MEGLUXIN 50 mg/ml

Authorised

- Flunixin meglumine

Product identification

Medicine name:
MEGLUXIN 50 mg/ml

Active substance:
Flunixin meglumine

Target species:
Pig
Cattle
Horse

Route of administration:
Intramuscular use
Intravenous use

Product details

Active substance and strength:
Flunixin meglumine
50.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Pig**
  - Meat and offal. 17 day

Intravenous use:

- **Cattle**
  - Meat and offal. 28 day

The product is not administered to horses whose milk is intended for human consumption.

- Meat and offal. 14 day
- Milk. 2 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM01AG90

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

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Authorisation status:
Valid

Authorised in:
Romania

Package description:
Available only in Romanian
Available only in Romanian
Available only in Romanian

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Directive No 2001/82/EC

Marketing authorisation holder:
Laboratorios Hipra S.A.

Marketing authorisation date:
22/04/2007

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

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:
160170

Date of authorisation status change:
29/10/2025

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

Documents

Combined File of all Documents

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