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# ZIPYRAN 50 MG / 50 MG / 150 MG TABLETS FOR DOGS

Ima dovoljenje za promet

- Praziquantel
- Febantel
- Pyrantel embonate

## Informacije o zdravilu

**Ime zdravila:**
ZIPYRAN 50 MG / 50 MG / 150 MG TABLETS FOR DOGS

**Učinkovina:**
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

**Pot uporabe:**
Peroralna uporaba

## Podatki o zdravilu

**Učinkovina / Jakost:**

Na voljo samo v English
50.00 milligram(s) / 1.00 Tableta

Na voljo samo v English
150.00 milligram(s) / 1.00 Tableta

Na voljo samo v English
144.00 milligram(s) / 1.00 Tableta

**Farmacevtska oblika:**

Tableta

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

QP52AA51

**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

**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

## 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:**
Laboratorios Calier S.A.

**Datum dovoljenja za promet z zdravilom:**
25/06/2011

**Proizvodna mesta, odgovorna za sproščanje serij:**
Laboratorios Calier S.A.
Beaphar B.V.
Beaphar B.V.

**Pristojni organ:**
Federal Office Of Consumer Protection And Food Safety

**Številka dovoljenja za promet z zdravilom:**
401431.00.00

**Datum spremembe statusa dovoljenja za promet:**
1/12/2016

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

**Številka postopka:**
FR/V/0224/001

**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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Na voljo samo v Estonian English French Lithuanian Portuguese 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

Combined File of all Documents

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