Multimin Solution for Injection for Cattle

Authorised

- Zinc oxide
- Manganese carbonate
- Sodium selenite
- Copper(II) carbonate

Product identification

Medicine name:
Multimin Solution for Injection for Cattle
Multimin (10 mg +15 mg + 60 mg + 5 mg)/ml Roztwór do wstrzykiwań

Active substance:
Zinc oxide

Manganese carbonate

Sodium selenite

Copper(II) carbonate

Target species:
Cattle

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

Zinc oxide
74.68 milligram(s) / 1.00 millilitre(s)

Manganese carbonate
20.92 milligram(s) / 1.00 millilitre(s)

Sodium selenite
10.95 milligram(s) / 1.00 millilitre(s)

Copper(II) carbonate
26.09 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Cattle**
  - Meat and offal. 28 day
  - Milk. 0 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA12CX99

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Poland

Package description:

Primary packaging: Clear Polyethylene Terephthalate (PET) bottle closed with grey bromobutyl rubber stopper sealed with aluminium cap.Package sizes: Cardboard box containing one vial of 500 ml

Primary packaging: Clear Polyethylene Terephthalate (PET) bottle closed with grey bromobutyl rubber stopper sealed with aluminium cap.Package sizes: Cardboard box

containing one vial of 100 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Warburton Technology Limited

Marketing authorisation date:
26/05/2021

Manufacturing sites for batch release:
Laboratoires Biove

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
3099

Date of authorisation status change:
26/05/2021

Reference member state:
Ireland

Procedure number:
IE/V/0322/001

Concerned member states:
Austria Belgium Bulgaria Croatia Denmark Estonia Finland France Greece Hungary Italy Latvia Liechtenstein Lithuania Luxembourg Netherlands Norway Poland Portugal Romania Slovakia Slovenia Spain Sweden United Kingdom (Northern Ireland)

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

Documents

Labelling

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

Package Leaflet

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

Summary of Product Characteristics

English (PDF)
Published on: 28/01/2022
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Combined File of all Documents
