

# Apoquel 16 mg - Film-coated tablet

Authorised

- Oclacitinib maleate

## Product identification

**Medicine name:**

Apoquel 16 mg - Film-coated tablet

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**Active substance:**

Oclacitinib maleate

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**Target species:**

Dog

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Oclacitinib maleate

21.50 milligram(s) / 1.00 Tablet

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**Pharmaceutical form:**

Film-coated tablet

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

QD11AH90

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**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)

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**Available in:**

Austria , Belgium , Bulgaria , Croatia , Czechia , Denmark , Estonia , Finland , Germany , Hungary , Iceland , Ireland , Italy , Latvia , Lithuania , Luxembourg , Netherlands , Poland , Romania , Slovakia , Spain , Sweden , United Kingdom (Northern Ireland)

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**Package description:**

Packaging:Blister (Alu/PVC/Aclar), Package\_size:20 tablets  
Packaging:Blister (Alu/PVC/Aclar), Package\_size:100 tablets  
Packaging:Blister (Alu/PVC/Aclar), Package\_size:50 tablets  
Packaging:Bottle (HDPE), Package\_size:20 tablets  
Packaging:Bottle (HDPE), Package\_size:100 tablets  
Packaging:Bottle (HDPE), Package\_size:50 tablets  
Packaging:Blister (PVC/PVDC/Alu), Package\_size:20 tablets  
Packaging:Blister (PVC/PVDC/Alu), Package\_size:50 tablets  
Packaging:Blister (PVC/PVDC/Alu), Package\_size:100 tablets

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Zoetis Belgium

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**Marketing authorisation date:**

12/09/2013

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**Manufacturing sites for batch release:**

Pfizer Italia S.r.l.

Zoetis Belgium

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**Responsible authority:**

European Commission

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

13/12/2021

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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)

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