

# Bovilis IBR marker Live

Pooblašeno

- Infectious Bovine rhinotracheitis virus, strain GK/D gE gene-deleted, Inactivated

## Product identification

**Ime zdravila:**

Bovilis IBR marker Live  
Bovilis IBR Marker live

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**Učinkovina:**

Na voljo samo v [English](#)

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**Ciljne živalske vrste:**

govedo

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**Pot uporabe:**

Nazalna uporaba  
intramuskularna uporaba

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## Product details

**Učinkovina / Jakost:**

Na voljo samo v [English](#)  
7.30 unit(s) / 1.00 Dose

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**Farmaceutska oblika:**

Liofilizat za suspenzijo za injiciranje

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**Withdrawal period by route of administration:**

**Nazalna uporaba:****• govedo**

- Milk. 0 day
- Meat and offal. 0 day

**intramuskularna uporaba:****• govedo**

- Milk. 0 day
- Meat and offal. 0 day

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

QI02AD01

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

Zdravilo, ki se izdaja na veterinarski recept

**Status dovoljenja:**

Valid

**Authorised in:**

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

**Opis ovojnine:**

Na voljo samo v [English](#)

Na voljo samo v [English](#)

Na voljo samo v [English](#)

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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](#) [French](#) [Italian](#) [Latvian](#) [Norwegian](#)

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**Imetnik dovoljenja za promet z zdravilom:**

Intervet Deutschland GmbH

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**Marketing authorisation date:**

2/02/2012

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

INTERVET INTERNATIONAL B.V.

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**Pristojni organ:**

Paul Ehrlich Institute

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**Številka dovoljenja :**

PEI.V.11616.01.1

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**Datum spremembe statusa dovoljenja:**

2/02/2012

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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:**

NL/V/0105/001

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**Zadevne države članice:**

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

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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

Povzetek glavnih značilnosti zdravila

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/600000039861>