EFFIPRO 67 MG SPOT-ON SOLUTION FOR SMALL DOGS

Authorised

- Fipronil

Product identification

Medicine name:
EFFIPRO 67 MG SPOT-ON SOLUTION FOR SMALL DOGS
EFFIPRO 67 mg Lösung zum Auftropfen für kleine Hunde

Active substance:
Fipronil

Target species:
Dog

Route of administration:
Cutaneous use

Product details

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

Pharmaceutical form:
Spot-on solution

Withdrawal period by route of administration:

Cutaneous use:

- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP53AX15

Legal status of supply:

Veterinary medicinal product not subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Germany

Package description:

Box containing 1 pipette of 0.67 mL
Box containing 2 pipettes of 0.67 mL
Box containing 3 pipettes of 0.67 mL
Box containing 4 pipettes of 0.67 mL
Box containing 6 pipettes of 0.67 mL
Box containing 8 pipettes of 0.67 mL
Box containing 12 pipettes of 0.67 mL
Box containing 24 pipettes of 0.67 mL
Box containing 30 pipettes of 0.67 mL
Box containing 60 pipettes of 0.67 mL
Box containing 90 pipettes of 0.67 mL
Box containing 150 pipettes of 0.67 mL
Box containing 1 blister pack of 1 pipette of 0.67 mL
Box containing 2 blister packs of 1 pipette of 0.67 mL
Box containing 4 blister packs of 1 pipette of 0.67 mL
Box containing 6 blister packs of 1 pipette of 0.67 mL
Box containing 8 blister packs of 1 pipette of 0.67 mL
Box containing 12 blister packs of 1 pipette of 0.67 mL
Box containing 24 blister packs of 1 pipette of 0.67 mL
Box containing 30 blister packs of 1 pipette of 0.67 mL
Box containing 60 blister packs of 1 pipette of 0.67 mL
Box containing 90 blister packs of 1 pipette of 0.67 mL

Box containing 150 blister packs of 1 pipette of 0.67 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:
9/03/2009

Manufacturing sites for batch release:
Virbac

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
401107.00.00

Date of authorisation status change:
24/03/2014

Reference member state:
France

Procedure number:
FR/V/0377/001

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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