

# KYNOSELEN SOLUTION INJECTABLE

Authorised

- Sodium selenite
- Potassium aspartate hemihydrate
- Magnesium aspartate dihydrate

## Product identification

**Medicine name:**

KYNOSELEN SOLUTION INJECTABLE

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**Active substance:**

Sodium selenite

Potassium aspartate hemihydrate

Magnesium aspartate dihydrate

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**Target species:**

Cattle

Pig

Cat

Sheep

Dog

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**Route of administration:**

Intramuscular use

Subcutaneous use

Intravenous use

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## Product details

### **Active substance and strength:**

Sodium selenite

0.50 milligram(s) / 1.00 millilitre(s)

Potassium aspartate hemihydrate

10.00 milligram(s) / 1.00 millilitre(s)

Magnesium aspartate dihydrate

15.00 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:**

#### **Intramuscular use:**

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#### **Cattle**

- Meat and offal. 4 day

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#### **Pig**

- Meat and offal. 4 day

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#### **Sheep**

- Meat and offal. 4 day

#### **Subcutaneous use:**

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#### **Cattle**

- Meat and offal. 4 day

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#### **Pig**

- Meat and offal. 4 day

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**Sheep**

- Meat and offal. 4 day

**Intravenous use:**

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**Cattle**

- Meat and offal. 4 day

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**Pig**

- Meat and offal. 4 day

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**Sheep**

- Meat and offal. 4 day

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

QA12CE99

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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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**Available in:**

France

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**Package description:**

Available only in French

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:**

Vetoquinol S.A.

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**Marketing authorisation date:**

30/06/1992

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**Manufacturing sites for batch release:**

Vetoquinol S.A.

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/1233902 0/1992

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**Date of authorisation status change:**

27/11/2018

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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

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Package Leaflet and Labelling

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