

VITALENE E 50 mg/ml emulsione iniettabile per bovini, equini, suini, ovini, caprini, cani, gatti e conigli

Authorised

- TOCOPHEROL

Product identification

Medicine name:

VITALENE E 50 mg/ml emulsione iniettabile per bovini, equini, suini, ovini, caprini, cani, gatti e conigli

Active substance:

TOCOPHEROL

Target species:

Cattle
Dog
Goat
Sheep
Horse
Cat
Rabbit
Pig

Route of administration:

Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:

TOCOPHEROL

45.56 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
- Milk. 0 hour

-

Sheep

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

-

Horse

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

-

Rabbit

- Meat and offal. 0 day

-

Pig

- Meat and offal. 0 day

Intravenous use:

-

Cattle

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

-

Goat

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

-

Sheep

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

-

Horse

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

-

Rabbit

- Meat and offal. 0 day

-

Pig

- Meat and offal. 0 day

Subcutaneous use:

-

Cattle

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

-

Goat

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

-

Sheep

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

-

Horse

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

-

Rabbit

- Meat and offal. 0 day

-

Pig

- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA11HA03

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:

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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