

# Engemycin Spray, 25mg/mL, cutaneous spray, suspension for cattle, sheep and pigs

Pooblašeno

- Oxytetracycline

## Product identification

### Ime zdravila:

Engemycin Spray, 25mg/mL, cutaneous spray, suspension for cattle, sheep and pigs  
Engemycin Spray, 25 mg/ml, δερματικό εκνέφωμα, εναιώρημα, για βοοειδή, πρόβατα  
και χοίρους

### Učinkovina:

Na voljo samo v [English](#)

### Ciljne živalske vrste:

govedo

Na voljo samo v [Bulgarian](#) [Spanish](#) [Czech](#) [Danish](#) [German](#) [Estonian](#) [Greek](#) [English](#)  
[French](#) [Italian](#) [Latvian](#) [Lithuanian](#) [Hungarian](#) [Dutch](#) [Romanian](#) [Finnish](#) [Swedish](#)  
[Icelandic](#) [Norwegian](#)

Na voljo samo v [Bulgarian](#) [Spanish](#) [Czech](#) [Danish](#) [German](#) [Estonian](#) [Greek](#) [English](#)  
[French](#) [Italian](#) [Latvian](#) [Lithuanian](#) [Hungarian](#) [Dutch](#) [Romanian](#) [Finnish](#) [Swedish](#)  
[Icelandic](#) [Norwegian](#)

### Pot uporabe:

Dermalna uporaba

## Product details

### **Učinkovina / Jakost:**

Na voljo samo v English

25.00 milligram(s) / 1.00 millilitre(s)

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### **Farmacevtska oblika:**

Dermalno pršilo, suspenzija

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

#### **Dermalna uporaba:**

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#### **govedo**

- Meat and offal. 0 day

- Milk. 0 day

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#### **Sheep**

- Meat and offal. 0 day

- Milk. 0 day

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#### **Pig**

- Meat and offal. 0 day

Stained part of the pig skin must be removed prior to the rest of the animal being used for human consumption

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### **Anatomsko-terapevtsko-kemična veterinarska oznaka (ATCvet):**

QD06AA03

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

Zdravilo, ki se izdaja na veterinarski recept

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### **Status dovoljenja:**

Valid

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**Authorised in:**

Na voljo samo v [Spanish](#) [Czech](#) [German](#) [Estonian](#) [Greek](#) [English](#) [French](#) [Italian](#) [Dutch](#) [Portuguese](#) [Slovak](#) [Swedish](#) [Icelandic](#) [Norwegian](#)

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**Opis ovojnine:**

Na voljo samo v [English](#)

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## Additional information

**Entitlement type:**

Na voljo samo v [English](#) [French](#) [Croatian](#) [Italian](#) [Latvian](#) [Finnish](#) [Swedish](#) [Icelandic](#) [Norwegian](#)

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**Pravna podlaga za izdajo dovoljenja za promet z zdravilom:**

Na voljo samo v [English](#) [Italian](#)

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**Imetnik dovoljenja za promet z zdravilom:**

Intervet International B.V.

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**Marketing authorisation date:**

11/04/2010

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

Intervet Productions S.r.l.

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**Pristojni organ:**

Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

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**Številka dovoljenja :**

CY00199V

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**Datum spremembe statusa dovoljenja:**

11/04/2010

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**Referenčna država članica:**

Na voljo samo v [Spanish](#) [Czech](#) [German](#) [Estonian](#) [English](#) [French](#) [Italian](#) [Dutch](#) [Portuguese](#) [Slovak](#) [Swedish](#) [Icelandic](#) [Norwegian](#)

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**Številka postopka:**

**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

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

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

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

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Na voljo samo v Estonian English French Portuguese Swedish Icelandic Norwegian

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Za poročila o domnevnih neželenih učinkih zdravil za uporabo v veterinarski medicini obiščite stran: [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

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

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**Source URL:** <https://medicines.health.europa.eu/veterinary/700000090839>