

Vetrimoxin LA, 150 mg/ml, injekcinė suspensija galvijams ir kiaulėms

Authorised

- Amoxicillin

Product identification

Medicine name:

Vetrimoxin LA, 150 mg/ml, injekcinė suspensija galvijams ir kiaulėms

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. 14 day

- Milk. 3 day

-

Pig

- Meat and offal. 12 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01CA04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Lithuania

Available in:

Lithuania

Package description:

Available only in [Lithuanian](#)

Available only in [Lithuanian](#)

Available only in [Lithuanian](#)

Available only in [Lithuanian](#)

Available only in [Lithuanian](#)

Available only in [Lithuanian](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder:

Ceva Sante Animale

Marketing authorisation date:

12/11/2002

Manufacturing sites for batch release:

Ceva Sante Animale

Vetem S.p.A.

Responsible authority:

State Food And Veterinary Service

Authorisation number:

LT/2/02/1488/001-008

Date of authorisation status change:

2/06/2025

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

Documents

Combined File of all Documents

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