PULSIX 600 mg/3000 mg spot-on solution for dogs over 40 kg up to 60 kg

Authorised

- Imidacloprid
- Permethrin (40:60)

Product identification

Medicine name:
PULSIX 600 mg/3000 mg spot-on solution for dogs over 40 kg up to 60 kg
PULSIX 600 mg + 3000 mg Lösung zum Auftropfen für Hunde über 40 kg bis 60 kg

Active substance:
Imidacloprid
Permethrin (40:60)

Target species:
Dog

Route of administration:
Spot-on use

Product details

Active substance and strength:
Imidacloprid

600.00 milligram(s) / 1.00 Pipette

Permethrin (40:60)
3000.00 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:
Germany

Package description:
Cardboard box containing 1 pipette of 6.0 ml with pouch
HISTORICAL Cardboard box containing 1 pipette of 6.0 ml without pouch
Cardboard box containing 2 pipettes of 6.0 ml with pouch
HISTORICAL Cardboard box containing 2 pipettes of 6.0 ml without pouch
Cardboard box containing 3 pipettes of 6.0 ml with pouch
HISTORICAL Cardboard box containing 3 pipettes of 6.0 ml without pouch
Cardboard box containing 4 pipettes of 6.0 ml with pouch
HISTORICAL Cardboard box containing 4 pipettes of 6.0 ml without pouch
Cardboard box containing 6 pipettes of 6.0 ml with pouch
HISTORICAL Cardboard box containing 6 pipettes of 6.0 ml without pouch
Cardboard box containing 24 pipettes of 6.0 ml with pouch
HISTORICAL Cardboard box containing 24 pipettes of 6.0 ml without pouch
HISTORICAL Cardboard box containing 12 pipettes of 6.0 ml without pouch
Cardboard box containing 12 pipettes of 6.0 ml with pouch

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:
Ab7 Sante

Marketing authorisation date:
8/11/2024

Manufacturing sites for batch release:
Ab7 Sante

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
V7006982.00.00

Date of authorisation status change:
8/11/2024

Reference member state:
Ireland

Procedure number:
IE/V/0667/005

Concerned member states:
France Germany Italy Netherlands Poland Portugal Spain

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

Documents

Combined File of all Documents

English (PDF)
Published on: 26/10/2025
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Summary of Product Characteristics