Paraclav Intramammary Suspension for Lactating Cows

Not authorised

- Amoxicillin trihydrate
- Potassium clavulanate
- Prednisolone

Product identification

Medicine name:
Paraclav Intramammary Suspension for Lactating Cows

Active substance:
Amoxicillin trihydrate

Potassium clavulanate

Prednisolone

Target species:
Cattle (lactating cow)

Route of administration:
Intramammary use

Product details

Active substance and strength:
Amoxicillin trihydrate

229.60 milligram(s) / 1.00 Applicator

Potassium clavulanate
59.60 milligram(s) / 1.00 Applicator

Prednisolone
10.00 milligram(s) / 1.00 Applicator

Pharmaceutical form:
Intramammary suspension

Withdrawal period by route of administration:
Intramammary use:

- **Cattle (lactating cow)**
  - Meat and offal. 7 day
  - Milk. 84 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51RV01

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

---

Authorisation status:
Surrendered

Authorised in:
United Kingdom (Northern Ireland)

Package description:
(ID4) 360 gram(s): unspecified outer container with 120 Applicator (low-density polyethylene) each with 3 gram(s)
(ID3) 72 gram(s): unspecified outer container with 24 Applicator (low-density polyethylene) each with 3 gram(s)
(ID2) 36 gram(s): unspecified outer container with 12 Applicator (low-density polyethylene) each with 3 gram(s)
(ID1) 9 gram(s): unspecified outer container with 3 Applicator (low-density polyethylene) each with 3 gram(s)

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 Limited

Marketing authorisation date:
3/10/2018

Manufacturing sites for batch release:
Norbrook Laboratories (Ireland) Limited
Norbrook Laboratories Limited

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 02000/4426

Date of authorisation status change:
11/07/2022

Reference member state:
Germany

Procedure number:
DE/V/0328/001

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

Documents

Summary of Product Characteristics

2402467-paren-20190329.rtf