Selehold 45 mg spot-on solution for cats 2.6–7.5 kg

Authorised

- Selamectin

Product identification

Medicine name:
Selehold 45 mg spot-on solution for cats 2.6–7.5 kg
Selehold 45 mg voor katten 2,6 - 7,5 kg 60 mg/ml Spot-on oplossing
Selehold 45 mg pour chats 2,6 - 7,5 kg 60 mg/ml Solution pour spot-on
Selehold 45 mg fur Katzen 2,6 - 7,5 kg 60 mg/ml Lösung zum Auftropfen

Active substance:
Selamectin

Target species:
Cat

Route of administration:
Spot-on use

Product details

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

Pharmaceutical form:
Spot-on solution

Withdrawal period by route of administration:

Spot-on use:

- **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:

Belgium

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. 3 ml pipette containing 0.75 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. 3 ml pipette containing 0.75 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. 3 ml pipette containing 0.75 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. 3 ml pipette containing 0.75 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:
11/01/2019

Manufacturing sites for batch release:
TAD Pharma GmbH
KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V537893

Date of authorisation status change:
11/01/2019

Reference member state:
Ireland

Procedure number:
IE/V/0394/002

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

Summary of Product Characteristics

English (PDF)
Published on: 3/05/2024
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Package Leaflet

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Labelling

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