Orbenin Dry Cow 500 mg Intramammary Suspension

Authorised

- Cloxacillin hemibenzathine

Product identification

Medicine name:
Orbenin Dry Cow 500 mg Intramammary Suspension

Active substance:
Cloxacillin hemibenzathine

Target species:
Cattle

Route of administration:
Intramammary use

Product details

Active substance and strength:
Cloxacillin hemibenzathine
1275.64 milligram(s) / 1.00 Syringe

Pharmaceutical form:
Intramammary suspension

Withdrawal period by route of administration:

Intramammary use:

- **Cattle**
  - Meat and offal. 28 day
  - Milk. 39 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51CF02

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

Authorisation status:
Valid

Authorised in:
Ireland

Package description:
White low density polyethylene intramammary syringe barrel containing 3 g of suspension with plunger. The closure is a white low density polyethylene push-fit combined dual nozzle and cap. Pack containing 24 syringes.
White low density polyethylene intramammary syringe barrel containing 3 g of suspension with plunger. The closure is a white low density polyethylene push-fit combined dual nozzle and cap. Pack containing 120 syringes.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Zoetis Belgium S.A.

Marketing authorisation date:
1/10/1997

Manufacturing sites for batch release:
Haupt Pharma Latina S.r.l.

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10387/045/001

Date of authorisation status change:
1/10/1997

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

Documents

Summary of Product Characteristics