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Felpreva 36.22 mg + 7.53 mg + 30.12 mg - Spot-on solution

Authorised

- Tigolaner
- Emodepside
- Praziquantel

Product identification

Medicine name:
Felpreva 36.22 mg + 7.53 mg + 30.12 mg - Spot-on solution

Active substance:
Tigolaner

Emodepside

Praziquantel

Target species:
Cat

Route of administration:
Spot-on use

Product details

Active substance and strength:
Tigolaner

36.22 milligram(s) / 1.00 Applicator

Emodepside
7.53 milligram(s) / 1.00 Applicator

Praziquantel
30.12 milligram(s) / 1.00 Applicator

Pharmaceutical form:
Spot-on solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52AA51

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

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)

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

Package description:
Packaging:Spot-on applicator with cap (PP), Package_size:10 spot-on applicators, Content:0.37 ml
Packaging:Spot-on applicator with cap (PP), Package_size:1 spot-on applicator, Content:0.37 ml
Packaging:Spot-on applicator with cap (PP), Package_size:20 spot-on applicators, Content:0.37 ml
Packaging:Spot-on applicator with cap (PP), Package_size:2 spot-on applicators, Content:0.37 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Vetoquinol S.A.

Marketing authorisation date:
11/11/2021

Manufacturing sites for batch release:
Vetoquinol S.A.
Vetoquinol Biowet Sp. z o.o.

Responsible authority:
European Commission

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

Date of authorisation status change:
11/11/2021

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: 23/10/2026

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