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Simparica Trio 12 mg + 0.24 mg + 50 mg - Chewable tablet

Authorised

- Sarolaner
- Moxidectin
- Pyrantel embonate

Product identification

Medicine name:
Simparica Trio 12 mg + 0.24 mg + 50 mg - Chewable tablet

Active substance:
Sarolaner

Moxidectin

Pyrantel embonate

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Sarolaner

Presentation_strength:12 mg Reference:Hse Index:0

Moxidectin

Presentation_strength:0.24 mg Reference:Ph. Eur. Index:1

Pyrantel embonate

Presentation_strength:50 mg Reference:Ph. Eur. Index:2

Pharmaceutical form:

Chewable tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AB52

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 , Germany , Greece , Hungary , Ireland , Italy , Latvia , Lithuania , Luxembourg , Netherlands , Poland , Romania , Slovakia , Slovenia , Spain , United Kingdom (Northern Ireland)

Package description:

Packaging:Blister (alu/alu), Package_size:1 tablet

Packaging:Blister (alu/alu), Package_size:3 tablets

Packaging:Blister (alu/alu), Package_size:6 tablets

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:
Zoetis Belgium

Marketing authorisation date:
17/09/2019

Manufacturing sites for batch release:
Corden Pharma GmbH
Zoetis Belgium

Responsible authority:
European Commission

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

Date of authorisation status change:
17/09/2019

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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