PROTECTIX 250 MG/1250 MG SPOT-ON SOLUTION FOR DOGS OVER 10 KG UP TO 25 KG

Authorised

- Imidacloprid
- Permethrin

Product identification

Medicine name:
PROTECTIX 250 MG/1250 MG SPOT-ON SOLUTION FOR DOGS OVER 10 KG UP TO 25 KG
ProtecTix 250 mg/1250 mg spot-on roztok pre psy od 10 kg do 25 kg.

Active substance:
Imidacloprid

Permethrin

Target species:
Dog

Route of administration:
Cutaneous use

Product details

Active substance and strength:
Imidacloprid

250.00 milligram(s) / 1.00 Pipette

Permethrin
1250.00 milligram(s) / 1.00 Pipette

Pharmaceutical form:
Spot-on solution

Withdrawal period by route of administration:

Cutaneous use:

- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP53AC54

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

Authorisation status:
Valid

Authorised in:
Slovakia

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:
This information is not available for this product.

Manufacturing sites for batch release:
Beaphar B.V.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/023/DC/24-S

Date of authorisation status change:
29/05/2024

Reference member state:
France

Procedure number:
FR/V/0433/003

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

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