

Selames 15 mg spot-on solution for cats and dogs ≤ 2.5 kg

Authorised

- Selamectin

Product identification

Medicine name:

Selames 15 mg spot-on solution for cats and dogs ≤ 2.5 kg

Selames 15 mg solution pour spot-on pour chats et chiens $\leq 2,5$ kg

Selames 15 mg spot-on oplossing voor katten en honden $\leq 2,5$ kg

Selames 15 mg Lösung zum Auftropfen für Katzen und Hunde $\leq 2,5$ kg

Active substance:

Selamectin

Target species:

Dog

Cat

Route of administration:

Spot-on use

Product details

Active substance and strength:

Selamectin

15.00 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 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:

6/03/2019

Manufacturing sites for batch release:

KRKA tovarna zdravil d.d. Novo mesto

TAD Pharma GmbH

Responsible authority:

Federal Agency For Medicines And Health Products

Authorisation number:

BE-V539733

Date of authorisation status change:

6/03/2019

Reference member state:

Ireland

Procedure number:

IE/V/0406/001

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: 3/05/2024

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Package Leaflet

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Labelling

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