

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

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:**

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Cattle

- Milk. 3 day
- Meat and offal. 18 day

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Pig

- Meat and offal. 20 day
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Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01CA04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Romania

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:

Przedsiębiorstwo Wielobranzowe Vet-Agro Sp. z o.o.

Marketing authorisation date:

12/08/2020

Manufacturing sites for batch release:

Przedsiębiorstwo Wielobranzowe Vet-Agro Sp. z o.o.

Responsible authority:

Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:

250117

Date of authorisation status change:

12/08/2025

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

Combined File of all Documents

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