Trovex 1 mg/ml suspension for injection for cattle, horses, pigs, cats and dogs

Authorised

- Dexamethasone isonicotinate

Product identification

Medicine name:
Trovex 1 mg/ml suspension for injection for cattle, horses, pigs, cats and dogs
Trovex 1 mg/ml suspension for injection for cattle, horses, pigs, cats and dogs

Active substance:
Dexamethasone isonicotinate

Target species:
Cattle
Dog
Horse
Cat
Pig

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Dexamethasone isonicotinate
1.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat and offal. 55 day
  - Milk. 60 hour
- **Dog**
- **Horse**
  - Meat and offal. 63 day
- **Cat**
- **Pig**
  - Meat and offal. 55 day

Subcutaneous use:

- **Dog**
- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH02AB02

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

Authorisation status:
Valid

Authorised in:
Ireland

Available in:
Ireland

Package description:
One amber, glass (Ph. Eur. Type I or Ph. Eur. siliconized Type II) multidose vial containing 50 ml of product, sealed with a grey siliconized bromobutyl rubber stopper and aluminium cap, in a cardboard boxPack sizes:Cardboard box containing 1 vial of 50 ml

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:
Emdoka

Marketing authorisation date:
12/11/2021

Manufacturing sites for batch release:
Divasa Farmavic S.A.

Responsible authority:
Health Products Regulatory Authority

Authorisation number:

VPA10534/009/001

Date of authorisation status change:
12/11/2021

Reference member state:
Ireland

Procedure number:
IE/V/0517/001

Concerned member states:
Austria Bulgaria Croatia Czechia Estonia Finland France Germany Greece Hungary Italy Luxembourg Netherlands Poland Portugal Romania Slovakia Slovenia United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

Combined File of all Documents