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Vectra Felis 423 mg/42.3 mg spot-on solution

Authorised

- Dinotefuran
- Pyriproxyfen

Product identification

Medicine name:
Vectra Felis 423 mg/42.3 mg spot-on solution

Active substance:
Dinotefuran

Pyriproxyfen

Target species:
Cat

Route of administration:
Spot-on use

Product details

Active substance and strength:
Dinotefuran
423.00 milligram(s) / 1.00 Applicator

Pyriproxyfen

42.30 milligram(s) / 1.00 Applicator

Pharmaceutical form:

Spot-on solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP53AX73

Legal status of supply:

Veterinary medicinal product not 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 , Cyprus , Czechia , Estonia , France , Germany , Greece , Hungary , Ireland , Italy , Latvia , Luxembourg , Netherlands , Poland , Portugal , Romania , Slovakia , Spain , United Kingdom (Northern Ireland)

Package description:

Packaging:Spot- on applicator (PE/Alu/PE), Package_size:1 spot-on applicator, Content:0.9 ml

Packaging:Spot- on applicator (PE/Alu/PE), Package_size:3 spot-on applicators, Content:0.9 ml

Packaging:Spot- on applicator (PE/Alu/PE), Package_size:6 spot-on applicators, Content:0.9 ml

Packaging:Spot- on applicator (PE/Alu/PE), Package_size:12 spot-on applicators, Content:0.9 ml

Packaging:Spot- on applicator (PE/Alu/PE), Package_size:4 spot-on applicators, Content:0.9 ml

Packaging:Spot- on applicator (PE/Alu/PE), Package_size:24 spot-on applicators, Content:0.9 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:
Ceva Sante Animale

Marketing authorisation date:
6/06/2014

Manufacturing sites for batch release:
Ceva Sante Animale
Ab7 Sante

Responsible authority:
European Commission

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

Date of authorisation status change:
6/06/2014

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: 13/03/2026

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