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**Source URL:** https://medicines.health.europa.eu/veterinary/en/600000001079

# Advocate 80 mg + 8 mg - Spot-on solution

Ima dovoljenje za promet

- Moxidectin
- Imidacloprid

## Informacije o zdravilu

**Ime zdravila:**
Advocate 80 mg + 8 mg - Spot-on solution

**Učinkovina:**
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

**Pot uporabe:**
Nanos na kožo

## Podatki o zdravilu

**Učinkovina / Jakost:**
Na voljo samo v English
Presentation_strength:8 mg Reference:Hse Index:0

Na voljo samo v English
Presentation_strength:80 mg Reference:Hse Index:1

**Farmacevtska oblika:**
Kožni nanos, raztopina

**Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):**
QP54AB52

**Pravni status za dobavo/izdajo zdravila:**
Zdravilo, ki se izdaja na veterinarski recept

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

**Ima dovoljenje za promet v:**
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Na voljo samo v Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

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

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**Opis ovojnine zdravila:**

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
Na voljo samo v English

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## 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

**Imetnik dovoljenja za promet z zdravilom:**
Elanco Animal Health GmbH

**Datum dovoljenja za promet z zdravilom:**
2/04/2003

**Proizvodna mesta, odgovorna za sproščanje serij:**
KVP Pharma+Veterinaer Produkte GmbH

**Pristojni organ:**
European Commission

**Številka dovoljenja za promet z zdravilom:**
Ta podatek za to zdravilo ni na voljo.

**Datum spremembe statusa dovoljenja za promet:**
2/04/2003

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

## Dokumenti

Combined File of all Documents

slovenščina (PDF)

Objavljeno na dan: 14/03/2026

Prenesi

ema-puar-advocate-v-076-var-ii-0041-g-en.pdf

ema-puar-advocate-v-076-var-ii-0039-g-en.pdf

ema-puar-advocate-v-076-par-en.pdf

ema-puar-advocate-v-076-var-ii-0026-g-en.pdf

ema-puar-advocate-v-076-var-ii-0022-en.pdf

ema-puar-advocate-v-000046-par-en.pdf