Bimoxyvet LA 150 mg/ml, Suspension for Injection for cattle, sheep and pigs

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:
Bimoxyvet LA 150 mg/ml, Suspension for Injection for cattle, sheep and pigs

Active substance:
Amoxicillin trihydrate

Target species:
Cattle
Sheep
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Amoxicillin trihydrate
172.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat and offal. 18 day
  - Milk. 72 hour
- **Sheep**
  - Meat and offal. 21 day
- **Pig**
  - Meat and offal. 21 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CA04

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

Authorisation status:
Valid

Authorised in:
Ireland

Package description:
Type I clear, glass vial sealed with a Type I bromobutyl rubber stopper and capped with aluminium overseal.Pack size:Vial of 250 ml
Type I clear, glass vial sealed with a Type I bromobutyl rubber stopper and capped with aluminium overseal.Pack size:Vial of 100 ml
Clear Polyethylene terephthalate (PET) vials with a Type I chlorobutyl stopper and aluminium cap with plastic flip off sealPack size:Vial of 100 ml

Clear Polyethylene terephthalate (PET) vials with a Type I chlorobutyl stopper and aluminium cap with plastic flip off sealPack size:Vial of 250 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Laboratorios Maymo S.A.U.

Marketing authorisation date:
10/02/2017

Manufacturing sites for batch release:
Laboratorios Syva S.A.

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10436/002/001

Date of authorisation status change:
10/02/2017

Reference member state:
Ireland

Procedure number:
IE/V/0362/001

Concerned member states:
Austria Belgium Denmark Finland France Italy Poland Portugal Romania Spain

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

Documents

Summary of Product Characteristics