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Broadline Fipronil 74.7 mg, (S)-methoprene 90 mg, eprinomectin 3.6 mg, praziquantel 74.7 mg - Spot-on solution

Authorised

- Eprinomectin
- Fipronil
- Praziquantel
- (S)-Methoprene

Product identification

Medicine name:
Broadline Fipronil 74.7 mg, (S)-methoprene 90 mg, eprinomectin 3.6 mg, praziquantel 74.7 mg - Spot-on solution

Active substance:
Eprinomectin

Fipronil

Praziquantel

(S)-Methoprene

Target species:
Cat

Route of administration:
Spot-on use

Product details

Active substance and strength:

Eprinomectin

Presentation_strength:3.6 mg Index:0

Fipronil

Presentation_strength:74.7 mg Index:1

Praziquantel

Presentation_strength:74.7 mg Index:2

(S)-Methoprene

Presentation_strength:90 mg Index:3

Pharmaceutical form:

Spot-on solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AA54

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Package description:

Packaging:Applicator (COC), Package_size:15 applicators, Content:0.9 ml

Packaging:Applicator (COC), Package_size:6 applicators, Content:0.9 ml

Packaging:Applicator (COC), Package_size:4 applicators, Content:0.9 ml

Packaging:Applicator (COC), Package_size:3 applicators, Content:0.9 ml

Packaging:Applicator (COC), Package_size:1 applicator, Content:0.9 ml

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Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Fixed combination application (Article 13b of Directive No 2001/82/EC)

Marketing authorisation holder:
Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:
4/12/2013

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
4/12/2013

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents

English (PDF)

Published on: 14/03/2026

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