

Imoxiclate 250 mg/62.5 mg spot-on solution for large dogs

Authorised

- Moxidectin
- Imidacloprid

Product identification

Medicine name:

Imoxiclate 250 mg/62.5 mg spot-on solution for large dogs

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Active substance:

Moxidectin

Imidacloprid

Target species:

Dog

Route of administration:

Spot-on use

Product details

Active substance and strength:

Moxidectin

62.50 milligram(s) / 1.00 Pipette

Imidacloprid
250.00 milligram(s) / 1.00 Pipette

Pharmaceutical form:

Spot-on solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AB52

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Package description:

A white polypropylene (PP) unit dose pipette with a closure with a spike composed of high density polyethylene (HDPE) or polyoxymethylene (POM) or polypropylene (PP) packed into a laminated triplex bag composed of polyester (PETP), aluminium(Al) and low density polyethylene (LDPE).Cardboard box containing 1 pipettes.

A white polypropylene (PP) unit dose pipette with a closure with a spike composed of high density polyethylene (HDPE) or polyoxymethylene (POM) or polypropylene (PP) packed into a laminated triplex bag composed of polyester (PETP), aluminium(Al) and low density polyethylene (LDPE).Cardboard box containing 3 pipettes.

A white polypropylene (PP) unit dose pipette with a closure with a spike composed of high density polyethylene (HDPE) or polyoxymethylene (POM) or polypropylene (PP) packed into a laminated triplex bag composed of polyester (PETP), aluminium(Al) and low density polyethylene (LDPE).Cardboard box containing 4 pipettes.

A white polypropylene (PP) unit dose pipette with a closure with a spike composed of high density polyethylene (HDPE) or polyoxymethylene (POM) or polypropylene (PP) packed into a laminated triplex bag composed of polyester (PETP), aluminium(Al) and low density polyethylene (LDPE).Cardboard box containing 6 pipettes.

A white polypropylene (PP) unit dose pipette with a closure with a spike composed of high density polyethylene (HDPE) or polyoxymethylene (POM) or polypropylene (PP) packed into a laminated triplex bag composed of polyester (PETP), aluminium(Al) and low density polyethylene (LDPE).Cardboard box containing 24 pipettes.

A white polypropylene (PP) unit dose pipette with a closure with a spike composed of high density polyethylene (HDPE) or polyoxymethylene (POM) or polypropylene (PP) packed into a laminated triplex bag composed of polyester (PETP), aluminium(Al) and low density polyethylene (LDPE).Cardboard box containing 48 pipettes.

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:

KRKA tovarna zdravil d.d. Novo mesto

Marketing authorisation date:

13/03/2020

Manufacturing sites for batch release:

KRKA tovarna zdravil d.d. Novo mesto

TAD Pharma GmbH

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA10774/067/003

Date of authorisation status change:

13/03/2020

Reference member state:

Ireland

Procedure number:

IE/V/0565/005

Concerned member states:

Germany

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

Documents

Summary of Product Characteristics