NUFLOR Swine 300 mg/ml Solution for injection

Suspended

This information is not available for this product.

Product identification

Medicine name:
NUFLOR Swine 300 mg/ml Solution for injection
NUFLOR 300 mg/ml SOLUCION INYECTABLE PORCINO

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:
Suspended

Authorised in:
Spain

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:
Merck Sharp & Dohme Animal Health S.L.

Marketing authorisation date:

28/09/2000

Manufacturing sites for batch release:
Trirx Segre

Responsible authority:
Spanish Agency For Medicines And Health Products

Authorisation number:
1349 ESP

Date of authorisation status change:
25/12/2021

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
Download

Package Leaflet

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Labelling

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