File downloaded on 2026-02-04
Source URL: https://medicines.health.europa.eu/veterinary/en/600000060538

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 450MG/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:

Greece

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 Hellas A.E.

Marketing authorisation date:
16/09/2015

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

Responsible authority:
National Organization For Medicines

Authorisation number:
61387/17-09-2015/K-0096505

Date of authorisation status change:
20/05/2024

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