

File downloaded on 2026-04-23

Source URL: <https://medicines.health.europa.eu/veterinary/en/600000088620>

Perfikan 268 mg/2400 mg spot-on solution for large

Authorised

- Permethrin
- Fipronil

Product identification

Medicine name:

Perfikan 268 mg/2400 mg spot-on solution for large

Active substance:

Permethrin

Fipronil

Target species:

Dog

Route of administration:

Spot-on use

Product details

Active substance and strength:

Permethrin

2398.00 milligram(s) / 1.00 Pipette

Fipronil

268.00 milligram(s) / 1.00 Pipette

Pharmaceutical form:

Spot-on solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP53AC54

Legal status of supply:

Veterinary medicinal product not subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Portugal

Package description:

The boxes contain individual pipette(s) placed in coloured overblister(s) made from polypropylene/cyclic olefin copolymer/polypropylene and closed with lid made from polyethylene-terephthalate/aluminium/polypropylene. Boxes 6 pipettes.

The boxes contain individual pipette(s) placed in coloured overblister(s) made from polypropylene/cyclic olefin copolymer/polypropylene and closed with lid made from polyethylene-terephthalate/aluminium/polypropylene. Boxes 4 pipettes.

The boxes contain individual pipette(s) placed in coloured overblister(s) made from polypropylene/cyclic olefin copolymer/polypropylene and closed with lid made from polyethylene-terephthalate/aluminium/polypropylene. Boxes of 2 pipettes.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Informed consent application (Article 13c of Directive No 2001/82/EC)

Marketing authorisation holder:

Alfamed

Marketing authorisation date:

6/06/2016

Manufacturing sites for batch release:

Alfamed

Responsible authority:

Directorate General For Food And Veterinary

Authorisation number:

1021/04/16DFVPT

Date of authorisation status change:

7/11/2020

Reference member state:

Portugal

Procedure number:

PT/V/0134/004

Concerned member states:

Italy Spain

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.