

# NUTRA B SOLUTION INJECTABLE POUR BOVINS OVINS EQUINS PORCINS CHIENS CHATS

Authorised

- Thiamine hydrochloride
- Ascorbic acid
- Nicotinamide
- Pyridoxine hydrochloride

## Product identification

### **Medicine name:**

NUTRA B SOLUTION INJECTABLE POUR BOVINS OVINS EQUINS PORCINS CHIENS CHATS

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### **Active substance:**

Thiamine hydrochloride

Ascorbic acid

Nicotinamide

Pyridoxine hydrochloride

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### **Target species:**

Cattle

Pig

Sheep

Equid

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**Route of administration:**

Intramuscular use

Intravenous use

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## Product details

**Active substance and strength:**

Thiamine hydrochloride

56.07 milligram(s) / 1.00 millilitre(s)

Ascorbic acid

50.00 milligram(s) / 1.00 millilitre(s)

Nicotinamide

50.00 milligram(s) / 1.00 millilitre(s)

Pyridoxine hydrochloride

60.79 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:****Intramuscular use:**

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**Cattle**

- Meat and offal. 0 day

- Milk. 0 day

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**Pig**

- Meat and offal. 0 day

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**Sheep**

- Meat and offal. 0 day

- Milk. 0 day

**Intravenous use:**

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**Cattle**

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

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**Pig**

- Meat and offal. 0 day

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**Equid**

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

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**Sheep**

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

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

QA11EB

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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:**

France

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**Package description:**

Available only in French

Available only in French

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Legal basis not covered by Directive 2001/82/EC

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**Marketing authorisation holder:**

Zoetis France

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**Marketing authorisation date:**

17/02/1992

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**Manufacturing sites for batch release:**

Zoetis Manufacturing & Research Spain S.L.

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/9202376 9/1992

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**Date of authorisation status change:**

17/02/2012

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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 and Labelling

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