

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 240 mg solução para unção punctiforme para cães com 20,1–40,0 kg

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:

Portugal

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:

15/11/2018

Manufacturing sites for batch release:

TAD Pharma GmbH

KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:

Directorate General For Food And Veterinary

Authorisation number:

1217/04/18DFVPT

Date of authorisation status change:

17/08/2022

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

Combined File of all Documents

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