Kelamoxil LA 150 mg/ml suspension for injection for cattle and pig

Not authorised

- Amoxicillin

Product identification

Medicine name:
Kelamoxil LA 150 mg/ml suspension for injection for cattle and pig
Kelamoxil LA 150 mg/ml suspenzija za injiciranje za govedo in prašiče

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

Authorisation status:
Surrendered

Authorised in:
Slovenia

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

Marketing authorisation date:
18/08/2023

Manufacturing sites for batch release:
KELA Kempisch Laboratorium Kela Laboratoria

Responsible authority:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:
DC/V/0786/001

Date of authorisation status change:
21/11/2025

Reference member state:
Netherlands

Procedure number:
NL/V/0390/001

Generic of:
600000004401

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 22/08/2023
Updated on: 23/08/2023
Download

Package Leaflet

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 and Labelling

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