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RAPISON 2 mg/ml soluzione iniettabile per bovini, equini, suini

Authorised

- Dexamethasone

Product identification

Medicine name:
RAPISON 2 mg/ml soluzione iniettabile per bovini, equini, suini

Active substance:
Dexamethasone

Target species:
Cattle
Horse
Pig

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

Product details

Active substance and strength:

Dexamethasone
2.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Milk. 60 hour
  - Meat and offal. 7 day
- **Horse**
  - Meat and offal. 11 day

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

- **Pig**
  - Meat and offal. 2 day

Intravenous use:

- **Cattle**
  - Milk. 60 hour
  - Meat and offal. 7 day
- **Horse**
  - Meat and offal. 11 day

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

- **Pig**
  - Meat and offal. 2 day

Subcutaneous use:

- **Cattle**
  - Milk. 60 hour
  - Meat and offal. 7 day
- **Horse**
  - Meat and offal. 11 day

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

- **Pig**
  - Meat and offal. 2 day

Intraarticular use:

- **Cattle**
  - Milk. 60 hour
  - Meat and offal. 7 day
- **Horse**
  - Meat and offal. 11 day

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

- **Pig**
  - Meat and offal. 2 day

Infiltration:

- **Cattle**
  - Milk. 60 hour
  - Meat and offal. 7 day

- **Horse**
  - Meat and offal. 11 day

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

- **Pig**
  - Meat and offal. 2 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH02AB02

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

Authorisation status:
Valid

Authorised in:
Italy

Package description:
Available only in Italian
Available only in Italian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Azienda Terapeutica Italiana A.T.I. S.r.l.

Marketing authorisation date:
24/02/1994

Manufacturing sites for batch release:
Fatro S.p.A.

Responsible authority:
Ministry Of Health

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
24/02/2009

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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