

Flunixin 50 mg/ml Solution for Injection for Cattle, Horses and Pigs

Authorised

- Flunixin meglumine

Product identification

Medicine name:

Flunixin 50 mg/ml Solution for Injection for Cattle, Horses and Pigs

Active substance:

Flunixin meglumine

Target species:

Cattle

Horse

Pig

Route of administration:

Intravenous use

Intramuscular use

Product details

Active substance and strength:

Flunixin meglumine

82.90 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intravenous use:

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Cattle

- Milk. 36 hour
- Meat and offal. 7 day

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Horse

- Milk. no withdrawal period

Do not use in mares producing milk for human consumption.

- Meat and offal. 7 day

Intramuscular use:

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Pig

- Meat and offal. 22 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QM01AG90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Portugal

Available in:

Portugal

Package description:

(ID1) 50 millilitre(s): unspecified outer container with 1 Vial (Glass) with 50 millilitre(s)

(ID2) 100 millilitre(s): unspecified outer container with 1 Vial (Glass) with 100 millilitre(s)

(ID3) 250 millilitre(s): unspecified outer container with 1 Vial (Glass) with 250 millilitre(s)

(ID4) 250 millilitre(s): unspecified outer container with 5 Vial (Glass) each with 50 millilitre(s)

(ID5) 500 millilitre(s): unspecified outer container with 10 Vial (Glass) each with 50 millilitre(s)

(ID6) 600 millilitre(s): unspecified outer container with 12 Vial (Glass) each with 50 millilitre(s)

(ID7) 500 millilitre(s): unspecified outer container with 5 Vial (Glass) each with 100 millilitre(s)

(ID8) 1000 millilitre(s): unspecified outer container with 10 Vial (Glass) each with 100 millilitre(s)

(ID9) 1200 millilitre(s): unspecified outer container with 12 Vial (Glass) each with 100 millilitre(s)

(ID10) 1250 millilitre(s): unspecified outer container with 5 Vial (Glass) each with 250 millilitre(s)

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:

Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:

30/12/2008

Manufacturing sites for batch release:

Norbrook Laboratories (Ireland) Limited

Norbrook Laboratories Limited

Responsible authority:

Directorate General For Food And Veterinary

Authorisation number:

133/01/08RFVPT

Date of authorisation status change:

26/01/2026

Reference member state:

Germany

Procedure number:

DE/V/0325/001

Concerned member states:

Iceland Netherlands Portugal Sweden United Kingdom (Northern Ireland)

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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