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# Multimin Solution for Injection for Cattle

Ima dovoljenje za promet

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

## Informacije o zdravilu

**Ime zdravila:**
Multimin Solution for Injection for Cattle

**Učinkovina:**
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English

**Ciljne živalske vrste:**
govedo

**Pot uporabe:**
Subkutana uporaba

## Podatki o zdravilu

**Učinkovina / Jakost:**

Na voljo samo v English
74.68 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
26.09 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
10.95 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
20.92 milligram(s) / 1.00 millilitre(s)

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**Farmacevtska oblika:**

Raztopina za injiciranje

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**Karenca glede na pot uporabe:**

**Subkutana uporaba:**

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

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**Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):**

QA12CX99

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**Pravni status za dobavo/izdajo zdravila:**

Zdravilo, ki se izdaja na veterinarski recept

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**Status dovoljenja za promet z zdravilom:**

Valid

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**Ima dovoljenje za promet v:**

Na voljo samo v Spanish Czech German Estonian English French Italian Lithuanian Dutch Portuguese Romanian Slovak Swedish Icelandic Norwegian

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**Na voljo v:**

Romania

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**Opis ovojnine zdravila:**

Na voljo samo v English

Na voljo samo v English

## Dodatne informacije

**Vrsta dovoljenja:**
Na voljo samo v English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

**Pravna podlaga za izdajo dovoljenja za promet z zdravilom:**
Na voljo samo v English Italian

**Imetnik dovoljenja za promet z zdravilom:**
Warburton Technology Limited

**Datum dovoljenja za promet z zdravilom:**
10/03/2021

**Proizvodna mesta, odgovorna za sproščanje serij:**
Laboratoires Biove

**Pristojni organ:**
Institute For Control Of Biological Products And Veterinary Medicines

**Številka dovoljenja za promet z zdravilom:**
260044

**Datum spremembe statusa dovoljenja za promet:**
9/03/2026

**Referenčna država članica:**
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

**Številka postopka:**
IE/V/0322/001

**Zadevne države članice:**

Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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Za poročila o domnevnih neželenih učinkih zdravil za uporabo v veterinaski medicini obiščite stran: www.adrreports.eu/vet

## Dokumenti

Combined File of all Documents

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.

Povzetek glavnih značilnosti zdravila

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.