

NAFPENZAL DC

(300+100+100)MG/3G

ΕΝΔΟΜΑΣΤΙΚΟ ΕΝΑΙΩΡΗΜΑ

Authorised

- Dihydrostreptomycin sulfate
- Nafcillin sodium monohydrate
- Benzylpenicillin procaine

Product identification

Medicine name:

NAFPENZAL DC (300+100+100)MG/3G ΕΝΔΟΜΑΣΤΙΚΟ ΕΝΑΙΩΡΗΜΑ

Active substance:

Dihydrostreptomycin sulfate

Nafcillin sodium monohydrate

Benzylpenicillin procaine

Target species:

Goat

Cattle

Sheep

Route of administration:

Intramammary use

Product details

Active substance and strength:

Dihydrostreptomycin sulfate

100.00 milligram(s) / 1.00 Syringe

Nafcillin sodium monohydrate

100.00 milligram(s) / 1.00 Syringe

Benzympenicillin procaine

300.00 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

-

Cattle

- Meat and offal. 16 day

- Milk. 48 hour

-

Sheep

- Meat and offal. 35 day

- Milk. 5 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:

Greece

Available in:

Greece

Package description:

Available only in Greek

Available only in Greek

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Complete application (stand-alone)

Marketing authorisation holder:

Intervet International B.V.

Marketing authorisation date:

23/02/1998

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

National Organization For Medicines

Authorisation number:

18132/09-03-2009/K-0112802

Date of authorisation status change:

8/07/2020

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