

# Surcalce injekció A.U.V.

Authorised

- Magnesium hypophosphite hexahydrate
- Calcium acetate
- Calcium gluconate

## Product identification

**Medicine name:**

Surcalce injekció A.U.V.

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**Active substance:**

Magnesium hypophosphite hexahydrate

Calcium acetate

Calcium gluconate

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**Target species:**

Cattle

Sheep

Goat

Pig

Horse

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**Route of administration:**

Intravenous use

Subcutaneous use

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## Product details

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

Magnesium hypophosphite hexahydrate

30.00 milligram(s) / 1.00 millilitre(s)

Calcium acetate

37.00 milligram(s) / 1.00 millilitre(s)

Calcium gluconate

465.00 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:**

#### **Intravenous use:**

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#### **Cattle**

- Milk. 0 day

- Meat and offal. 0 day

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#### **Sheep**

- Milk. 0 day

- Meat and offal. 0 day

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#### **Goat**

- Milk. 0 day

- Meat and offal. 0 day

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#### **Pig**

- Meat and offal. 0 day

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**Horse**

- Meat and offal. 0 day

**Subcutaneous use:**

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**Cattle**

- Milk. 0 day

- Meat and offal. 0 day

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**Sheep**

- Milk. 0 day

- Meat and offal. 0 day

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**Goat**

- Milk. 0 day

- Meat and offal. 0 day

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**Pig**

- Meat and offal. 0 day

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

QA12AA20

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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:**

Hungary

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**Package description:**

Available only in Hungarian

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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:**

V.M.D.

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**Marketing authorisation date:**

3/06/1999

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**Manufacturing sites for batch release:**

Alvetra U. Werfft GmbH

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**Responsible authority:**

Directorate Of Veterinary Medicinal Products

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

3/06/1999

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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

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