EFFIPRO 50 MG SPOT-ON SOLUTION FOR CATS

Authorised

- Fipronil

Product identification

Medicine name:
EFFIPRO 50mg/pipette ΔΙΑΛΥΜΑ ΓΙΑ ΕΠΙΧΥΣΗ ΣΕ ΣΗΜΕΙΟ
EFFIPRO 50 MG SPOT-ON SOLUTION FOR CATS

Active substance:
Fipronil

Target species:
Cat

Route of administration:
Cutaneous use

Product details

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

Pharmaceutical form:
Spot-on solution

Withdrawal period by route of administration:

Cutaneous use:

- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP53AX15

Legal status of supply:

This information is not available for this product.

Authorisation status:

Valid

Authorised in:

Greece

Package description:

Box containing 2 pipettes of 0.5 mL
Box containing 3 pipettes of 0.5 mL
Box containing 4 pipettes of 0.5 mL
Box containing 6 pipettes of 0.5 mL
Box containing 8 pipettes of 0.5 mL
Box containing 12 pipettes of 0.5 mL
Box containing 24 pipettes of 0.5 mL
Box containing 30 pipettes of 0.5 mL
Box containing 60 pipettes of 0.5 mL
Box containing 90 pipettes of 0.5 mL
Box containing 150 pipettes of 0.5 mL
Box containing 1 blister pack of 1 pipette of 0.5 mL
Box containing 2 blister packs of 1 pipette of 0.5 mL
Box containing 3 blister packs of 1 pipette of 0.5 mL
Box containing 4 blister packs of 1 pipette of 0.5 mL
Box containing 6 blister packs of 1 pipette of 0.5 mL
Box containing 8 blister packs of 1 pipette of 0.5 mL
Box containing 12 blister packs of 1 pipette of 0.5 mL
Box containing 24 blister packs of 1 pipette of 0.5 mL
Box containing 30 blister packs of 1 pipette of 0.5 mL
Box containing 60 blister packs of 1 pipette of 0.5 mL
Box containing 90 blister packs of 1 pipette of 0.5 mL

Box containing 150 blister packs of 1 pipette of 0.5 mL
Box containing 1 pipette of 0.5 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:
24/04/2009

Manufacturing sites for batch release:
Virbac

Responsible authority:
National Organization For Medicines

Authorisation number:
19358/08-03-2017/K-0178202

Date of authorisation status change:
8/03/2017

Reference member state:
France

Procedure number:
FR/V/0376/001

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

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