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Credelio 112 mg - Chewable tablet (dogs)

Authorised

- Lotilaner
- Lotilaner

Product identification

Medicine name:
Credelio 112 mg - Chewable tablet (dogs)

Active substance:
Lotilaner
Lotilaner

Target species:
Dog
Dog

Route of administration:
Oral use
Oral use

Product details

Active substance and strength:
Lotilaner
Presentation_strength:112.5 mg Reference:Hse Index:0

Lotilaner

Presentation_strength:112.5 mg Reference:Hse Index:0

Pharmaceutical form:

Chewable tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP53BE04

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 , Finland , France , Germany , Greece , Hungary , Italy , Latvia , Lithuania , Luxembourg , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden

Package description:

Packaging:Blister (Alu/Alu), Package_size:18 tablets

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

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

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

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

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

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

Packaging:Blister (Alu/Alu), Package_size:18 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:
Elanco GmbH

Marketing authorisation date:
25/04/2017

Manufacturing sites for batch release:
Elanco France S.A.S.

Responsible authority:
European Commission

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

Date of authorisation status change:
25/04/2017

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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