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# Wellplus Flavoured Tablets

Ima dovoljenje za promet

- Praziquantel
- Pyrantel embonate
- Febantel

## Informacije o zdravilu

**Ime zdravila:**
Wellplus Flavoured Tablets

**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
144.00 milligram(s) / 1.00 Tableta

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

**Farmacevtska oblika:**

Tableta

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

QP52AA30

**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

## 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:**
Divasa Farmavic S.A.

**Datum dovoljenja za promet z zdravilom:**
26/02/2015

**Proizvodna mesta, odgovorna za sproščanje serij:**
Divasa Farmavic S.A.

**Pristojni organ:**
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

**Številka dovoljenja za promet z zdravilom:**
2409

**Datum spremembe statusa dovoljenja za promet:**
26/02/2015

**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/0163/001

**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

Povzetek glavnih značilnosti zdravila

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

Navodilo za uporabo

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

Označevanje

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

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