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

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:

Valid

Authorised in:

Luxembourg

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:

5/09/2023

Manufacturing sites for batch release:

Kela - Kempisch Laboratorium - Kela Laboratoria

Responsible authority:

Ministere De La Sante Division De La Pharmacie Et Des Medicaments

Authorisation number:

V188/05/09/2023

Date of authorisation status change:

5/09/2023

Reference member state:

Netherlands

Procedure number:

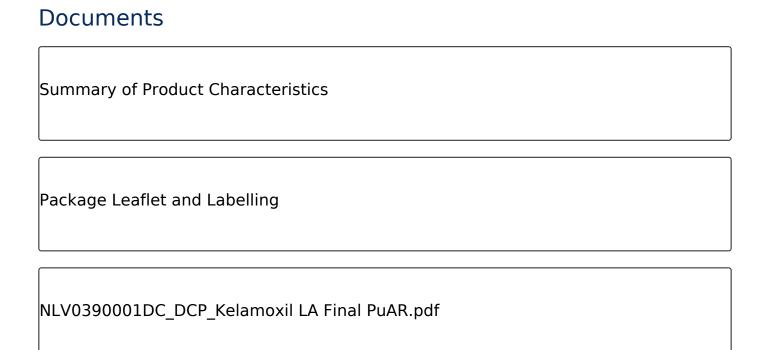
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Concerned member states:

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)

This information is not available for this product.

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



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