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Neovit B Complex oldatos injekció szarvasmarhák, lovak, juhok, sertések, kutyák, macskák és házityúkok részére

Authorised

- Nicotinamide
- Dexpanthenol
- Cyanocobalamin
- RIBOFLAVIN SODIUM PHOSPHATE
- Thiamine hydrochloride
- Pyridoxine hydrochloride

Product identification

Medicine name:
Neovit B Complex oldatos injekció szarvasmarhák, lovak, juhok, sertések, kutyák, macskák és házityúkok részére

Active substance:
Nicotinamide

Dexpanthenol

Cyanocobalamin

RIBOFLAVIN SODIUM PHOSPHATE

Thiamine hydrochloride

Pyridoxine hydrochloride

Target species:
Cattle
Sheep
Horse
Pig
Chicken
Dog
Cat

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Nicotinamide
25.00 milligram(s) / 1.00 millilitre(s)

Dexpanthenol
5.00 milligram(s) / 1.00 millilitre(s)

Cyanocobalamin
10.00 microgram(s) / 1.00 millilitre(s)

RIBOFLAVIN SODIUM PHOSPHATE
5.40 milligram(s) / 1.00 millilitre(s)

Thiamine hydrochloride
10.00 milligram(s) / 1.00 millilitre(s)

Pyridoxine hydrochloride
4.00 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
- **Sheep**
  - Meat and offal. 0 day
- **Horse**
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day
- **Chicken**
  - Meat and offal. 0 day

Subcutaneous use:

- **Cattle**
  - Meat and offal. 0 day
- **Sheep**
  - Meat and offal. 0 day
- **Horse**
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day
- **Chicken**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA11EA

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Hungary

Available in:
Hungary

Package description:
Available only in Hungarian
Available only in Hungarian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
KELA Kempisch Laboratorium Kela Laboratoria

Marketing authorisation date:
2/03/1998

Manufacturing sites for batch release:
KELA Kempisch Laboratorium Kela Laboratoria

Responsible authority:
Directorate Of Veterinary Medicinal Products

Authorisation number:

This information is not available for this product.

Date of authorisation status change:
2/03/1998

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet