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# Hedylon 25 mg tablete za pse

Ima dovoljenje za promet

- Prednisolone

## Informacije o zdravilu

**Ime zdravila:**
Hedylon 25 mg tablete za pse

**Učinkovina:**
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:**
Peroralna uporaba

## Podatki o zdravilu

**Učinkovina / Jakost:**
Na voljo samo v English
25.00 milligram(s) / 1.00 Tableta

**Farmacevtska oblika:**

Tableta

**Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):**
QH02AB06

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

**Na voljo v:**
Slovenia

**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

## 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 Italian Latvian Norwegian

**Imetnik dovoljenja za promet z zdravilom:**
Industrial Veterinaria S.A.

**Datum dovoljenja za promet z zdravilom:**
27/02/2019

**Proizvodna mesta, odgovorna za sproščanje serij:**
Industrial Veterinaria S.A.
aniMedica GmbH
aniMedica Herstellungs GmbH

**Pristojni organ:**
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

**Številka dovoljenja za promet z zdravilom:**
DC/V/0650/002

**Datum spremembe statusa dovoljenja za promet:**
27/02/2019

**Referenčna država članica:**
Na voljo samo v Spanish Czech German Estonian English French Croatian Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

**Številka postopka:**
ES/V/0292/002

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

## Dokumenti

Povzetek glavnih značilnosti zdravila

slovenščina (PDF)
Objavljeno na dan: 17/04/2026
Prenesi

Navodilo za uporabo

slovenščina (PDF)
Objavljeno na dan: 17/04/2026
Prenesi

Označevanje

slovenščina (PDF)
Objavljeno na dan: 17/04/2026
Prenesi

Combined File of all Documents

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

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