

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
- Milk. no withdrawal period

Not authorised for use in horses producing milk for human consumption.

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
- Milk. no withdrawal period

Not authorised for use in horses producing milk for human consumption.

-

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
- Milk. no withdrawal period

Not authorised for use in horses producing milk for human consumption.

-

Pig

- Meat and offal. 6 day

Periarticular use:

-

Horse

- Meat and offal. 8 day
- Milk. no withdrawal period

Not authorised for use in horses producing milk for human consumption.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH02AB02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Denmark

Available in:

Denmark

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:

3/01/2019

Manufacturing sites for batch release:

Industrial Veterinaria S.A.

aniMedica GmbH

Responsible authority:

Danish Medicines Agency

Authorisation number:

60463

Date of authorisation status change:

3/01/2019

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