ZOOLOBELIN 1,8 mg/ml soluzione iniettabile per bovini, equini, suini, ovini, cani

Authorised

- Lobeline

Product identification

Medicine name:
ZOOLOBELIN 1,8 mg/ml soluzione iniettabile per bovini, equini, suini, ovini, cani

Active substance:
Lobeline

Target species:
Cattle
Dog
Sheep
Pig
Horse

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
Lobeline
1.80 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Milk. 0 hour
  - Meat and offal. 0 day
- **Dog**
  - Unspecified. 0 day
- **Sheep**
  - Milk. 0 hour
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day
- **Horse**
  - Milk. 0 hour
  - Meat and offal. 0 day

Intravenous use:

-

Cattle

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

•

Dog

- Unspecified. 0 day

•

Sheep

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

•

Pig

- Meat and offal. 0 day

•

Horse

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

Subcutaneous use:

•

Cattle

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

•

Dog

- Unspecified. 0 day

•

Sheep

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

- **Pig**
  - Meat and offal. 0 day
- **Horse**
  - Milk. 0 hour
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QV04CV01

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:
19/12/1961

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:
1/01/2009

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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