# Bovilis Cryptium (--) - Emulsion for injection

Authorised

- Cryptosporidium parvum, glycoprotein gp40

## Product identification

**Medicine name:**
Bovilis Cryptium (--) - Emulsion for injection

**Active substance:**
Cryptosporidium parvum, glycoprotein gp40

**Target species:**
Cattle

**Route of administration:**
Subcutaneous use

## Product details

**Active substance and strength:**
Cryptosporidium parvum, glycoprotein gp40
1.00 unit(s) / 1.00 Vial

**Pharmaceutical form:**
Emulsion for injection

**Withdrawal period by route of administration:**

**Subcutaneous use:**

- **Cattle**
  - Not applicable. 0 day Zero days

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

QI02AO02

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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:**

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

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**Available in:**

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Ireland , Italy , Latvia , Lithuania , Luxembourg , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Spain , Sweden , United Kingdom (Northern Ireland)

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**Package description:**

Packaging:Vial (PET), Package_size:1 vial (5 doses), Content:10 ml

Packaging:Vial (glass), Package_size:1 vial (5 doses), Content:10 ml

Packaging:Vial (glass), Package_size:10 vial (10 x 1 doses), Content:2 ml

Packaging:Vial (PET), Package_size:1 vial (50 doses), Content:100 ml

Packaging:Vial (PET), Package_size:1 vial (20 doses), Content:40 ml

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## Additional information

**Entitlement type:**

Marketing Authorisation

**Legal basis of product authorisation:**
Full application - new active substance (Article 8 of Regulation (EU) 2019/6)

**Marketing authorisation holder:**
Intervet International B.V.

**Marketing authorisation date:**
23/11/2023

**Manufacturing sites for batch release:**
Intervet International B.V.

**Responsible authority:**
European Commission

**Authorisation number:**
This information is not available for this product.

**Date of authorisation status change:**
23/11/2023

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

## Documents

Combined File of all Documents

English (PDF)
Published on: 19/12/2024
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