VITALENE C, 200 mg/ml, soluzione iniettabile per bovini, equini, suini, ovini, caprini, cani, gatti e conigli

Authorised

- Ascorbic acid

Product identification

Medicine name:
VITALENE C, 200 mg/ml, soluzione iniettabile per bovini, equini, suini, ovini, caprini, cani, gatti e conigli

Active substance:
Ascorbic acid

Target species:
Cattle
Dog
Goat
Sheep
Cat
Rabbit
Pig
Horse (food producing)

Route of administration:

Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
Ascorbic acid
200.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Goat**
  - Meat and offal. 0 day
- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Rabbit**
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day

- **Horse (food producing)**
  - Meat and offal. 0 day
  - Milk. 0 hour

Intravenous use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 hour

- **Goat**
  - Milk. 0 day
  - Meat and offal. 0 day

- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 hour

- **Rabbit**
  - Meat and offal. 0 day

- **Pig**
  - Meat and offal. 0 day

- **Horse (food producing)**
  - Meat and offal. 0 day
  - Milk. 0 hour

Subcutaneous use:

-

Cattle

- Meat and offal. 0 day
- Milk. 0 hour

•

Goat

- Meat and offal. 0 day
- Milk. 0 day

•

Sheep

- Meat and offal. 0 day
- Milk. 0 hour

•

Rabbit

- Meat and offal. 0 day

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Pig

- Meat and offal. 0 day

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Horse (food producing)

- Meat and offal. 0 day
- Milk. 0 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA11GA01

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

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:
Fatro S.p.A.

Marketing authorisation date:
4/06/1997

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:
4/06/1997

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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