# Selames 240 mg spot-on solution for dogs 20.1–40.0 kg

Authorised

- Selamectin

## Product identification

**Medicine name:**
Selames 240 mg spot-on solution for dogs 20.1–40.0 kg
Selames voor honden 20,1-40,0 kg 240 mg Spot-on oplossing
Selames pour chiens 20,1-40,0 kg 240 mg Solution pour spot-on
Selames fur Hunde 20,1-40,0 kg 240 mg Lösung zum Auftropfen

**Active substance:**
Selamectin

**Target species:**
Dog

**Route of administration:**
Spot-on use

## Product details

**Active substance and strength:**
Selamectin
240.00 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:**
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.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.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.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.Cardboard box containing 15 pipettes.

## 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:**
7/12/2018

**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-V537093

**Date of authorisation status change:**
7/12/2018

**Reference member state:**
Ireland

**Procedure number:**
IE/V/0407/004

**Concerned member states:**
Belgium France Germany Italy Netherlands Portugal 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)

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Package Leaflet

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Labelling

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