

Flunixin Injection

Authorised

- Flunixin meglumine

Product identification

Medicine name:

Flunixin Injection

Active substance:

Flunixin meglumine

Target species:

Cattle

Pig

Horse

Route of administration:

Intramuscular use

Intravenous use

Product details

Active substance and strength:

Flunixin meglumine

82.96 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:**Intramuscular use:**

-

Cattle

- Meat and offal. 31 day
- Milk. 36 hour

-

Pig

- Meat and offal. 24 day

Intravenous use:

-

Cattle

- Meat and offal. 4 day
- Milk. 24 hour

-

Horse

- Meat and offal. 5 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QM01AG90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Available in:

Ireland

Package description:

Flunixin Injection is supplied in 50ml type I clear glass vial, complete with bromobutyl bung and aluminium cap.

Flunixin Injection is supplied in 100ml type I clear glass vial, complete with bromobutyl bung and aluminium cap.

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:

23/04/1997

Manufacturing sites for batch release:

Norbrook Laboratories Limited

Norbrook Manufacturing Limited

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA22664/046/001

Date of authorisation status change:

23/04/1997

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

Documents

Summary of Product Characteristics