CITRAMOX L.A. 150 mg/ml suspension for injection for cattle and pigs

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:
CITRAMOX L.A. 150 mg/ml suspension for injection for cattle and pigs
CITRAMOX L.A. 150 MG/ML SUSPENSION INJECTABLE POUR BOVINS ET PORCINS

Active substance:
Amoxicillin trihydrate

Target species:
Cattle
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Amoxicillin trihydrate
172.20 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Milk. 3 day
  - Meat and offal. 18 day
- **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:
France

Package description:
Cardboard box with multi-layer polypropylene_ethylene vinyl alcohol_ polypropylene vial closed with bromobutyl rubber stopper and aluminium and plastic flip capsule, of capacity 250 ml
Cardboard box with multi-layer polypropylene_ethylene vinyl alcohol_ polypropylene vial closed with bromobutyl rubber stopper and aluminium and plastic flip capsule, of capacity 100 ml

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Laboratorios Karizoo S.A.

Marketing authorisation date:
13/08/2020

Manufacturing sites for batch release:
Przedsiebiorstwo Wielobranzowe Vet-Agro Sp. z o.o.

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/7104511 9/2020

Date of authorisation status change:
13/08/2020

Reference member state:
Netherlands

Procedure number:
NL/V/0323/001

Concerned member states:
Belgium Bulgaria Croatia Cyprus Czechia Denmark France Germany Greece Hungary Ireland Italy Lithuania Poland Portugal Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland)

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.

Package Leaflet and Labelling

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