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 kožni nanos, raztopina za mačke

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:
Veterinary medicinal product subject to veterinary prescription

---

Authorisation status:
Valid

Authorised in:
Slovenia

Package description:
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French

Available only in French
Available only in French
Available only in French

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:
12/03/2009

Manufacturing sites for batch release:
Virbac

Responsible authority:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:
DC/V/0104/005

Date of authorisation status change:
12/03/2009

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

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.
