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Frontpro 136 mg - Chewable tablet

Authorised

- Afoxolaner

Product identification

Medicine name:
Frontpro 136 mg - Chewable tablet

Active substance:
Afoxolaner

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Afoxolaner

Presentation_strength:136 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:
Austria , Belgium , Czechia , France , Hungary , Poland , Slovakia , Spain

Package description:
Packaging:Blister (Aclar/PVC/Alu), Package_size:3 tablets
Packaging:Blister (Aclar/PVC/Alu), Package_size:6 tablets
Packaging:Blister (Aclar/PVC/Alu), Package_size:1 tablet

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: 14/03/2026
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