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TOLFEDOL, 40 mg/ml, solution for injection for cattle, pigs, cats and dogs

Authorised

- Tolfenamic acid

Product identification

Medicine name:
TOLFEDOL, 40 mg/ml, solution for injection for cattle, pigs, cats and dogs

Active substance:
Tolfenamic acid

Target species:
Cattle
Dog
Pig
Cat

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
Tolfenamic acid
40.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat and offal. 12 day
  - Milk. 0 hour
- **Pig**
  - Meat and offal. 16 day

Intravenous use:

- **Cattle**
  - Meat and offal. 4 day
  - Milk. 24 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM01AG02

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

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Authorisation status:
Valid

Authorised in:

France

Package description:
Amber polypropylene vial of 250 ml provided with a pink bromobutyl stopper and aluminium seal with a flip-off sealing.Each vial is packaged in an outer carton.
Amber polypropylene vial of 100 ml provided with a grey bromobutyl stopper and aluminium seal with a flip-off sealing.Each vial is packaged in an outer carton.
Amber polypropylene vial of 50 ml provided with a grey bromobutyl stopper and aluminium seal with a flip-off sealing.Each vial is packaged in an outer carton.
Amber polypropylene vial of 20 ml provided with a grey bromobutyl stopper and aluminium seal with a flip-off sealing.Each vial is packaged in an outer carton.

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:
S P Veterinaria S.A.

Marketing authorisation date:
18/06/2015

Manufacturing sites for batch release:
S P Veterinaria S.A.

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/6409545 3/2015

Date of authorisation status change:
26/01/2021

Reference member state:

Ireland

Procedure number:
IE/V/0344/001

Concerned member states:
France Germany Portugal Romania Slovakia Spain

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 25/09/2024
Updated on: 14/03/2026
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Package Leaflet and Labelling

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

Combined File of all Documents