PULSIX 400 mg/2000 mg spot-on solution for dogs over 25 kg up to 40 kg

Authorised

- Imidacloprid
- Permethrin (40:60)

Product identification

Medicine name:
PULSIX 400 mg/2000 mg spot-on solution for dogs over 25 kg up to 40 kg
PULSIX 400 mg + 2000 mg Lösung zum Auftropfen für Hunde über 25 kg bis 40 kg

Active substance:
Imidacloprid
Permethrin (40:60)

Target species:
Dog

Route of administration:
Spot-on use

Product details

Active substance and strength:
Imidacloprid

400.00 milligram(s) / 1.00 Pipette

Permethrin (40:60)
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:
Germany

Package description:
HISTORICAL Cardboard box containing 1 pipette of 4.0 ml without pouch
Cardboard box containing 2 pipettes of 4.0 ml with pouch
HISTORICAL Cardboard box containing 2 pipettes of 4.0 ml without pouch
Cardboard box containing 3 pipettes of 4.0 ml with pouch
HISTORICAL Cardboard box containing 3 pipettes of 4.0 ml without pouch
Cardboard box containing 4 pipettes of 4.0 ml with pouch
HISTORICAL Cardboard box containing 4 pipettes of 4.0 ml without pouch
Cardboard box containing 6 pipettes of 4.0 ml with pouch
HISTORICAL Cardboard box containing 6 pipettes of 4.0 ml without pouch
Cardboard box containing 24 pipettes of 4.0 ml with pouch
HISTORICAL Cardboard box containing 24 pipettes of 4.0 ml without pouch
Cardboard box containing 1 pipette of 4.0 ml with pouch
HISTORICAL Cardboard box containing 12 pipettes of 4.0 ml without pouch
Cardboard box containing 12 pipettes of 4.0 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:
V7006981.00.00

Date of authorisation status change:
8/11/2024

Reference member state:
Ireland

Procedure number:
IE/V/0667/004

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)
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Summary of Product Characteristics