Lamoxsan 150 mg/ml suspension for injection for cattle and pigs

Authorised

- Amoxicillin

Product identification

Medicine name:
Lamoxsan 150 mg/ml suspension for injection for cattle and pigs

Active substance:
Amoxicillin

Target species:
Cattle
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Amoxicillin
150.00 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
- **Pig**
  - Meat and offal. 20 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CA04

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

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Authorisation status:
Valid

Authorised in:
Belgium

Package description:
Clear PET vial of 100 ml closed with type I laminated chlorobutyl rubber stopper and aluminium cap in a carton box.
Clear PET vial of 250 ml closed with type I laminated chlorobutyl rubber stopper and aluminium cap in a carton box.
Clear type II glass vial of 100 ml closed with type I laminated chlorobutyl rubber stopper and aluminium cap in a carton box.
Clear type II glass vial of 250 ml closed with type I laminated chlorobutyl rubber stopper and aluminium cap in a carton box.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:
Alfasan Nederland B.V.

Marketing authorisation date:
13/06/2024

Manufacturing sites for batch release:
Alfasan Nederland B.V.

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
13/06/2024

Reference member state:
Netherlands

Procedure number:
NL/V/0391/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland France Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg Norway Poland Portugal Romania Slovakia Slovenia Spain Sweden United Kingdom (Northern Ireland)

Generic of:
600000004401

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

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents

NLV0391001DC_DCP_Lamoxsan final PuAR.pdf