NAFPENZAL DC

Authorised

- Benzylpenicillin procaine
- Nafcillin sodium

Product identification

Medicine name:
NAFPENZAL DC

Active substance:
Benzylpenicillin procaine

Nafcillin sodium

Target species:
Goat
Sheep
Cattle (dairy cow)

Route of administration:
Intramammary use

Product details

Active substance and strength:
Benzylpenicillin procaine
300.00 milligram(s) / 1.00 Syringe

Nafcillin sodium
100.00 milligram(s) / 1.00 Syringe

Pharmaceutical form:
Intramammary suspension

Withdrawal period by route of administration:

Intramammary use:

- **Goat**
  - Meat and offal. 35 day
  - Milk. 9 day
- **Sheep**
  - Meat and offal. 35 day
  - Milk. 90 day
- **Cattle (dairy cow)**
  - Meat and offal. 48 hour
  - Meat and offal. 16 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51GA90
QJ51RC
QJ51RC22

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

Authorisation status:
Valid

Authorised in:
Cyprus

Package description:
Available only in Greek
Available only in Greek

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
30/04/2004

Manufacturing sites for batch release:
INTERVET INTERNATIONAL B.V.

Responsible authority:
Ministry Of Agriculture Rural Development And Environment

Authorisation number:
CY00072V

Date of authorisation status change:
9/06/2017

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

Documents

Summary of Product Characteristics

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