

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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