PROTECTIX 100 MG/500 MG SPOT-ON SOLUTION FOR DOGS OVER 4 KG UP TO 10 KG

Authorised

- Imidacloprid
- Permethrin

Product identification

Medicine name:
PROTECTIX 100 MG/500 MG SPOT-ON SOLUTION FOR DOGS OVER 4 KG UP TO 10 KG

Active substance:
Imidacloprid

Permethrin

Target species:
Dog

Route of administration:
Cutaneous use

Product details

Active substance and strength:
Imidacloprid
100.00 milligram(s) / 1.00 Pipette

Permethrin
500.00 milligram(s) / 1.00 Pipette

Pharmaceutical form:
Spot-on solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP53AC54

Legal status of supply:
Veterinary medicinal product not subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Slovenia

Package description:
Pack containing 1 PET/PE/aluminium/surlyn sachet containing one white polypropylene unit dose pipette
Pack containing 2 PET/PE/aluminium/surlyn sachets each containing one white polypropylene unit dose pipette
Pack containing 3 PET/PE/aluminium/surlyn sachets each containing one white polypropylene unit dose pipette
Pack containing 4 PET/PE/aluminium/surlyn sachets each containing one white polypropylene unit dose pipette
Pack containing 6 PET/PE/aluminium/surlyn sachets each containing one white polypropylene unit dose pipette
Pack containing 12 PET/PE/aluminium/surlyn sachets each containing one white polypropylene unit dose pipette
Pack containing 24 PET/PE/aluminium/surlyn sachets each containing one white polypropylene unit dose pipette

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Beaphar B.V.

Marketing authorisation date:
24/01/2024

Manufacturing sites for batch release:
Beaphar B.V.

Responsible authority:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:
DC/V/0801/002

Date of authorisation status change:
24/01/2024

Reference member state:
France

Procedure number:
FR/V/0433/002

Concerned member states:
Croatia Cyprus Czechia Germany Greece Hungary Italy Latvia Lithuania Malta Netherlands Poland Portugal Romania Slovakia Slovenia Spain

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

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents

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