MEGANYL 50 mg/ml Solution for Injection for cattle, pigs and horses.

Authorised

- Flunixin meglumine

Product identification

Medicine name:
MEGANYL 50 mg/ml Solution for Injection for cattle, pigs and horses.

Active substance:
Flunixin meglumine

Target species:
Cattle
Horse
Pig

Route of administration:
Intravenous use
Intramuscular use

Product details

Active substance and strength:
Flunixin meglumine

83.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intravenous use:

- **Cattle**
  - Meat and offal. 4 day
  - Milk. 24 hour
- **Horse**
  - Meat and offal. 4 day
  - Milk. no withdrawal period

Milk: Not authorised for use in mares producing milk for human consumption.

- **Cattle**
  - Meat and offal. 4 day
  - Milk. 24 hour
- **Horse**
  - Meat and offal. 4 day
  - Milk. no withdrawal period

Milk: Not authorised for use in mares producing milk for human consumption.

Intramuscular use:

- **Pig**
  - Meat and offal. 24 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM01AG90

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

Authorisation status:
Valid

Authorised in:
France

Package description:
box containing 1 vial of 250 ml
box containing 1 vial of 100 ml

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:
Laboratorios Syva S.A.

Marketing authorisation date:
9/05/2019

Manufacturing sites for batch release:
Laboratorios Syva S.A.

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/8615169 0/2019

Date of authorisation status change:
27/08/2020

Reference member state:

Spain

Procedure number:
ES/V/0249/001

Concerned member states:
Belgium Bulgaria Cyprus France Greece Hungary Italy Poland Portugal Romania

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

Documents

Package Leaflet and Labelling

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

Summary of Product Characteristics

English (PDF)
Published on: 5/04/2023
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Package Leaflet

Labelling

Combined File of all Documents