

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

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:

Bimoxivet 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:

Denmark

Package description:

Type I clear, glass vial sealed with a Type I bromobutyl rubber stopper and capped with aluminium overseal.Pack size:Vial of 100 ml

Type I clear, glass vial sealed with a Type I bromobutyl rubber stopper and capped with aluminium overseal.Pack size:Vial of 250 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:

15/02/2017

Manufacturing sites for batch release:

Laboratorios Syva S.A.

Responsible authority:

Danish Medicines Agency

Authorisation number:

57386

Date of authorisation status change:

15/02/2017

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

English (PDF)

Published on: 5/07/2026

[Download](#)

Combined File of all Documents