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Frontpro 11.3 mg - Chewable tablet

Authorised

- Afoxolaner

Product identification

Medicine name:

Frontpro 11.3 mg - Chewable tablet

Active substance:

Afoxolaner

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Afoxolaner

Presentation_strength:11.3 mg Reference:Hse Index:0

Pharmaceutical form:

Chewable tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP53BE01

Legal status of supply:

Veterinary medicinal product not 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:

Belgium , Czechia , France , Hungary , Poland , Slovakia , Spain

Package description:

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

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

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

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Informed consent application (Article 13c of Directive No 2001/82/EC)

Marketing authorisation holder:

Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:

20/05/2019

Manufacturing sites for batch release:

Boehringer Ingelheim Animal Health France

Responsible authority:

European Commission

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

20/05/2019

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: 7/01/2026

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