

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

Authorised

- Amoxicillin

## Product identification

### **Medicine name:**

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

Kelamoxil LA 150 mg/ml szuszpenziós injekció szarvarmarhák és sertések részére

### **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

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**Withdrawal period by route of administration:****Intramuscular use:**

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**Cattle**

- Meat and offal. 18 day
- Milk. 72 hour

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**Pig**

- Meat and offal. 20 day

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ01CA04

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Hungary

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**Available in:**

Hungary

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**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.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 18 of Regulation (EU) 2019/6)

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**Marketing authorisation holder:**

KELA Kempisch Laboratorium Kela Laboratoria

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**Marketing authorisation date:**

9/10/2023

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**Manufacturing sites for batch release:**

KELA Kempisch Laboratorium Kela Laboratoria

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**Responsible authority:**

Directorate Of Veterinary Medicinal Products

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

9/10/2023

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0390/001

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**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)

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**Generic of:**

600000004401

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

Package Leaflet and Labelling

NLV0390001DC\_DCP\_Kelamoxil LA Final PuAR.pdf