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Bovaclox DC Xtra Intramammary Suspension

Authorised

- Ampicillin trihydrate
- Cloxacillin hemibenzathine

Product identification

Medicine name:

Bovaclox DC Xtra Intramammary Suspension

Active substance:

Ampicillin trihydrate

Cloxacillin hemibenzathine

Target species:

Cattle

Route of administration:

Intramammary use

Product details

Active substance and strength:

Ampicillin trihydrate 346.45 milligram(s) / 1.00 Syringe

Cloxacillin hemibenzathine

Pharmaceutical form:

Intramammary suspension

Withdrawal period by route of administration: Intramammary use:

Cattle

- Meat and offal. 28 day
- Milk. 156 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ51CR50

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Available in:

Ireland

Package description:

Bovaclox DC Xtra is presented in high density polyethylene syringes (with high density polyethylene caps) containing 5.4g of an off-white non aqueous suspension. These intramammary syringes will be supplied in buckets of 120 syringes Bovaclox DC Xtra is presented in high density polyethylene syringes (with high density polyethylene caps) containing 5.4g of an off-white non aqueous suspension. These intramammary syringes will be supplied in cartons of 24 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:

Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:

11/06/1993

Manufacturing sites for batch release:

Norbrook Laboratories Limited Norbrook Manufacturing Limited

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA22664/038/001

Date of authorisation status change:

11/06/1993

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

Documents

Summary of Product Characteristics