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# ECHINACEA COMPOSITUM, injekcinis tirpalas

Ima dovoljenje za promet

- ECHINACEA D3
- ACONITUM NAPELLUS D4
- SULFUR D8
- LACHESIS MUTUS D 10
- BRYONIA CRETICA D6
- HYDRARGYRUM BICHLORATUM D6
- PHOSPHORUS D8
- ARNICA MONTANA D6

## Informacije o zdravilu

**Ime zdravila:**
ECHINACEA COMPOSITUM, injekcinis tirpalas

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

**Ciljne živalske vrste:**

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

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**Pot uporabe:**
Subkutana uporaba

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## Podatki o zdravilu

**Učinkovina / Jakost:**
Na voljo samo v English
0.50 millilitre(s) / 1.00 Ampula

Na voljo samo v English
0.10 millilitre(s) / 1.00 Ampula

Na voljo samo v English
0.10 millilitre(s) / 1.00 Ampula

Na voljo samo v English
0.10 millilitre(s) / 1.00 Ampula

Na voljo samo v English
0.10 millilitre(s) / 1.00 Ampula

Na voljo samo v English

0.10 millilitre(s) / 1.00 Ampula

Na voljo samo v English
0.10 millilitre(s) / 1.00 Ampula

Na voljo samo v English
0.10 millilitre(s) / 1.00 Ampula

**Farmacevtska oblika:**
Raztopina za injiciranje

**Pravni status za dobavo/izdajo zdravila:**
Zdravilo, ki se izdaja brez veterinarskega recepta

**Status dovoljenja za promet z zdravilom:**
Valid

**Ima dovoljenje za promet v:**
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

**Opis ovojnine zdravila:**
Na voljo samo v Lithuanian
Na voljo samo v Lithuanian

## Dodatne informacije

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

**Imetnik dovoljenja za promet z zdravilom:**
Biologische Heilmittel Heel GmbH

**Datum dovoljenja za promet z zdravilom:**
23/03/2003

**Proizvodna mesta, odgovorna za sproščanje serij:**
Biologische Heilmittel Heel GmbH

**Pristojni organ:**
State Food And Veterinary Service

**Številka dovoljenja za promet z zdravilom:**
LT/2/03/1531/001-002

**Datum spremembe statusa dovoljenja za promet:**
28/06/2026

Za poročila o domnevnih neželenih učinkih zdravil za uporabo v veterinaski medicini obiščite stran: www.adrreports.eu/vet

## Dokumenti

Navodilo za uporabo

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