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EFFIPRO 402 MG SPOT-ON SOLUTION FOR VERY LARGE DOGS

- Fipronil

Authorised

Product identification

Medicine name:

EFFIPRO 402 MG SPOT-ON SOLUTION FOR VERY LARGE DOGS

Active substance:

- Fipronil

Target species:

- Dog

Route of administration:

- Cutaneous use

Product details

Active substance and strength:

- Fipronil
402.00
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:

- Latvia

Package description:

- Box containing 1 pipette of 4.02 mL
- Box containing 150 blister packs of 1 pipette of 4.02 mL
- Box containing 90 blister packs of 1 pipette of 4.02 mL
- Box containing 60 blister packs of 1 pipette of 4.02 mL
- Box containing 30 blister packs of 1 pipette of 4.02 mL
- Box containing 24 blister packs of 1 pipette of 4.02 mL
- Box containing 12 blister packs of 1 pipette of 4.02 mL
- Box containing 8 blister packs of 1 pipette of 4.02 mL
- Box containing 6 blister packs of 1 pipette of 4.02 mL
- Box containing 4 blister packs of 1 pipette of 4.02 mL
- Box containing 3 blister packs of 1 pipette of 4.02 mL
- Box containing 2 blister packs of 1 pipette of 4.02 mL
- Box containing 1 blister pack of 1 pipette of 4.02 mL
- Box containing 150 pipettes of 4.02 mL
- Box containing 90 pipettes of 4.02 mL
- Box containing 60 pipettes of 4.02 mL
- Box containing 30 pipettes of 4.02 mL
- Box containing 24 pipettes of 4.02 mL
- Box containing 12 pipettes of 4.02 mL
- Box containing 8 pipettes of 4.02 mL
- Box containing 6 pipettes of 4.02 mL
- Box containing 4 pipettes of 4.02 mL
- Box containing 3 pipettes of 4.02 mL
- Box containing 2 pipettes of 4.02 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:

- Food And Veterinary Service

Authorisation number:

- V/DCP/22/0033

Date of authorisation status change:

- 9/03/2009

Reference member state:

- France

Procedure number:

- FR/V/0377/004

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

Product information

Combined File of all Documents

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Latvian (PDF)

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