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Ubrolexin intramammary suspension for lactating dairy cows

Authorised

- Cefalexin monohydrate
- KANAMYCIN MONOSULPHATE

Product identification

Medicine name:
Ubrolexin intramammary suspension for lactating dairy cows

Active substance:
Cefalexin monohydrate
KANAMYCIN MONOSULPHATE

Target species:
Cattle

Route of administration:
Intramammary use

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

Pharmaceutical form:
Intramammary suspension

Withdrawal period by route of administration:
Intramammary use:

- **Cattle**
  - Meat and offal. 10 day
  - Milk. 5 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51RD01

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Italy

Package description:
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.
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.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:
22/10/2008

Manufacturing sites for batch release:
Univet Limited

Responsible authority:
Ministry Of Health

Authorisation number:
104073

Date of authorisation status change:
26/10/2012

Reference member state:
Ireland

Procedure number:
IE/V/0221/001

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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Documents

Combined File of all Documents

English (PDF)
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Summary of Product Characteristics