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# FRONTLINE TRI-ACT 33.8 MG / 252.4 MG SPOT-ON SOLUTION FOR DOGS 2- 5 KG

Ima dovoljenje za promet

- Fipronil
- Permethrin

## Informacije o zdravilu

**Ime zdravila:**
FRONTLINE TRI-ACT 33.8 MG / 252.4 MG SPOT-ON SOLUTION FOR DOGS 2- 5 KG

**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:**
Dermalna uporaba

## Podatki o zdravilu

**Učinkovina / Jakost:**

Na voljo samo v English
33.80 milligram(s) / 1.00 Pipeta

Na voljo samo v English
252.40 milligram(s) / 1.00 Pipeta

**Farmacevtska oblika:**
Kožni nanos, raztopina

**Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):**
QP53AX65

**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

**Na voljo v:**
Austria

**Opis ovojnine zdravila:**
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:**

Boehringer Ingelheim Animal Health France

**Datum dovoljenja za promet z zdravilom:**
16/09/2014

**Proizvodna mesta, odgovorna za sproščanje serij:**
Boehringer Ingelheim Animal Health France

**Pristojni organ:**
Austrian Agency For Health And Food Safety

**Številka dovoljenja za promet z zdravilom:**
835757

**Datum spremembe statusa dovoljenja za promet:**
16/09/2014

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

**Številka postopka:**
FR/V/0266/001

**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

Navodilo za uporabo

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

Povzetek glavnih značilnosti zdravila

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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