

Beviplex vet. Injektionsvätska, lösning

Authorised

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

Product identification

Medicine name:

Beviplex vet. Injektionsvätska, lösning

Active substance:

Cyanocobalamin

Dexpanthenol

Nicotinamide

Pyridoxine hydrochloride

RIBOFLAVIN SODIUM PHOSPHATE

Thiamine hydrochloride

Target species:

Sheep

Dog

Horse

Cattle

Mink

Pig

Route of administration:

Intramuscular use

Intravenous use

Subcutaneous use

Product details

Active substance and strength:

Cyanocobalamin

5.00 microgram(s) / 1.00 millilitre(s)

Dexpanthenol

5.00 milligram(s) / 1.00 millilitre(s)

Nicotinamide

25.00 milligram(s) / 1.00 millilitre(s)

Pyridoxine hydrochloride

2.00 milligram(s) / 1.00 millilitre(s)

RIBOFLAVIN SODIUM PHOSPHATE

3.00 milligram(s) / 1.00 millilitre(s)

Thiamine hydrochloride

4.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Sheep

- Milk. 0 day

- Meat and offal. 0 day

-

Horse

- Meat and offal. 0 day

•

Cattle

- Milk. 0 day

- Meat and offal. 0 day

•

Pig

- Meat and offal. 0 day

Intravenous use:

•

Sheep

- Milk. 0 day

- Meat and offal. 0 day

•

Horse

- Meat and offal. 0 day

•

Cattle

- Milk. 0 day

- Meat and offal. 0 day

•

Pig

- Meat and offal. 0 day

Subcutaneous use:

•

Sheep

- Milk. 0 day

- Meat and offal. 0 day

-

Horse

- Meat and offal. 0 day

-

Cattle

- Milk. 0 day

- Meat and offal. 0 day

-

Pig

- 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:

Sweden

Available in:

Sweden

Package description:

Available only in [Swedish](#)

Available only in [Swedish](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Pharmaxim AB

Marketing authorisation date:

1/02/1963

Manufacturing sites for batch release:

Bela-Pharm GmbH & Co. KG

Responsible authority:

Swedish Medical Products Agency

Authorisation number:

6670

Date of authorisation status change:

1/02/1963

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

Documents

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.