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
Flunixin N-vet 50 mg/ml Injektionsvätska, lösning

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:
Sweden

Available in:

Sweden

Package description:
Available only in Swedish
Available only in Swedish
Available only in Swedish
Available only in Swedish
Available only in Swedish
Available only in Swedish
Available only in Swedish
Available only in Swedish
Available only in Swedish
Available only in Swedish

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:
27/10/2009

Manufacturing sites for batch release:
Norbrook Laboratories (Ireland) Limited
Norbrook Laboratories Limited

Responsible authority:
Swedish Medical Products Agency

Authorisation number:
27763

Date of authorisation status change:

27/10/2009

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

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.