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NexGard Combo 3.6 mg + 1.2 mg + 24.9 mg - Spot-on solution

Authorised

- Esafoxolaner
- Eprinomectin
- Praziquantel

Product identification

Medicine name:
NexGard Combo 3.6 mg + 1.2 mg + 24.9 mg - Spot-on solution

Active substance:
Esafoxolaner

Eprinomectin

Praziquantel

Target species:
Cat

Route of administration:
Spot-on use

Product details

Active substance and strength:
Esafoxolaner

Presentation_strength:3.6 mg Reference:Hse Index:0

Eprinomectin

Presentation_strength:1.2 mg Reference:Hse Index:1

Praziquantel

Presentation_strength:24.9 mg Reference:Ph.Eur. Index:2

Pharmaceutical form:

Spot-on solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AA54

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 , Czechia , Denmark , Estonia , Finland , France , Latvia , Lithuania , Luxembourg , Norway , Poland , Slovakia , Spain , Sweden

Package description:

Packaging:Spot-on applicator (COC), Package_size:15 applicators, Content:0.3 ml

Packaging:Spot-on applicator (COC), Package_size:4 applicators, Content:0.3 ml

Packaging:Spot-on applicator (COC), Package_size:3 applicators, Content:0.3 ml

Packaging:Spot-on applicator (COC), Package_size:1 applicator, Content:0.3 ml

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:
6/01/2021

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:
6/01/2021

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: 13/03/2026
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