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

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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)
Published on: 26/02/2024
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