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Simparica 5 mg - Chewable tablet

Authorised

- Sarolaner

Product identification

Medicine name:
Simparica 5 mg - Chewable tablet

Active substance:
Sarolaner

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Sarolaner

Presentation_strength:5 mg Reference:In house Index:0

Pharmaceutical form:
Chewable tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP53BE03

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 , Cyprus , Czechia , Denmark , Finland , Germany , Greece , Hungary , Ireland , Italy , Luxembourg , Netherlands , Poland , Romania , Slovakia , Spain , Sweden , United Kingdom (Northern Ireland)

Package description:
Packaging:Blister (alu/alu), Package_size:6 tablets
Packaging:Blister (alu/alu), Package_size:3 tablets
Packaging:Blister (alu/alu), Package_size:1 tablet

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:
6/11/2015

Manufacturing sites for batch release:
Zoetis Belgium

Responsible authority:

European Commission

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

Date of authorisation status change:
6/11/2015

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: 14/03/2026
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