Vectra 3D 87 mg dinotefuran, 7.7 mg pyriproxyfen, 635 mg permethrin - Spot-on solution

Authorised

- Permethrin
- Pyriproxyfen
- Dinotefuran

Product identification

Medicine name:
Vectra 3D 87 mg dinotefuran, 7.7 mg pyriproxyfen, 635 mg permethrin - Spot-on solution

Active substance:
Permethrin

Pyriproxyfen

Dinotefuran

Target species:
Dog

Route of administration:
Spot-on use

Product details

Active substance and strength:

Permethrin

Presentation_strength:635 mg Reference:Hse Index:0

Pyriproxyfen

Presentation_strength:7.7 mg Reference:Hse Index:1

Dinotefuran

Presentation_strength:87 mg Reference:Hse Index:2

Pharmaceutical form:

Spot-on solution

Withdrawal period by route of administration:

Spot-on use:

- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP53A

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)

Package description:

Packaging:Spot-on applicator (PE/Alu/PE), Package_size:24 applicators, Content:1.6 ml

Packaging:Spot-on applicator (PE/Alu/PE), Package_size:4 applicators, Content:1.6 ml

Packaging:Spot-on applicator (PE/Alu/PE), Package_size:12 applicators, Content:1.6 ml

Packaging:Spot-on applicator (PE/Alu/PE), Package_size:6 applicators, Content:1.6 ml

Packaging:Spot-on applicator (PE/Alu/PE), Package_size:3 applicators, Content:1.6 ml

Packaging:Spot-on applicator (PE/Alu/PE), Package_size:1 applicator, Content:1.6 ml

Packaging:Spot-on applicator (PE/Alu/PE), Package_size:48 applicators, Content:1.6 ml

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
CEVA Santé Animale

Marketing authorisation date:
4/12/2013

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:
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: 20/04/2023
Download

ema-puar-vectra-3d-v-002555-referral-a82-0023-en.pdf

ema-puar-vectra-3d-v-2555-par-en.pdf
