

Selehold 360 mg spot-on solution for dogs 40.1 60.0 kg

Authorised

- Selamectin

Product identification

Medicine name:

Selehold 360 mg spot-on solution for dogs 40.1  60.0 kg

Selehold 360 mg šķīdums pilināšanai uz ādas suņiem 40,1–60,0 kg

Active substance:

Selamectin

Target species:

Dog

Route of administration:

Spot-on use

Product details

Active substance and strength:

Selamectin

360.00 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:

QP54AA05

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Latvia

Package description:

Translucent polypropylene unit-dose pipette with polyethylene or polyoxymethylene or polypropylene closure with spike packaged into a laminated triplex bag composed of polyester, aluminium and polyethylene. 6 ml pipette containing 3.0 ml of solution. Cardboard box containing 1 pipette.

Translucent polypropylene unit-dose pipette with polyethylene or polyoxymethylene or polypropylene closure with spike packaged into a laminated triplex bag composed of polyester, aluminium and polyethylene. 6 ml pipette containing 3.0 ml of solution. Cardboard box containing 3 pipettes.

Translucent polypropylene unit-dose pipette with polyethylene or polyoxymethylene or polypropylene closure with spike packaged into a laminated triplex bag composed of polyester, aluminium and polyethylene. 6 ml pipette containing 3.0 ml of solution. Cardboard box containing 6 pipettes.

Translucent polypropylene unit-dose pipette with polyethylene or polyoxymethylene or polypropylene closure with spike packaged into a laminated triplex bag composed of polyester, aluminium and polyethylene. 6 ml pipette containing 3.0 ml of solution. Cardboard box containing 15 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:

4/12/2018

Manufacturing sites for batch release:

Krka d.d. Novo Mesto

Tad Pharma GmbH

Responsible authority:

Food And Veterinary Service

Authorisation number:

V/DCP/18/0072

Date of authorisation status change:

4/12/2018

Reference member state:

Ireland

Procedure number:

IE/V/0395/005

Concerned member states:

Belgium Bulgaria Croatia Czechia Estonia France Greece Hungary Italy
Latvia Lithuania Netherlands Poland Portugal Romania Slovakia Slovenia
Spain United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

Combined File of all Documents

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