BOVICEF DC 250 mg Intramammary Suspension for Cattle

Not authorised

- Cefalonium dihydrate

Product identification

Medicine name:
BOVICEF DC 250 mg Intramammary Suspension for Cattle

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51DB90

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

Authorisation status:
Surrendered

Authorised in:
Hungary

Package description:
Single dose 3g white polyethylene intramammary syringe with a polyethylene cap. Pack sizes: 120 intramammary syringes in a bucket. All pack sizes contain cleaning towels.
Single dose 3g white polyethylene intramammary syringe with a polyethylene cap. Pack sizes: 24 intramammary syringes in a carton. All pack sizes contain cleaning towels.

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:
Zoetis Hungary Kft.

Marketing authorisation date:
17/07/2015

Manufacturing sites for batch release:
Cross Vetpharm Group Limited

Responsible authority:
Directorate Of Veterinary Medicinal Products

Authorisation number:
3661/X/15 NÉBIH ÁTI

Date of authorisation status change:
6/12/2022

Reference member state:
Ireland

Procedure number:
IE/V/0530/001

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

Documents

Summary of Product Characteristics