

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:**

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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