

# Ubrolexin intramammary suspension for lactating dairy COWS

Authorised

- Cefalexin monohydrate
- KANAMYCIN MONOSULPHATE

## Product identification

**Medicine name:**

Ubrolexin intramammary suspension for lactating dairy cows

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**Active substance:**

Cefalexin monohydrate

KANAMYCIN MONOSULPHATE

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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:**

Cefalexin monohydrate

210.36 milligram(s) / 1.00 Syringe

KANAMYCIN MONOSULPHATE

120247.00 international unit(s) / 1.00 Syringe

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**Pharmaceutical form:**

Intramammary suspension

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**Withdrawal period by route of administration:**

**Intramammary use:**

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**Cattle**

- Meat and offal. 10 day
  - Milk. 5 day
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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ51RD01

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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:**

Germany

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**Available in:**

Germany

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**Package description:**

Cardboard box with 20 single use intramammary syringes and 20 teat wipes (containing isopropanol 70%). Each 10 g syringe contains 12 ml intramammary suspension and consists of a barrel with plunger and sealed sterile tip, all made of low density polyethylene.

Cardboard box with 10 single use intramammary syringes and 10 teat wipes (containing isopropanol 70%). Each 10 g syringe contains 12 ml intramammary suspension and consists of a barrel with plunger and sealed sterile tip, all made of low density polyethylene.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Well-established use application (Article 13a of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Boehringer Ingelheim Vetmedica GmbH

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**Marketing authorisation date:**

1/08/2008

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**Manufacturing sites for batch release:**

Univet Limited

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**Responsible authority:**

Federal Office Of Consumer Protection And Food Safety

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**Authorisation number:**

401157.00.00

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**Date of authorisation status change:**

22/11/2012

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0221/001

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**Concerned member states:**

Austria Belgium Cyprus Czechia Estonia France Germany Greece Hungary  
Italy Latvia Lithuania Luxembourg Netherlands Poland Portugal Romania  
Slovakia Slovenia Spain United Kingdom (Northern Ireland)

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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

Combined File of all Documents

English (PDF)

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