PULSIX 40 mg/200 mg spot-on solution for dogs over 1.5 kg up to 4 kg

Authorised

- Imidacloprid
- Permethrin (40:60)

Product identification

Medicine name:
PULSIX 40 mg/200 mg spot-on solution for dogs over 1.5 kg up to 4 kg
Pulsix (40 mg + 200 mg)/pipetkę 0,4 ml Roztwór do nakrapiania

Active substance:
Imidacloprid
Permethrin (40:60)

Target species:
Dog

Route of administration:
Spot-on use

Product details

Active substance and strength:
Imidacloprid

40.00 milligram(s) / 1.00 Pipette

Permethrin (40:60)
200.00 milligram(s) / 1.00 Pipette

Pharmaceutical form:
Spot-on solution

Withdrawal period by route of administration:
Spot-on 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:
Poland

Package description:
Cardboard box containing 1 pipette of 0.4 ml with pouch
HISTORICAL Cardboard box containing 1 pipette of 0.4 ml without pouch
Cardboard box containing 2 pipettes of 0.4 ml with pouch
HISTORICAL Cardboard box containing 2 pipettes of 0.4 ml without pouch
HISTORICAL Cardboard box containing 3 pipettes of 0.4 ml without pouch
Cardboard box containing 4 pipettes of 0.4 ml with pouch
HISTORICAL Cardboard box containing 4 pipettes of 0.4 ml without pouch
Cardboard box containing 6 pipettes of 0.4 ml with pouch
HISTORICAL Cardboard box containing 6 pipettes of 0.4 ml without pouch
Cardboard box containing 24 pipettes of 0.4 ml with pouch
HISTORICAL Cardboard box containing 24 pipettes of 0.4 ml without pouch
Cardboard box containing 3 pipettes of 0.4 ml with pouch
HISTORICAL Cardboard box containing 12 pipettes of 0.4 ml without pouch
Cardboard box containing 12 pipettes of 0.4 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:
21/01/2025

Manufacturing sites for batch release:
Ab7 Sante

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
3378

Date of authorisation status change:
21/01/2025

Reference member state:
Ireland

Procedure number:
IE/V/0667/001

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

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.