Dexafast 2 mg/ml solution for injection for horses, cattle, goats, pigs, dogs and cats

Authorised

- Dexamethasone sodium phosphate

Product identification

Medicine name:
Dexafast 2 mg/ml solution for injection for horses, cattle, goats, pigs, dogs and cats

Active substance:
Dexamethasone sodium phosphate

Target species:
Horse
Cattle
Dog
Goat
Cat
Pig

Route of administration:
Intraarticular use
Intramuscular use
Intravenous use
Periarticular use
Subcutaneous use

Product details

Active substance and strength:
Dexamethasone sodium phosphate
2.63 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intraarticular use:

- **Horse**
  - Meat and offal. 8 day

Intramuscular use:

- **Cattle**
  - Meat and offal. 8 day
  - Milk. 72 hour
- **Goat**
  - Meat and offal. 8 day
  - Milk. 72 hour
- **Horse**
  - Meat and offal. 8 day
- **Pig**
  - Meat and offal. 2 day

Intravenous use:

-

Cattle

- Meat and offal. 8 day
- Milk. 72 hour

•

Goat

- Meat and offal. 8 day
- Milk. 72 hour

•

Horse

- Meat and offal. 8 day

•

Pig

- Meat and offal. 6 day

Periarticular use:

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Horse

- Meat and offal. 8 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH02AB02

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

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Authorisation status:
Valid

Authorised in:
Czechia

Available in:
Czechia

Package description:

Clear glass (Type I Ph. Eur.) vial of 100 ml closed with a bromobutyl rubber stopper and sealed with an aluminium cap.Carton of 12 x 100 ml
Clear glass (Type I Ph. Eur.) vial of 100 ml closed with a bromobutyl rubber stopper and sealed with an aluminium cap.Carton of 6 x 100 ml
Clear glass (Type I Ph. Eur.) vial of 100 ml closed with a bromobutyl rubber stopper and sealed with an aluminium cap.Carton of 1 x 100 ml
Clear glass (Type I Ph. Eur.) vial of 50 ml closed with a bromobutyl rubber stopper and sealed with an aluminium cap.Carton of 12 x 50 ml
Clear glass (Type I Ph. Eur.) vial of 50 ml closed with a bromobutyl rubber stopper and sealed with an aluminium cap.Carton of ml, 6 x 50 ml
Clear glass (Type I Ph. Eur.) vial of 50 ml closed with a bromobutyl rubber stopper and sealed with an aluminium cap.Carton of 1 x 50 ml
Clear glass (Type I Ph. Eur.) vial of 20 ml closed with a bromobutyl rubber stopper and sealed with an aluminium cap.Carton of 12 x 20 ml
Clear glass (Type I Ph. Eur.) vial of 20 ml closed with a bromobutyl rubber stopper and sealed with an aluminium cap.Carton of 6 x 20 ml
Clear glass (Type I Ph. Eur.) vial of 20 ml closed with a bromobutyl rubber stopper and sealed with an aluminium cap.Carton of 1 x 20 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:
Industrial Veterinaria S.A.

Marketing authorisation date:
13/07/2023

Manufacturing sites for batch release:
aniMedica GmbH
Industrial Veterinaria S.A.

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/028/23-C

Date of authorisation status change:
13/07/2023

Reference member state:
Ireland

Procedure number:
IE/V/0390/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland France Germany Greece Hungary Italy Latvia Lithuania Netherlands Poland Portugal Romania Slovakia Slovenia Spain 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

Summary of Product Characteristics

English (PDF)
Published on: 25/01/2026
Updated on: 13/03/2026

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Package Leaflet

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Labelling

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