Selehold 15 mg spot-on solution for cats and dogs = 2.5 kg

Authorised

- Selamectin

Product identification

Medicine name:
Selehold 15 mg spot-on solution for cats and dogs = 2.5 kg
Selehold 15 mg roztok na kvapkanie na kožu pre mačky a psy ≤ 2,5 kg

Active substance:
Selamectin

Target species:
Dog
Cat

Route of administration:
Spot-on use

Product details

Active substance and strength:
Selamectin
15.00 milligram(s) / 1.00 Pipette

Pharmaceutical form:
Spot-on solution

Withdrawal period by route of administration:

Spot-on use:

- **Dog**
- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AA05

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Slovakia

Package description:

Translucent polypropylene unit-dose pipette with polyethylene or polyoxymethylene or polypropylene closure with spikepackaged into a laminated triplex bag composed of polyester, aluminium and polyethylene. 1 ml pipette containing 0.25 ml of solution.Cardboard box containing 1 pipette.

Translucent polypropylene unit-dose pipette with polyethylene or polyoxymethylene or polypropylene closure with spikepackaged into a laminated triplex bag composed of polyester, aluminium and polyethylene. 1 ml pipette containing 0.25 ml of solution.Cardboard box containing 3 pipettes.

Translucent polypropylene unit-dose pipette with polyethylene or polyoxymethylene or polypropylene closure with spikepackaged into a laminated triplex bag composed of polyester, aluminium and polyethylene. 1 ml pipette containing 0.25 ml of solution.Cardboard box containing 6 pipettes.

Translucent polypropylene unit-dose pipette with polyethylene or polyoxymethylene or polypropylene closure with spikepackaged into a laminated triplex bag composed of polyester, aluminium and polyethylene. 1 ml pipette containing 0.25 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:
12/12/2018

Manufacturing sites for batch release:
Krka d.d. Novo Mesto
Tad Pharma GmbH

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/061/DC/18-S

Date of authorisation status change:
12/12/2018

Reference member state:
Ireland

Procedure number:
IE/V/0394/001

Concerned member states:
Belgium Bulgaria Croatia Czechia Estonia France Germany 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

Combined File of all Documents

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Summary of Product Characteristics
