

Nafpenzal DC 100 mg suspension intramammaire

Authorised

- Nafcillin sodium
- Dihydrostreptomycin sulfate
- Benzylpenicillin procaine

Product identification

Medicine name:

Nafpenzal DC 100 mg suspension intramammaire

Active substance:

Nafcillin sodium

Dihydrostreptomycin sulfate

Benzylpenicillin procaine

Target species:

Cattle (dairy cattle)

Route of administration:

Intramammary use

Product details

Active substance and strength:

Nafcillin sodium

100.00 milligram(s) / 1.00 Syringe

Dihydrostreptomycin sulfate

100.00 milligram(s) / 1.00 Syringe

Benzylpenicillin procaine

300.00 milligram(s) / 1.00 Syringe

Pharmaceutical form:

Intramammary suspension

Withdrawal period by route of administration:

Intramammary use:

-

Cattle (dairy cattle)

- Meat and offal. 5 day

- Milk. no withdrawal period

Treatment to calving interval \geq 46 days: 48 hours; Treatment to calving interval $<$ 46 days: 46 days after treatment

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ51RC23

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Luxembourg

Available in:

Luxembourg

Package description:

20 Intramammary syringes with 3g of Intramammary suspension

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 International B.V.

Marketing authorisation date:

1/08/1972

Manufacturing sites for batch release:

Intervet International GmbH

Responsible authority:

Ministry Of Health And Social Security

Authorisation number:

V 817/02/11/0746

Date of authorisation status change:

1/01/2000

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.