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 Oplossing voor injectie
Nuflor Minidose 450 mg/ml Solution injectable
Nuflor Minidose 450 mg/ml Injektionslösung

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:
Belgium

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:
18/08/2008

Manufacturing sites for batch release:
Intervet International GmbH
Trirx Segre
Vet Pharma Friesoythe GmbH

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V321964

Date of authorisation status change:
13/02/2024

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

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

Package Leaflet

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

Labelling

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