

# 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

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**Active substance:**

Dexamethasone isonicotinate

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**Target species:**

Cattle

Dog

Horse

Cat

Pig

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**Route of administration:**

Intramuscular use

Subcutaneous use

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## Product details

**Active substance and strength:**

Dexamethasone isonicotinate  
1.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Suspension for injection

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**Withdrawal period by route of administration:****Intramuscular use:**

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**Cattle**

- Meat and offal. 55 day
- Milk. 60 hour

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**Horse**

- Meat and offal. 63 day

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**Pig**

- Meat and offal. 55 day
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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QH02AB02

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

France

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**Available in:**

France

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**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

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Emdoka

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**Marketing authorisation date:**

12/08/2021

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**Manufacturing sites for batch release:**

Divasa Farmavic S.A.

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/4610056 6/2021

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**Date of authorisation status change:**

12/08/2021

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0517/001

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**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](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

English (PDF)

Published on: 5/10/2025

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Combined File of all Documents

Package Leaflet and Labelling

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