

Flunazine 50 mg/ml Solution for Injection for cattle, horses and pigs

Authorised

- Flunixin meglumine

Product identification

Medicine name:

Flunazine 50 mg/ml Solution for Injection for cattle, horses and pigs

Active substance:

Flunixin meglumine

Target species:

Cattle

Pig

Horse

Route of administration:

Intramuscular use

Intravenous 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:

Intramuscular use:

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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

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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:

Germany

Package description:

50 ml clear Type I glass Multidose vial, with Bromobutyl rubber bung.
100 ml clear Type I glass Multidose vial, with Bromobutyl rubber bung.

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:

Bimeda Animal Health Limited

Marketing authorisation date:

18/12/2011

Manufacturing sites for batch release:

Labiana Life Sciences S.A.

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:

401619.00.00

Date of authorisation status change:

18/12/2011

Reference member state:

Ireland

Procedure number:

IE/V/0125/001

Concerned member states:

Belgium Germany Italy Luxembourg Netherlands Spain

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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