

STIMULFOS soluzione iniettabile per bovini, ovini, equini, suini, cani e gatti

Authorised

- Pyridoxine hydrochloride
- Riboflavin
- Thiamine hydrochloride
- Toldimfos
- Cyanocobalamin
- Nicotinamide
- Calcium pantothenate

Product identification

Medicine name:

STIMULFOS soluzione iniettabile per bovini, ovini, equini, suini, cani e gatti

Active substance:

Pyridoxine hydrochloride

Riboflavin

Thiamine hydrochloride

Toldimfos

Cyanocobalamin

Nicotinamide

Calcium pantothenate

Target species:

Cattle

Horse
Pig
Sheep
Dog
Cat

Route of administration:

Intramuscular use
Subcutaneous use
Intravenous use

Product details

Active substance and strength:

Pyridoxine hydrochloride
0.50 milligram(s) / 1.00 millilitre(s)
Riboflavin
1.00 milligram(s) / 1.00 millilitre(s)
Thiamine hydrochloride
20.00 milligram(s) / 1.00 millilitre(s)
Toldimfos
20.00 milligram(s) / 1.00 millilitre(s)
Cyanocobalamin
2.50 milligram(s) / 1.00 millilitre(s)
Nicotinamide
10.00 milligram(s) / 1.00 millilitre(s)
Calcium pantothenate
2.50 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 day

•

Horse

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

•

Pig

- Meat and offal. 0 day

•

Sheep

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

•

Dog

•

Cat

Subcutaneous use:

•

Cattle

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

•

Horse

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

•

Pig

- Meat and offal. 0 day

-

Sheep

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

-

Dog

-

Cat

Intravenous use:

-

Cattle

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

-

Horse

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

-

Pig

- Meat and offal. 0 day

-

Sheep

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

-

Dog

-

Cat

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA12CX90

Legal status of supply:

Veterinary medicinal product not subject to veterinary prescription

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:

Teknofarma S.r.l.

Marketing authorisation date:

9/04/1987

Manufacturing sites for batch release:

Teknofarma S.r.l.

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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