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

Authorised

- Flunixin meglumine

Product identification

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

Active substance:
Flunixin meglumine

Target species:
Cattle
Pig
Horse

Route of administration:
Intramuscular use
Intravenous 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:
Intramuscular use:

- **Cattle**
  - Meat and offal. 31 day
  - Milk. 36 hour
- **Pig**
  - Meat and offal. 24 day

Intravenous use:

- **Cattle**
  - Meat and offal. 4 day
  - Milk. 24 hour
- **Horse**
  - Meat and offal. 5 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM01AG90

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

Authorisation status:
Valid

Authorised in:
Spain

Package description:
50 ml clear Type I glass Multidose vial, with Bromobutyl rubber bung.
100 ml clear Type I glass Multidose vial, with Bromobutyl rubber bung.

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:
Bimeda Animal Health Limited

Marketing authorisation date:
9/05/2002

Manufacturing sites for batch release:
Labiana Life Sciences S.A.

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
1450 ESP

Date of authorisation status change:
9/09/2020

Reference member state:
Ireland

Procedure number:
IE/V/0125/001

Concerned member states:
Belgium Germany Italy Luxembourg Netherlands 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: 11/05/2025
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Package Leaflet

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Labelling

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Combined File of all Documents