PULSIX 100 mg/500 mg spot-on solution for dogs over 4 kg up to 10 kg

Authorised

- Imidacloprid
- Permethrin (40:60)

Product identification

Medicine name:
PULSIX 100 mg/500 mg spot-on solution for dogs over 4 kg up to 10 kg
PULSIX 100 MG/500 MG SOLUTION POUR SPOT-ON POUR CHIENS DE 4 KG A 10 KG

Active substance:
Imidacloprid
Permethrin (40:60)

Target species:
Dog

Route of administration:
Spot-on use

Product details

Active substance and strength:
Imidacloprid
100.00 milligram(s) / 1.00 Pipette

Permethrin (40:60)
500.00 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:
France

Package description:
Cardboard box containing 1 pipette of 1.0 ml with pouch
Cardboard box containing 1 pipette of 1.0 ml without pouch
Cardboard box containing 2 pipettes of 1.0 ml without pouch
Cardboard box containing 2 pipettes of 1.0 ml with pouch
Cardboard box containing 3 pipettes of 1.0 ml with pouch
Cardboard box containing 3 pipettes of 1.0 ml without pouch
Cardboard box containing 4 pipettes of 1.0 ml with pouch
Cardboard box containing 4 pipettes of 1.0 ml without pouch
Cardboard box containing 6 pipettes of 1.0 ml with pouch
Cardboard box containing 6 pipettes of 1.0 ml without pouch
Cardboard box containing 24 pipettes of 1.0 ml with pouch
Cardboard box containing 24 pipettes of 1.0 ml without pouch
Cardboard box containing 12 pipettes of 1.0 ml without pouch
Cardboard box containing 12 pipettes of 1.0 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:
27/09/2024

Manufacturing sites for batch release:
Ab7 Sante

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/0297003 1/2024

Date of authorisation status change:
27/09/2024

Reference member state:
Ireland

Procedure number:
IE/V/0667/002

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

Package Leaflet and Labelling

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

Summary of Product Characteristics

English (PDF)
Published on: 20/10/2024
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