Alfamox LA 150 mg/ml Suspension for Injection

Authorised

- Amoxicillin trihydrate

Product identification

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

Active substance:
Amoxicillin trihydrate

Target species:
Cattle
Sheep
Pig

Route of administration:
Intramuscular 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. 28 day
  - Milk. 5 day
- **Sheep**
  - Meat and offal. 28 day
  - Milk. 5 day
- **Pig**
  - Meat and offal. 28 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:

12 x 100 ml clear, type II, glass vials closed with a nitryl stopper and aluminium seal.
100 ml clear, type II, glass vial closed with a nitryl stopper and aluminium seal.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Informed consent application (Article 13c of Directive No 2001/82/EC)

Marketing authorisation holder:
Alfa Med Limited

Marketing authorisation date:
22/01/1999

Manufacturing sites for batch release:
Alfa Med Limited

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10894/004/001

Date of authorisation status change:
22/01/1999

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

Documents

Summary of Product Characteristics