MEFLOSYL 50 mg/ml Solution for injection for cattle, pigs and horses

Authorised

- Flunixin meglumine

Product identification

Medicine name:
MEFLOSYL 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**
  - Milk. no withdrawal period

Not authorised for use in animals producing milk for human consumption.

  - Meat and offal. 5 day

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:
Slovakia

Package description:

(ID1) 100 millilitre(s): Box (cardboard) with 1 Vial (Glass type I) with 100 millilitre(s), closed with Lid (aluminium) and Stopper (bromobutyl rubber)
(ID2) 50 millilitre(s): Box (cardboard) with 1 Vial (Glass type I) with 50 millilitre(s), closed with Lid (aluminium) and Stopper (bromobutyl rubber)
(ID3) 250 millilitre(s): Box (cardboard) with 1 Vial (Glass type I) with 250 millilitre(s), closed with Lid (aluminium) and Stopper (bromobutyl rubber)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Zoetis Ceska Republika s.r.o.

Marketing authorisation date:
13/05/1999

Manufacturing sites for batch release:
Zoetis Manufacturing & Research Spain, S.L.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/047/99-S

Date of authorisation status change:
13/05/1999

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

Documents

Combined File of all Documents

English (PDF)
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