

Combiclav Intramammary Suspension for Lactating Cows

Authorised

- Prednisolone
- Potassium clavulanate
- Amoxicillin trihydrate

Product identification

Medicine name:

Combiclav Intramammary Suspension for Lactating Cows

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
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Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ51RV01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Croatia

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:

6/08/2019

Manufacturing sites for batch release:

Norbrook Manufacturing Limited

Norbrook Laboratories Limited

Responsible authority:

Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:

UP/I-322-05/19-01/409

Date of authorisation status change:

25/04/2025

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

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Combined File of all Documents