

NAFPENZAL DC

Authorised

- Benzylpenicillin procaine
- Nafcillin
- Dihydrostreptomycin

Product identification

Medicine name:

NAFPENZAL DC

Active substance:

Benzylpenicillin procaine

Nafcillin

Dihydrostreptomycin

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

100.00 milligram(s) / 1.00 Syringe

Dihydrostreptomycin

100.00 milligram(s) / 1.00 Syringe

Pharmaceutical form:

Intramammary suspension

Withdrawal period by route of administration:

Intramammary use:

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Goat

- Meat and offal. 35 day

- Milk. 9 day

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Sheep

- Meat and offal. 35 day

- Milk. 90 day

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Cattle (dairy cow)

- Meat and offal. 48 hour

- Meat and offal. 16 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ51RC

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:

Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:

CY00072V

Date of authorisation status change:

10/06/2017

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.