

Nuflor Minidose 450 mg/ml solution for injection for cattle

Not
authorised

- Florfenicol

Product identification

Medicine name:

Nuflor Minidose 450 mg/ml solution for injection for cattle
Nuflor Minidose 450 mg/ml ενέσιμο διάλυμα για βοοειδή

Active substance:

Florfenicol

Target species:

Cattle

Route of administration:

Subcutaneous use
Intramuscular use

Product details

Active substance and strength:

Florfenicol
450.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:**Subcutaneous use:**

-

Cattle

- Milk. no withdrawal period

Not permitted for use in lactating animals producing milk for human consumption.

- Meat and offal. 64 day

Intramuscular use:

-

Cattle

- Milk. no withdrawal period

Not permitted for use in lactating animals producing milk for human consumption.

- Meat and offal. 37 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01BA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Cyprus

Package description:

(ID3) 250 millilitre(s): unspecified outer container with 1 Vial (Glass) with 250 millilitre(s), closed with Stopper (bromobutyl rubber`)

(ID2) 100 millilitre(s): unspecified outer container with 1 Vial (Glass) with 100 millilitre(s), closed with Stopper (bromobutyl rubber`)

(ID1) 50 millilitre(s): unspecified outer container with 1 Vial (Glass) with 50 millilitre(s), closed with Stopper (bromobutyl rubber`)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

Intervet International B.V.

Marketing authorisation date:

7/12/2008

Manufacturing sites for batch release:

Intervet International GmbH

Trirx Segre

Vet Pharma Friesoythe GmbH

Responsible authority:

Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:

CY00163V

Date of authorisation status change:

16/03/2021

Reference member state:

Germany

Procedure number:

DE/V/0122/001

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

Documents

Summary of Product Characteristics

Combined File of all Documents

English (PDF)

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