

Trymox LA 150 mg/ml Suspension for Injection for Cattle, Sheep, Pigs, Dogs and Cats

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:

Trymox LA 150 mg/ml Suspension for Injection for Cattle, Sheep, Pigs, Dogs and Cats

Active substance:

Amoxicillin trihydrate

Target species:

Cattle

Dog

Sheep

Cat

Pig

Route of administration:

Intramuscular use

Subcutaneous use

Product details

Active substance and strength:

Amoxicillin trihydrate

172.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Cattle

- Meat and offal. 39 day

- Milk. 108 hour

-

Sheep

- Meat and offal. 29 day

-

Pig

- Meat and offal. 42 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01CA04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Norway

Package description:

100 ml clear polyethylene terephthalate vial sealed with nitrile bung and aluminium overseal. Pack size: 12 x 100ml vials

250 ml clear polyethylene terephthalate vial sealed with nitrile bung and aluminium overseal. Pack size: 6 x 250 ml vial

100 ml clear polyethylene terephthalate vial sealed with nitrile bung and aluminium overseal. Pack size: 1 x 100ml vial

250 ml clear polyethylene terephthalate vial sealed with nitrile bung and aluminium overseal. Pack size: 1 x 250 ml vial

250 ml clear, colourless Type II glass vial, closed with nitrile rubber bung and aluminium overseal. Pack Sizes: 6 x 250 ml vials

100 ml clear, colourless Type II glass vial, closed with nitrile rubber bung and aluminium overseal. Pack Sizes: 12 x 100 ml vials

50 ml clear, colourless Type II glass vial, closed with nitrile rubber bung and aluminium overseal. Pack Sizes: 12 x 50 ml vials

250 ml clear, colourless Type II glass vial, closed with nitrile rubber bung and aluminium overseal. Pack size: 1 x 250 ml vial

100 ml clear, colourless Type II glass vial, closed with nitrile rubber bung and aluminium overseal. Pack size: 1 x 100 ml vial

50 ml clear, colourless Type II glass vial, closed with nitrile rubber bung and aluminium overseal. Pack Sizes: 1 x 50 ml vial

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:

Univet Limited

Marketing authorisation date:

21/02/2019

Manufacturing sites for batch release:

Univet Limited

Responsible authority:

Norwegian Medical Products Agency

Authorisation number:

18-12222

Date of authorisation status change:

21/02/2019

Reference member state:

Ireland

Procedure number:

IE/V/0608/001

Concerned member states:

Austria Belgium Bulgaria Croatia Cyprus Denmark Estonia Finland France
Germany Greece Hungary Italy Latvia Liechtenstein Lithuania Netherlands
Norway Poland Portugal Romania Slovakia Slovenia Sweden
United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 26/01/2026

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Package Leaflet

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