing authorisation date:
2/02/1990

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

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/5170868 1/1990

Date of authorisation status change:
2/02/2010

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

Documents

Summary of Product Characteristics

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Package Leaflet and Labelling

This document does not exist in this language (English). You can find it in another language below.