Finadyne Transdermal 50 mg/ml pour-on solution for cattle

Not authorised

- Flunixin meglumine

Product identification

Medicine name:
Finadyne Transdermal 50 mg/ml pour-on solution for cattle
Finadyne transdermal 50 mg/ml solução para unção contínua para bovinos

Active substance:
Flunixin meglumine

Target species:
Cattle

Route of administration:
Pour-on use

Product details

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

Pharmaceutical form:
Pour-on solution

Withdrawal period by route of administration:

Pour-on use:

- **Cattle**
  - Meat and offal. 7 day
  - Milk. 36 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM01AG90

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

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Authorisation status:
Revoked

Authorised in:
Portugal

Package description:
High density polyethylene (HDPE) bottles with polypropylene (PP) closures which have a peelable foil laminate induction innerseal and a liner. The bottles are equipped with a graduated dosing chamber and are supplied individually in a cardboard carton.Container size: 100 ml
High density polyethylene (HDPE) bottles with polypropylene (PP) closures which have a peelable foil laminate induction innerseal and a liner. The bottles are equipped with a graduated dosing chamber and are supplied individually in a cardboard carton.Container size: 250 ml
High density polyethylene (HDPE) bottles with polypropylene (PP) closures which have a peelable foil laminate induction innerseal and a liner. The bottles are equipped with a graduated dosing chamber and are supplied individually in a cardboard carton.Container size: 1000 ml

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Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
MSD Animal Health Lda.

Marketing authorisation date:
23/06/2014

Manufacturing sites for batch release:
Vet Pharma Friesoythe GmbH

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
812/01/14DFVPT

Date of authorisation status change:
2/11/2023

Reference member state:
Spain

Procedure number:
ES/V/0451/001

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

Documents

Summary of Product Characteristics