

# 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

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**Active substance:**

Zinc oxide

Copper(II) carbonate

Sodium selenite

Manganese carbonate

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**Target species:**

Cattle

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**Route of administration:**

Subcutaneous use

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## 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)

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### **Pharmaceutical form:**

Solution for injection

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### **Withdrawal period by route of administration:**

#### **Subcutaneous use:**

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#### **Cattle**

- Meat and offal. 28 day

- Milk. 0 hour

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### **Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QA12CX99

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### **Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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### **Authorisation status:**

Valid

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### **Authorised in:**

Ireland

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### **Available in:**

Ireland

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**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

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Warburton Technology Limited

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**Marketing authorisation date:**

10/07/2015

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**Manufacturing sites for batch release:**

Laboratoires Biove

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**Responsible authority:**

Health Products Regulatory Authority

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**Authorisation number:**

VPA10431/001/001

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**Date of authorisation status change:**

10/07/2015

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0322/001

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**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)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

Summary of Product Characteristics