ALIVIOS, flunixin meglumina, soluzione iniettabile per bovini, suini ed equini

Authorised

- Flunixin meglumine

Product identification

Medicine name:
ALIVIOS, flunixin meglumina, soluzione iniettabile per bovini, suini ed equini

Active substance:
Flunixin meglumine

Target species:
Cattle
Pig
Horse

Route of administration:
Intramuscular use
Intravenous use

Product details

Active substance and strength:
Flunixin meglumine
82.95 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Milk. 48 hour pari a 4 mungiture
  - Meat and offal. 7 day
- **Pig**
  - Meat and offal. 18 day
- **Horse**
  - Meat and offal. 7 day

Uso non autorizzato in equidi che producono alimenti per il consumo umano

Intravenous use:

- **Cattle**
  - Milk. 48 hour pari a 4 mungiture
  - Meat and offal. 7 day
- **Pig**
  - Meat and offal. 18 day
- **Horse**
  - Meat and offal. 7 day

Uso non autorizzato in equidi che producono alimenti per il consumo umano

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Anatomical therapeutic chemical veterinary (ATCvet) codes:

QM01AG90

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Italy

Available in:
Italy

Package description:
Available only in Italian
Available only in Italian
Available only in Italian
Available only in Italian
Available only in Italian
Available only in Italian
Available only in Italian
Available only in Italian
Available only in Italian
Available only in Italian
Available only in Italian
Available only in Italian

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:
Fatro S.p.A.

Marketing authorisation date:
4/06/2003

Manufacturing sites for batch release:
Fatro S.p.A.

Responsible authority:
Ministry Of Health

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
29/01/2019

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

Documents

Combined File of all Documents

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