

Credelio Plus 225 mg + 8.44 mg - Chewable tablet

Authorised

- Lotilaner
- Milbemycin oxime

Product identification

Medicine name:

Credelio Plus 225 mg + 8.44 mg - Chewable tablet

Active substance:

Lotilaner

Milbemycin oxime

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Lotilaner

Presentation_strength:225 mg Reference:In house Index:0

Milbemycin oxime

Presentation_strength:8.44 mg Reference:Ph. Eur. Index:1

Pharmaceutical form:

Chewable tablet

Withdrawal period by route of administration:**Oral use:**

- Dog
-

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AB51

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 , France , Germany , Greece , Hungary , Italy , Netherlands , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , United Kingdom (Northern Ireland)

Package description:

Packaging:Blister (alu), Package_size:18 tablets

Packaging:Blister (alu), Package_size:6 tablets

Packaging:Blister (alu), Package_size:1 tablet

Packaging:Blister (alu), Package_size:3 tablets

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Fixed combination application (Article 13b of Directive No 2001/82/EC)

Marketing authorisation holder:

Elanco GmbH

Marketing authorisation date:

14/04/2021

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:

14/04/2021

To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet

Documents

Combined File of all Documents

English (PDF)

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