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# Apoquel 5.4 mg - Chewable tablet

Authorised

- Oclacitinib maleate

## Product identification

**Medicine name:**
Apoquel 5.4 mg - Chewable tablet

**Active substance:**
Oclacitinib maleate

**Target species:**
Dog

**Route of administration:**
Oral use

## Product details

**Active substance and strength:**
Oclacitinib maleate
7.26 milligram(s) / 1.00 Tablet

**Pharmaceutical form:**
Chewable tablet

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QD11AH90

**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 , Germany , Greece , Hungary , Ireland , Italy , Latvia , Lithuania , Luxembourg , Netherlands , Poland , Portugal , Romania , Slovakia , Spain , Sweden ,
United Kingdom (Northern Ireland)

**Package description:**
Packaging:Blister (Alu/PVC/Aclar), Package_size:100 tablets
Packaging:Blister (Alu/PVC/Aclar), Package_size:20 tablets
Packaging:Blister (Alu/PVC/Aclar), Package_size:50 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:**
Zoetis Belgium

**Marketing authorisation date:**
12/09/2013

**Manufacturing sites for batch release:**
Pfizer Italia S.r.l.
Zoetis Belgium

**Responsible authority:**
European Commission

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

**Date of authorisation status change:**
13/12/2021

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: 28/04/2026
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