Advantix Spot-on solution for dogs over 10 kg up to 25 kg

Pooblaščeno

- Imidacloprid
- Permethrin (Cis:Trans 40:60)

Product identification

Ime zdravila:
Advantix Spot-on solution for dogs over 10 kg up to 25 kg
ADVANTIX CHIEN MOYEN

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 Swedish Icelandic Norwegian

Pot uporabe:
Nanos na kožo

Product details

Učinkovina / Jakost:
Na voljo samo v English
250.00 milligram(s) / 1.00 Merilna kapalka

Na voljo samo v English

1250.00 milligram(s) / 1.00 Merilna kapalka

Farmacevtska oblika:

Kožni nanos, raztopina

Withdrawal period by route of administration:

Nanos na kožo:

- **Dog**

Anatomsko-terapevtsko-kemična veterinarska oznaka (ATCvet):

QP53AC54

Pravni status za dobavo / izdajo zdravila:

Zdravilo, ki se izdaja brez veterinarskega recepta

Status dovoljenja:

Valid

Authorised in:

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

Available in:

France

Opis ovojnine:

Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
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Na voljo samo v English
Na voljo samo v English

Additional information

Entitlement type:
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

Imetnik dovoljenja za promet z zdravilom:
Elanco GmbH

Marketing authorisation date:
27/01/2004

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

Pristojni organ:
French Agency For Food, Environmental And Occupational Health & Safety

Številka dovoljenja :
FR/V/0721081 9/2004

Datum spremembe statusa dovoljenja:
27/01/2009

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

Številka postopka:
IT/V/0115/001

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 Latvian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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Na voljo samo v Estonian English French Swedish Icelandic Norwegian

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To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents

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

Povzetek glavnih značilnosti zdravila

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

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