

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

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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:

Romania

Available in:

Romania

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:

11/01/2022

Manufacturing sites for batch release:

Divasa Farmavic S.A.

Responsible authority:

Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:

220008

Date of authorisation status change:

3/09/2025

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

Combined File of all Documents

English (PDF)

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Summary of Product Characteristics