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 spot-on oplossing voor katten en honden ≤ 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:
Netherlands

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:
11/02/2019

Manufacturing sites for batch release:
KRKA tovarna zdravil d.d. Novo mesto
TAD Pharma GmbH

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 122203

Date of authorisation status change:
26/01/2022

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

Combined File of all Documents

This document does not exist in this language (English). You can find it in another language below.

Summary of Product Characteristics