Terrexine DC 250 mg Intramammary Suspension for Dry Cows

Authorised

- Cefalonium dihydrate

Product identification

Medicine name:
Terrexine DC 250 mg Intramammary Suspension for Dry Cows
Arentor DC 250 mg suspensie voor intramammair gebruik bij droogstaande koeien

Active substance:
Cefalonium dihydrate

Target species:
Cattle

Route of administration:
Intramammary use

Product details

Active substance and strength:
Cefalonium dihydrate
269.63 milligram(s) / 1.00 Syringe

Pharmaceutical form:
Intramammary suspension

Withdrawal period by route of administration:
Intramammary use:

- **Cattle**
  - Meat and offal. 21 day
  - Milk. 96 hour

96 hours after calving if the dry period is longer than 54 days. 58 days following the treatment if the dry period is less than or equal to 54 days

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51DB90

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

Authorisation status:
Valid

Authorised in:
Netherlands

Package description:
A 3g coloured low density polyethylene intramammary syringe with a coloured low density polyethylene dual capPack sizes:Buckets of 120 intramammary syringes and 120 individually wrapped cleaning towels containing isopropyl alcohol.
A 3g coloured low density polyethylene intramammary syringe with a coloured low density polyethylene dual capPack sizes:Cartons of 20 intramammary syringes and 20 individually wrapped 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:
Univet Limited

Marketing authorisation date:
2/07/2019

Manufacturing sites for batch release:
Univet Limited

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 124023

Date of authorisation status change:
27/01/2022

Reference member state:
Ireland

Procedure number:
IE/V/0522/001

Concerned member states:
Netherlands

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents

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

Summary of Product Characteristics