

# 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

---

**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

---

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QP54AA05

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Slovakia

---

**Available in:**

Slovakia

---

**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. 3 ml pipette containing 0.75 ml of solution. Cardboard box containing 15 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. 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 spike packaged 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 spike packaged 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.

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

Generic application (Article 13(1) 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:**

TAD Pharma GmbH

KRKA tovarna zdravil d.d. Novo mesto

---

**Responsible authority:**

Institute For State Control Of Veterinary Biologicals And Medicaments

---

**Authorisation number:**

96/062/DC/18-S

---

**Date of authorisation status change:**

12/12/2018

---

**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](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

This document does not exist in this language (English). You can find it in another language below.

Summary of Product Characteristics