

# Apoquel 16 mg - Chewable tablet

Authorised

- Oclacitinib maleate

## Product identification

**Medicine name:**

Apoquel 16 mg - Chewable tablet

---

**Active substance:**

Oclacitinib maleate

---

**Target species:**

Dog

---

**Route of administration:**

Oral use

---

## Product details

**Active substance and strength:**

Oclacitinib maleate

21.50 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 SA

---

**Responsible authority:**

European Commission

---

**Authorisation number:**

This information is not available for this product.

---

**Date of authorisation status change:**

5/04/2023

---

To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

English (PDF)

Published on: 28/04/2026

[Download](#)

ema-puar-apoquel-v-2688-var-x-0019-en.pdf

ema-puar-apoquel-v-2688-par-en.pdf