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# KYNOSELEN SOLUTION INJECTABLE

Authorised

- Sodium selenite
- Potassium aspartate hemihydrate
- Magnesium aspartate dihydrate

## Product identification

**Medicine name:**
KYNOSELEN SOLUTION INJECTABLE

**Active substance:**
Sodium selenite
Potassium aspartate hemihydrate
Magnesium aspartate dihydrate

**Target species:**
Cattle
Pig
Cat
Sheep
Dog

**Route of administration:**
Intramuscular use
Subcutaneous use
Intravenous use

## 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:**

- **Cattle**
  - Meat and offal. 4 day

- **Pig**
  - Meat and offal. 4 day

- **Sheep**
  - Meat and offal. 4 day

**Subcutaneous use:**

- **Cattle**
  - Meat and offal. 4 day

- **Pig**
  - Meat and offal. 4 day

- **Sheep**
  - Meat and offal. 4 day

**Intravenous use:**

- **Cattle**
  - Meat and offal. 4 day

- **Pig**
  - Meat and offal. 4 day

- **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

**Legal basis of product authorisation:**
Legal basis not covered by Directive 2001/82/EC

**Marketing authorisation holder:**
Vetoquinol S.A.

**Marketing authorisation date:**
30/06/1992

**Manufacturing sites for batch release:**
Vetoquinol S.A.

**Responsible authority:**
French Agency For Food, Environmental And Occupational Health & Safety

**Authorisation number:**
FR/V/1233902 0/1992

**Date of authorisation status change:**
27/11/2018

To consult adverse reactions on veterinary medicinal products please go to 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.