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RAPIDEXON

Authorised

- Dexamethasone sodium phosphate

Product identification

Medicine name:

RAPIDEXON

Active substance:

Dexamethasone sodium phosphate

Target species:

Cattle

Dog

Goat

Cat

Pig

Horse

Route of administration:

Intramuscular use

Intravenous use

Intraarticular 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:

Intramuscular use:

•

Cattle

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

•

Goat

- Milk. 14 hour
- Meat and offal. 60 day

•

Pig

- Meat and offal. 2 day

•

Horse

- Meat and offal. 8 day

Usò non autorizzato in equidi che producono latte per il consumo umano

Intravenous use:

•

Cattle

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

•

Goat

- Milk. 14 hour
- Meat and offal. 60 day

- Milk. 14 hour
- Meat and offal. 60 day

-

Goat

- Milk. 14 hour
- Meat and offal. 60 day
- Milk. 14 hour
- Meat and offal. 60 day

-

Pig

- Meat and offal. 2 day

-

Horse

- Meat and offal. 8 day

Use non autorizzato in equidi che producono latte per il consumo umano

Intraarticular use:

-

Horse

- Meat and offal. 8 day

Use non autorizzato in equidi che producono latte per il consumo umano

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH02AB02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Italy

Available in:

Italy

Package description:

Available only in [Italian](#)

Available only in [Italian](#)

Available only in [Italian](#)

Available only in [Italian](#)

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:

Eurovet Animal Health B.V.

Marketing authorisation date:

2/03/2007

Manufacturing sites for batch release:

Eurovet Animal Health B.V.

Responsible authority:

Ministry Of Health

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

2/03/2007

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents

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