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 ì
- **Goat**
  - Milk. 14 hour
  - Meat and offal. 60 day
- **Pig**
  - Meat and offal. 2 day
- **Horse**
  - Meat and offal. 8 day

Uso 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

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

Intraarticular use:

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

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

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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:
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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