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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 Dexafast

Active substance:

Dexamethasone sodium phosphate

Target species:

Horse

Cattle

Dog

Goat

Cat

Pig

Route of administration:

Intraarticular use

Intramuscular use

Intravenous use

Periarticular 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:

Available in:	
Authorised in: Germany	
Authorisation status: Valid	
Legal status of supply: Veterinary medicinal product subject to veterinary prescription	
Anatomical therapeutic chemical veterinary (ATCvet) codes: QH02AB02	
Horse - Meat and offal. 8 day	
Periarticular use: •	
Pig - Meat and offal. 6 day	
•	
Horse - Meat and offal. 8 day	
•	
- Milk. 72 hour	
Goat - Meat and offal. 8 day	
•	
- Milk. 72 hour	
- Meat and offal. 8 day	
• Cattle	

Germany

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 6×100 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 1×100 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 12×50 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 1×50 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 $12 \times 20 \text{ ml}$

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 $1 \times 20 \text{ ml}$

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:

23/01/2019

Manufacturing sites for batch release:

Industrial Veterinaria S.A. aniMedica GmbH

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:

402526.00.00

Date of authorisation status change:

23/01/2019

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

English (PDF)

Published on: 9/11/2025

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