

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

Selames 120 mg Lösung zum Auftropfen für Hunde 10,1 - 20,0 kg

Active substance:

Selamectin

Target species:

Dog

Route of administration:

Spot-on use

Product details

Active substance and strength:

Selamectin

120.00 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:

Germany

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 1 pipette.

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 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 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:

5/11/2018

Manufacturing sites for batch release:

TAD Pharma GmbH
KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:

402515.00.00

Date of authorisation status change:

5/11/2018

Reference member state:

Ireland

Procedure number:

IE/V/0407/003

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)

Published on: 18/01/2026

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Package Leaflet

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