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

Authorised

- Flunixin meglumine

Product identification

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

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:

Valid

Authorised in:

Italy

Available in:

Italy

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:
Intervet International B.V.

Marketing authorisation date:
5/06/2014

Manufacturing sites for batch release:
Vet Pharma Friesoythe GmbH

Responsible authority:
Ministry Of Health

Authorisation number:
104637

Date of authorisation status change:
5/06/2014

Reference member state:
Spain

Procedure number:
ES/V/0451/001

Concerned member states:
Austria Belgium Bulgaria Cyprus Czechia Denmark Estonia Finland France Germany Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Netherlands Norway Poland Portugal Romania Slovakia Slovenia Sweden United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics