

# Synulox Lactating Cow

## Intramammary suspension.

Authorised

- Prednisolone
- Potassium clavulanate
- Amoxicillin trihydrate

## Product identification

**Medicine name:**

Synulox Lactating Cow Intramammary suspension.

---

**Active substance:**

Prednisolone

Potassium clavulanate

Amoxicillin trihydrate

---

**Target species:**

Cattle

---

**Route of administration:**

Intramammary use

---

## Product details

**Active substance and strength:**

Prednisolone

10.00 milligram(s) / 1.00 Syringe

Potassium clavulanate

59.56 milligram(s) / 1.00 Syringe

Amoxicillin trihydrate

230.00 milligram(s) / 1.00 Syringe

---

**Pharmaceutical form:**

Intramammary suspension

---

**Withdrawal period by route of administration:**

**Intramammary use:**

- 

**Cattle**

- Milk. 60 hour

---

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

QJ51RV01

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Slovenia

---

**Available in:**

Slovenia

---

**Package description:**

Low density polyethylene syringes packed in cartons containing 12 syringes.

Low density polyethylene syringes packed in cartons containing 24 syringes.

Low density polyethylene syringes packed in cartons containing 300 syringes.

**\*\*HISTORICAL\*\*** Low density polyethylene syringes packed in cartons containing 3 syringes.

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

---

**Marketing authorisation holder:**

Zoetis Belgium

---

**Marketing authorisation date:**

31/01/2001

---

**Manufacturing sites for batch release:**

Haupt Pharma Latina S.r.l.

---

**Responsible authority:**

Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

---

**Authorisation number:**

NP/V/0326/001

---

**Date of authorisation status change:**

31/01/2001

---

**Reference member state:**

Ireland

---

**Procedure number:**

IE/V/0605/001

---

**Concerned member states:**

Austria Bulgaria Cyprus Czechia France Greece Hungary Italy Latvia  
Lithuania Netherlands Norway Poland Portugal Romania Slovakia Slovenia  
Spain United Kingdom (Northern Ireland)

---

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