

# Combiclav Intramammary Suspension for Lactating Cows

Authorised

- Prednisolone
- Potassium clavulanate
- Amoxicillin trihydrate

## Product identification

**Medicine name:**

Combiclav Intramammary Suspension for Lactating Cows  
Combiclav intramamarna suspenzija za krave v laktaciji

---

**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

229.61 milligram(s) / 1.00 Syringe

---

**Pharmaceutical form:**

Intramammary suspension

---

**Withdrawal period by route of administration:**

**Intramammary use:**

- 

**Cattle**

- Meat and offal. 7 day

- Meat and offal. 42 day

Combined therapy: this product (intramammary use) and Noroclav Injection for Cattle and Dogs

- Milk. 84 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

---

**Package description:**

Single dose 3g white LDPE syringes with a white LDPE dual push-fit cap. Buckets of 120 syringes, including 120 individually wrapped teat cleaning towels containing isopropyl alcohol.

Single dose 3g white LDPE syringes with a white LDPE dual push-fit cap. Cartons of 24 syringes, including 24 individually wrapped teat cleaning towels containing isopropyl alcohol.

Single dose 3g white LDPE syringes with a white LDPE dual push-fit cap. Cartons of 12 syringes, including 12 individually wrapped teat cleaning towels containing isopropyl alcohol.

Single dose 3g white LDPE syringes with a white LDPE dual push-fit cap. Cartons of 3 syringes, including 3 individually wrapped teat cleaning towels containing isopropyl alcohol.

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

---

**Marketing authorisation holder:**

Norbrook Laboratories (Ireland) Limited

---

**Marketing authorisation date:**

7/11/2018

---

**Manufacturing sites for batch release:**

Norbrook Manufacturing Limited

Norbrook Laboratories Limited

---

**Responsible authority:**

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

---

**Authorisation number:**

DC/V/0640/001

---

**Date of authorisation status change:**

7/11/2018

---

**Reference member state:**

Ireland

---

**Procedure number:**

IE/V/0535/001

---

**Concerned member states:**

Belgium Bulgaria Croatia Czechia Hungary Italy Portugal 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

English (PDF)

Published on: 29/12/2024

[Download](#)

Combined File of all Documents

Package Leaflet

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

## Labelling

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