Albiotic 330 mg/100 mg Intramammary Solution

Authorised

- NEOMYCIN SULFATE
- Lincomycin hydrochloride

Product identification

Medicine name:
Albiotic 330 mg/100 mg Intramammary Solution

Active substance:
NEOMYCIN SULFATE
Lincomycin hydrochloride

Target species:
Cattle

Route of administration:
Intramammary use

Product details

Active substance and strength:
NEOMYCIN SULFATE
115.96 milligram(s) / 1.00 Syringe
Lincomycin hydrochloride

359.63 milligram(s) / 1.00 Syringe

Pharmaceutical form:
Intramammary solution

Withdrawal period by route of administration:
Intramammary use:

- **Cattle**
  - Meat and offal. 48 hour
  - Milk. 72 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51RF03

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

Authorisation status:
Valid

Authorised in:
Ireland

Package description:
Sterile aqueous solution in 10 ml high density polyethylene syringes with a fixed cannula (plastets), packaged as 100plastets in an outer cardboard box also containing alcohol pads for teat disinfection. Cap LDPE, brombutyl rubber stopper. Sterile aqueous solution in 10 ml high density polyethylene syringe with a fixed cannula (plastet), packaged as 1 plastet in an outer cardboard box also containing alcohol pad for teat disinfection. Cap LDPE, brombutyl rubber stopper.

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:
HuVepharma

Marketing authorisation date:
31/03/2006

Manufacturing sites for batch release:
Biovet AD

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10782/028/001

Date of authorisation status change:
31/03/2006

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

Documents

Summary of Product Characteristics