

Imoxicate 40 mg/10 mg spot-on solution for small dogs

Authorised

This information is not available for this product.

Product identification

Medicine name:

Imoxicate 40 mg/10 mg spot-on solution for small dogs

Imoxicate 40 mg/10 mg Lösung zum Auftropfen für kleine Hunde

Active substance:

This information is not available for this product.

Target species:

Dog

Route of administration:

Spot-on use

Product details

Active substance and strength:

This information is not available for this product.

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:

Germany

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 pipette.

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:

30/12/2019

Manufacturing sites for batch release:

Tad Pharma GmbH

Krka d.d. Novo Mesto

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:

402663.00.00

Date of authorisation status change:

30/12/2019

Reference member state:

Ireland

Procedure number:

IE/V/0565/003

Concerned member states:

Germany

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

Documents

Summary of Product Characteristics

Source URL: <https://medicines.health.europa.eu/veterinary/600000052240>