

Multimin Solution for Injection for Cattle

Authorised

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

Product identification

Medicine name:

Multimin Solution for Injection for Cattle

MULTIMIN injekčný roztok pre hovädzí dobytok

Active substance:

Zinc oxide

Copper(II) carbonate

Sodium selenite

Manganese 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)

Copper(II) carbonate

26.09 milligram(s) / 1.00 millilitre(s)

Sodium selenite

10.95 milligram(s) / 1.00 millilitre(s)

Manganese carbonate

20.92 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

Authorisation status:

Valid

Authorised in:

Slovakia

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:

7/04/2021

Manufacturing sites for batch release:

Laboratoires Biove

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/001/MR/21-S

Date of authorisation status change:

7/04/2021

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 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

Combined File of all Documents

English (PDF)

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Summary of Product Characteristics