# 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:**

- **Cattle**
  - Milk. 36 hour
  - Meat and offal. 7 day
- **Horse**
  - Milk. no withdrawal period

Do not use in mares producing milk for human consumption.

  - Meat and offal. 7 day

**Intramuscular use:**

- **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:**
(ID10) 1250 millilitre(s): unspecified outer container with 5 Vial (Glass) each with 250 millilitre(s)
(ID9) 1200 millilitre(s): unspecified outer container with 12 Vial (Glass) each with 100 millilitre(s)
(ID8) 1000 millilitre(s): unspecified outer container with 10 Vial (Glass) each with 100 millilitre(s)
(ID7) 500 millilitre(s): unspecified outer container with 5 Vial (Glass) each with 100 millilitre(s)
(ID6) 600 millilitre(s): unspecified outer container with 12 Vial (Glass) each with 50 millilitre(s)
(ID5) 500 millilitre(s): unspecified outer container with 10 Vial (Glass) each with 50 millilitre(s)
(ID4) 250 millilitre(s): unspecified outer container with 5 Vial (Glass) each with 50 millilitre(s)
(ID3) 250 millilitre(s): unspecified outer container with 1 Vial (Glass) with 250 millilitre(s)
(ID2) 100 millilitre(s): unspecified outer container with 1 Vial (Glass) with 100 millilitre(s)
(ID1) 50 millilitre(s): unspecified outer container with 1 Vial (Glass) with 50 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

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