EFFIPRO 50 MG SPOT-ON SOLUTION FOR CATS

Authorised

- Fipronil

Product identification

Medicine name:
EFFIPRO 50 MG SPOT-ON SOLUTION FOR CATS
Effipro 50 mg spot-on, opløsning

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP53AX15

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

Authorisation status:
Valid

Authorised in:
Denmark

Package description:
Box containing 90 pipettes of 0.5 mL
Box containing 60 pipettes of 0.5 mL
Box containing 3 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 8 pipettes of 0.5 mL
Box containing 4 pipettes of 0.5 mL
Box containing 2 pipettes of 0.5 mL
Box containing 150 pipettes of 0.5 mL
Box containing 6 pipettes of 0.5 mL
Box containing 1 pipette of 0.5 mL
Box containing 24 blister packs of 1 pipette of 0.5 mL
Box containing 4 blister packs of 1 pipette of 0.5 mL
Box containing 12 blister packs of 1 pipette of 0.5 mL
Box containing 8 blister packs of 1 pipette of 0.5 mL
Box containing 3 blister packs of 1 pipette of 0.5 mL
Box containing 6 blister packs of 1 pipette of 0.5 mL
Box containing 2 blister packs of 1 pipette of 0.5 mL
Box containing 30 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 blister pack of 1 pipette of 0.5 mL
Box containing 60 blister packs of 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:
30/04/2009

Manufacturing sites for batch release:
Virbac

Responsible authority:
Danish Medicines Agency

Authorisation number:
42990

Date of authorisation status change:
13/03/2025

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)

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Documents

Summary of Product Characteristics

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