Combiclav Intramammary Suspension for Lactating Cows

Authorised

- Prednisolone
- Potassium clavulanate
- Amoxicillin trihydrate

Product identification

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

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

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Authorisation status:

Valid

Authorised in:

Belgium

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 Manufacturing Limited
Norbrook Laboratories Limited

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
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Package Leaflet

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Combined File of all Documents

Labelling

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