

Selames 60 mg spot-on solution for dogs 5.1–10.0 kg

Authorised

- Selamectin

Product identification

Medicine name:

Selames 60 mg spot-on solution for dogs 5.1–10.0 kg

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Active substance:

Selamectin

Target species:

Dog

Route of administration:

Spot-on use

Product details

Active substance and strength:

Selamectin

60.00 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:

Ireland

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:

21/12/2018

Manufacturing sites for batch release:

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

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA10774/056/002

Date of authorisation status change:

21/12/2018

Reference member state:

Ireland

Procedure number:

IE/V/0407/002

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

Combined File of all Documents