

# Selames 120 mg spot-on solution for dogs 10.1–20.0 kg

Authorised

- Selamectin

## Product identification

**Medicine name:**

Selames 120 mg spot-on solution for dogs 10.1–20.0 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

120.00 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:**

Belgium

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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. 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. 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. 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. 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:**

7/12/2018

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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:**

Federal Agency For Medicines And Health Products

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**Authorisation number:**

BE-V537084

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**Date of authorisation status change:**

7/12/2018

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0407/003

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**Concerned member states:**

Belgium France Germany Italy Netherlands Portugal 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

English (PDF)

Published on: 18/01/2026

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Package Leaflet

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Labelling

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