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STRECTIS 68 MG/34 MG SPOT-ON SOLUTION FOR CATS 0.5-5 KG

Awtorizzat

- Fipronil
- Methoprene

Product identification

Isem tal-mediċina:
STRECTIS 68 MG/34 MG SPOT-ON SOLUTION FOR CATS 0.5-5 KG
Strectis 68 mg/34 mg, Lösung zum Auftropfen für Katzen von 0,5-5 kg

Sustanza attiva:
Disponibbli biss fi English
Disponibbli biss fi English

Speċi li fuqhom ser jintuża l-prodott:
Disponibbli biss fi Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian

Metodu ta’ amministrazzjoni:
Użu għall-ġilda

Product details

Sustanza attiva / Qawwa:
Disponibbli biss fi English

68.00 milligram(s) / 1.00 Pipette

Disponibbli biss fi English
34.00 milligram(s) / 1.00 Pipette

Forma farmaċewtika:

Soluzzjoni għall-użu lokalizzat

Withdrawal period by route of administration:

Użu għall-ġilda:

- **Cat**

Kodiċi veterinarju anatomiku terapewtiku kimiku (ATCvet):

QP53AX65

Status legali tal-provvista:

Disponibbli biss fi Czech Estonian English French Italian Latvian Lithuanian Portuguese Romanian Slovenian Finnish Swedish Icelandic Norwegian

Status tal-awtorizzazzjoni:

Valid

Authorised in:

Disponibbli biss fi Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Deskrizzjoni tal-pakkett:

Disponibbli biss fi French
Disponibbli biss fi French
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Disponibbli biss fi French
Disponibbli biss fi French
Disponibbli biss fi French

Additional information

Entitlement type:

Disponibbli biss fi English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

Bażi ġuridika tal-awtorizzazzjoni tal-prodott:
Disponibbli biss fi English Italian Latvian Lithuanian Norwegian

Id-detentur tal-awtorizzazzjoni għat-tqegħid fis-suq:
Ceva Tiergesundheit GmbH

Marketing authorisation date:
16/12/2015

Siti ta’ manifattura b’rilaxx tal-lott:
Ceva Sante Animale
Klocke Verpackungs-Service GmbH

Awtorità responsabbli:
Federal Office Of Consumer Protection And Food Safety

Numru tal-awtorizzazzjoni:
402220.00.00

Data tal-bidla fl-istatus tal-awtorizzazzjoni:
2/05/2020

Stat Membru ta’ referenza:
Disponibbli biss fi Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Numru tal-proċedura:
FR/V/0346/002

Stati Membri Kkonċernati:
Disponibbli biss fi Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Disponibbli biss fi Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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Disponibbli biss fi Spanish Czech German Estonian English French Croatian Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Disponibbli biss fi Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

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

Documents

Sommarju tal-karatteristiċi tal-prodott

Dan id-dokument ma jeżistix f'dan il-lingwa (Malti). Tista 'ssibha f'lingwa oħra hawn taħt.

Fuljett ta’ tagħrif

Dan id-dokument ma jeżistix f'dan il-lingwa (Malti). Tista 'ssibha f'lingwa oħra hawn taħt.