Betamox LA 150 mg/ml Suspension for Injection

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:
Betamox LA 150 mg/ml Suspension for Injection

Active substance:
Amoxicillin trihydrate

Target species:
Cattle
Dog
Sheep
Cat
Pig

Route of administration:
Intramuscular use
Subcutaneous use

Product details

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

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. 39 day
  - Milk. 108 hour
- **Dog**
- **Sheep**
  - Meat and offal. 29 day
- **Cat**
- **Pig**
  - Meat and offal. 42 day

Subcutaneous use:

- **Dog**
- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CA01

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

---

Authorisation status:
Valid

Authorised in:

Ireland

Available in:
Ireland

Package description:
The veterinary medicinal product is supplied in 50 ml Type II glass vial sealed with nitryl rubber bung and aluminium overseal OR 50 ml clear polyethylene terephthalate (PET) vial sealed with nitryl rubber bung and aluminium overseal.
The veterinary medicinal product is supplied in 100 ml Type II glass vial sealed with nitryl rubber bung and aluminium overseal OR 100 ml clear polyethylene terephthalate (PET) vial sealed with nitryl rubber bung and aluminium overseal.
The veterinary medicinal product is supplied in 250 ml clear polyethylene terephthalate (PET) vial sealed with nitryl rubber bung and aluminium overseal.

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:
1/10/1987

Manufacturing sites for batch release:
Norbrook Laboratories Limited
Norbrook Manufacturing Limited

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA22664/010/001

Date of authorisation status change:
1/10/1987

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

Documents

Summary of Product Characteristics