NEUROFISIN 10 U.I./ml soluzione iniettabile per bovini, equini, ovini, caprini, suini, cani, gatti

Authorised

- Oxytocin acetate

Product identification

Medicine name:
NEUROFISIN 10 U.I./ml soluzione iniettabile per bovini, equini, ovini, caprini, suini, cani, gatti

Active substance:
Oxytocin acetate

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

Route of administration:
Top-dressing use
Intramuscular use
Intravenous use

Subcutaneous use

Product details

Active substance and strength:
Oxytocin acetate
10.00 international unit(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Top-dressing use:

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

Intramuscular use:

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

- **Dog**
  - Unspecified. 0 day

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

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

- Milk. 0 hour

- **Cat**
  - Unspecified. 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

- **Dog**
  - Unspecified. 0 day

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

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

- **Cat**

- Unspecified. 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

- **Dog**
  - Unspecified. 0 day

- **Goat**
  - Milk. 0 hour

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

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

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

- Milk. 0 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH01BB02

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:
20/10/1994

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:
20/10/1994

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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