

PROTECTIX 400 MG/2000 MG SPOT-ON SOLUTION FOR DOGS OVER 25 KG UP TO 40 KG

Authorised

- Imidacloprid
- Permethrin

Product identification

Medicine name:

PROTECTIX 400 MG/2000 MG SPOT-ON SOLUTION FOR DOGS OVER 25 KG UP TO 40 KG

ProtecTix 400 mg/2000 mg spot-on oplossing voor honden van meer dan 25 kg en tot 40 kg

Active substance:

Imidacloprid

Permethrin

Target species:

Dog

Route of administration:

Cutaneous use

Product details

Active substance and strength:

Imidacloprid

400.00 milligram(s) / 1.00 Pipette

Permethrin

2000.00 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:

Netherlands

Available in:

Netherlands

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:

22/01/2024

Manufacturing sites for batch release:

Beaphar B.V.

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 128715

Date of authorisation status change:

14/06/2024

Reference member state:

France

Procedure number:

FR/V/0433/004

Concerned member states:

Croatia Cyprus Czechia Germany Greece Hungary Italy Latvia Lithuania

Malta Netherlands Poland Portugal Romania Slovakia Slovenia Spain

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Documents

Combined File of all Documents

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