

# Selehold 30 mg spot-on solution for dogs 2.6–5.0 kg

Authorised

- Selamectin

## Product identification

**Medicine name:**

Selehold 30 mg spot-on solution for dogs 2.6–5.0 kg

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**Active substance:**

Selamectin

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**Target species:**

Dog

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**Route of administration:**

Spot-on use

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## Product details

**Active substance and strength:**

Selamectin

30.00 milligram(s) / 1.00 Pipette

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**Pharmaceutical form:**

Spot-on solution

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

QP54AA05

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Latvia

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**Available in:**

Latvia

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**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. 1 ml pipette containing 0.25 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. 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 spike packaged 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 spike packaged 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.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

KRKA tovarna zdravil d.d. Novo mesto

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**Marketing authorisation date:**

4/12/2018

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**Manufacturing sites for batch release:**

TAD Pharma GmbH

KRKA tovarna zdravil d.d. Novo mesto

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**Responsible authority:**

Food And Veterinary Service

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**Authorisation number:**

V/DCP/18/0068

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**Date of authorisation status change:**

4/12/2018

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0395/001

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**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)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

Combined File of all Documents

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