REMUVET 250 MG/1250 MG SPOT-ON SOLUTION FOR DOGS OVER 10 KG UP TO 25 KG

Authorised

- Imidacloprid
- Permethrin

Product identification

Medicine name:
REMUVET 250 MG/1250 MG SPOT-ON SOLUTION FOR DOGS OVER 10 KG UP TO 25 KG
REMUVET 250 mg/1250 mg SOLUCION SPOT-ON PARA PERROS DE MAS DE 10 KG HASTA 25 Kg

Active substance:
Imidacloprid

Permethrin

Target species:
Dog

Route of administration:
Cutaneous use

Product details

Active substance and strength:

Imidacloprid
250.00 milligram(s) / 1.00 Pipette

Permethrin
1250.00 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:
Spain

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:
25/04/2024

Manufacturing sites for batch release:
Beaphar B.V.

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
4286 ESP

Date of authorisation status change:
26/04/2024

Reference member state:
France

Procedure number:
FR/V/0440/003

Concerned member states:
Italy 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.