NUFLOR Swine 300 mg/ml Solution for injection

Authorised

This information is not available for this product.

Product identification

Medicine name:
NUFLOR Swine 300 mg/ml Solution for injection
Nuflor Swine 300 mg/ml Oplossing voor injectie
Nuflor Swine 300 mg/ml Solution injectable
Nuflor Swine 300 mg/ml Injektionslösung

Active substance:
This information is not available for this product.

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
This information is not available for this product.

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Pig**

  - Meat and offal. 18 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01BA90

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Belgium

Package description:
Pack Size 20 ml colourless Type I glass vial closed with Grey bromobutyl rubber stopper with aluminium seal.
Pack Size 50 ml colourless Type I glass vial closed with Grey bromobutyl rubber stopper with aluminium seal.
Pack Size 100 ml colourless Type I glass vial closed with Grey bromobutyl rubber stopper with aluminium seal.
Pack Size 250 ml colourless Type I glass vial closed with Grey bromobutyl rubber stopper with aluminium seal.
Pack Size 500 ml colourless Type I glass vial closed with Grey bromobutyl rubber stopper with aluminium seal.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
15/01/2001

Manufacturing sites for batch release:
Trirx Segre

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V220927

Date of authorisation status change:
15/01/2001

Reference member state:
Ireland

Procedure number:
IE/V/0593/001

Concerned member states:
Austria Belgium Greece Italy Luxembourg Netherlands Portugal Spain

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 11/02/2022
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Package Leaflet

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