PROTECTIX 40 MG/200 MG SPOT-ON SOLUTION FOR DOGS UP TO 4 KG

Authorised

- Imidacloprid
- Permethrin

Product identification

Medicine name:
PROTECTIX 40 MG/200 MG SPOT-ON SOLUTION FOR DOGS UP TO 4 KG

Active substance:
Imidacloprid

Permethrin

Target species:
Dog

Route of administration:
Cutaneous use

Product details

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

Permethrin
200.00 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:
Romania

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:
4/11/2024

Manufacturing sites for batch release:
Beaphar B.V.

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:
240141

Date of authorisation status change:
4/11/2024

Reference member state:
France

Procedure number:
FR/V/0433/001

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

Combined File of all Documents

English (PDF)
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