

# Simparica Trio 6 mg + 0.12 mg + 25 mg - Chewable tablet

Authorised

- Sarolaner
- Moxidectin
- Pyrantel embonate

## Product identification

**Medicine name:**

Simparica Trio 6 mg + 0.12 mg + 25 mg - Chewable tablet

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**Active substance:**

Sarolaner

Moxidectin

Pyrantel embonate

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**Target species:**

Dog

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Sarolaner

Presentation\_strength:6 mg Reference:Hse Index:0

Moxidectin

Presentation\_strength:0.12 mg Reference:Ph. Eur. Index:1

Pyrantel embonate

Presentation\_strength:25 mg Reference:Ph. Eur. Index:2

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**Pharmaceutical form:**

Chewable tablet

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QP54AB52

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**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)

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**Available in:**

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Greece , Hungary , Ireland , Italy , Latvia , Lithuania , Luxembourg , Netherlands , Poland , Romania , Slovakia , Spain , United Kingdom (Northern Ireland)

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**Package description:**

Packaging:Blister (alu/alu), Package\_size:3 tablets

Packaging:Blister (alu/alu), Package\_size:1 tablet

Packaging:Blister (alu/alu), Package\_size:6 tablets

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Zoetis Belgium

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**Marketing authorisation date:**

17/09/2019

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**Manufacturing sites for batch release:**

Corden Pharma GmbH

Zoetis Belgium

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**Responsible authority:**

European Commission

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

17/09/2019

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

English (PDF)

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