# 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

**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
- **Horse**
  - Meat and offal. 63 day
- **Pig**
  - Meat and offal. 55 day

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QH02AB02

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

**Authorisation status:**
Valid

**Authorised in:**
Greece

**Available in:**

Greece

**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:**
8/11/2021

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

**Responsible authority:**
National Organization For Medicines

**Authorisation number:**
104378 / 09-11-2021 / K-0245601

**Date of authorisation status change:**
8/11/2021

**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)

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

## Documents

Summary of Product Characteristics

English (PDF)
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Combined File of all Documents

Package Leaflet

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