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MEGLUVET 50 mg/ml SOLUCIÓN INYECTABLE

Authorised

- Flunixin meglumine

Product identification

Medicine name:
MEGLUVET 50 mg/ml SOLUCIÓN INYECTABLE

Active substance:
Flunixin meglumine

Target species:
Cattle
Horse

Route of administration:
Intravenous use

Product details

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

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intravenous use:

- **Cattle**
  - Meat and offal. 4 day
  - Milk. 24 hour
- **Horse**
  - Meat and offal. 4 day
  - Milk. no withdrawal period

Leche: Su uso no está autorizado en yeguas en lactación cuya leche se utiliza para consumo humano

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Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM01AG90

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

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Authorisation status:
Valid

Authorised in:
Spain

Available in:
Spain

Package description:
Available only in Spanish
Available only in Spanish

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:
Super's Diana S.L.

Marketing authorisation date:
27/03/2008

Manufacturing sites for batch release:
Super's Diana S.L.

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
1863 ESP

Date of authorisation status change:
28/03/2008

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

Documents

Summary of Product Characteristics

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Package Leaflet

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

Labelling

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