

# INTROVIT B-complex, solution for injection for cattle, sheep, goats, pigs and horses

Authorised

- Choline chloride
- Biotin
- Nicotinamide
- Dexpanthenol
- Cyanocobalamin
- Pyridoxine hydrochloride
- RIBOFLAVIN SODIUM PHOSPHATE
- Thiamine hydrochloride

## Product identification

**Medicine name:**

INTROVIT B-complex, solution for injection for cattle, sheep, goats, pigs and horses

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**Active substance:**

Choline chloride

Biotin

Nicotinamide

Dexpanthenol

Cyanocobalamin

Pyridoxine hydrochloride

RIBOFLAVIN SODIUM PHOSPHATE

Thiamine hydrochloride

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**Target species:**

Cattle

Sheep

Goat

Pig

Horse

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**Route of administration:**

Subcutaneous use

Intramuscular use

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## Product details

**Active substance and strength:**

Choline chloride

10.00 milligram(s) / 1.00 millilitre(s)

Biotin

100.00 microgram(s) / 1.00 millilitre(s)

Nicotinamide

50.00 milligram(s) / 1.00 millilitre(s)

Dexpanthenol

12.50 milligram(s) / 1.00 millilitre(s)

Cyanocobalamin

20.00 microgram(s) / 1.00 millilitre(s)

Pyridoxine hydrochloride

5.00 milligram(s) / 1.00 millilitre(s)

RIBOFLAVIN SODIUM PHOSPHATE

5.00 milligram(s) / 1.00 millilitre(s)

Thiamine hydrochloride

10.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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**Withdrawal period by route of administration:**

**Subcutaneous use:**

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**Cattle**

- Meat and offal. 0 day Мляко: нула дни

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**Sheep**

- Meat and offal. 0 day Мляко: нула дни

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**Goat**

- Meat and offal. 0 day Мляко: нула дни

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**Pig**

- Meat and offal. 0 day

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**Horse**

- Meat and offal. 0 day

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QA11EA

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Bulgaria

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**Package description:**

Available only in Bulgarian

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Interchemie Werken De Adelaar B.V.

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**Marketing authorisation date:**

8/11/2009

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**Manufacturing sites for batch release:**

Interchemie Werken De Adelaar Eesti AS

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**Responsible authority:**

Bulgarian Food Safety Authority

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**Authorisation number:**

0022-2395

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**Date of authorisation status change:**

8/11/2009

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

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

Package Leaflet

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.