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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
  - Milk. no withdrawal period

Not authorised for use in sheep producing milk for human consumption.

- **Pig**
  - Meat and offal. 42 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CA04

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

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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 Czechia 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
Updated on: 14/03/2026

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Package Leaflet

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