# Combiclav Intramammary Suspension for Lactating Cows

Authorised

- Prednisolone
- Amoxicillin trihydrate
- Potassium clavulanate

## Product identification

**Medicine name:**
Combiclav Intramammary Suspension for Lactating Cows
Combiclav Suspensie voor intramammair gebruik
Combiclav Suspension intramammaire
Combiclav Suspension zur intramammären Anwendung

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**Active substance:**
Prednisolone

Amoxicillin trihydrate

Potassium clavulanate

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**Target species:**
Cattle

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**Route of administration:**
Intramammary use

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## Product details

**Active substance and strength:**

Prednisolone
10.00 milligram(s) / 1.00 Syringe

Amoxicillin trihydrate
229.61 milligram(s) / 1.00 Syringe

Potassium clavulanate
59.56 milligram(s) / 1.00 Syringe

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**Pharmaceutical form:**
Intramammary suspension

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**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

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QJ51RV01

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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:**
Belgium

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**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:**
12/12/2018

**Manufacturing sites for batch release:**
Norbrook Laboratories Limited
Norbrook Manufacturing Ltd

**Responsible authority:**
Federal Agency For Medicines And Health Products

**Authorisation number:**
BE-V537342

**Date of authorisation status change:**
12/12/2018

**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

## Documents

Summary of Product Characteristics

English (PDF)
Published on: 29/12/2024
Download

Package Leaflet

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

Combined File of all Documents

Labelling

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

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