Imoxicate 100 mg/25 mg spot-on solution for medium dogs

Authorised

- Imidacloprid
- Moxidectin

Product identification

Medicine name:
Imoxicate 100 mg/25 mg spot-on solution for medium dogs
Imoxicate 100 mg/25 mg spot-on solution for medium dogs

Active substance:
Imidacloprid

Moxidectin

Target species:
Dog

Route of administration:
Spot-on use

Product details

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

Moxidectin
25.00 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:
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) orpolyoxymethylene (POM) or polypropylene (PP) packed into a laminated triplex bag composed of polyester (PETP), aluminium(Al) and low density polyetylene (LDPE).Cardboard box containing 1 pipette.
A white polypropylene (PP) unit dose pipette with a closure with a spike composed of high density polyethylene (HDPE) orpolyoxymethylene (POM) or polypropylene (PP) packed into a laminated triplex bag composed of polyester (PETP), aluminium(Al) and low density polyetylene (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) orpolyoxymethylene (POM) or polypropylene (PP) packed into a laminated triplex bag composed of polyester (PETP), aluminium(Al) and low density polyetylene (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) orpolyoxymethylene (POM) or polypropylene (PP) packed into a laminated triplex bag composed of polyester (PETP), aluminium(Al) and low density polyetylene (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) orpolyoxymethylene (POM) or polypropylene (PP) packed into a laminated triplex bag composed of polyester (PETP), aluminium(Al) and low density polyetylene (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) orpolyoxymethylene (POM) or polypropylene (PP) packed into a laminated triplex bag composed of polyester (PETP), aluminium(Al) and low density polyetylene (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:
TAD Pharma GmbH
KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10774/067/002

Date of authorisation status change:
13/03/2020

Reference member state:
Ireland

Procedure number:
IE/V/0565/004

Concerned member states:
Germany

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

Documents

Summary of Product Characteristics