EFFIPRO 268 MG SPOT-ON SOLUTION FOR LARGE DOGS

Authorised

- Fipronil

Product identification

Medicine name:
EFFIPRO 268 MG SPOT-ON SOLUTION FOR LARGE DOGS
EFFIPRO 268 mg spot-on oplossing voor grote honden

Active substance:
Fipronil

Target species:
Dog

Route of administration:
Cutaneous use

Product details

Active substance and strength:
Fipronil
268.00 milligram(s) / 1.00 Pipette

Pharmaceutical form:
Spot-on solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP53AX15

Legal status of supply:
Veterinary medicinal product not subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Netherlands

Package description:
Box containing 2 pipettes of 2.68 mL
Box containing 3 pipettes of 2.68 mL
Box containing 4 pipettes of 2.68 mL
Box containing 6 pipettes of 2.68 mL
Box containing 8 pipettes of 2.68 mL
Box containing 12 pipettes of 2.68 mL
Box containing 24 pipettes of 2.68 mL
Box containing 30 pipettes of 2.68 mL
Box containing 60 pipettes of 2.68 mL
Box containing 90 pipettes of 2.68 mL
Box containing 150 pipettes of 2.68 mL
Box containing 1 blister pack of 1 pipette of 2.68 mL
Box containing 2 blister packs of 1 pipette of 2.68 mL
Box containing 3 blister packs of 1 pipette of 2.68 mL
Box containing 4 blister packs of 1 pipette of 2.68 mL
Box containing 6 blister packs of 1 pipette of 2.68 mL
Box containing 8 blister packs of 1 pipette of 2.68 mL
Box containing 12 blister packs of 1 pipette of 2.68 mL
Box containing 24 blister packs of 1 pipette of 2.68 mL
Box containing 30 blister packs of 1 pipette of 2.68 mL
Box containing 60 blister packs of 1 pipette of 2.68 mL
Box containing 90 blister packs of 1 pipette of 2.68 mL
Box containing 150 blister packs of 1 pipette of 2.68 mL
Box containing 1 pipette of 2.68 mL

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Virbac

Marketing authorisation date:
26/11/2008

Manufacturing sites for batch release:
Virbac

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 102117

Date of authorisation status change:
24/01/2024

Reference member state:
France

Procedure number:
FR/V/0377/003

Concerned member states:
Austria Belgium Bulgaria Cyprus Czechia Denmark Estonia Finland Germany Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Netherlands Poland Portugal Romania Slovakia Slovenia Spain Sweden 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.