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
Trymox LA 150 mg/ml Injektionssuspension für Rinder, Schafe, Schweine, Hunde, Katzen

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:
Austria

Package description:

Available only in German
Available only in German
Available only in German
Available only in German
Available only in German
Available only in German
250 ml clear polyethylene terephthalate vial sealed with nitrile bung and aluminium overseal. Pack size: 1 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: 6 x 250 ml vial
100 ml clear polyethylene terephthalate vial sealed with nitrile bung and aluminium overseal. Pack size: 12 x 100ml vials

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:
12/03/2019

Manufacturing sites for batch release:
Univet Limited

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
838793

Date of authorisation status change:
12/03/2019

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: 11/02/2022
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Labelling

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Package Leaflet

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