

Apoquel 5.4 mg - Film-coated tablet

Authorised

- Oclacitinib maleate

Product identification

Medicine name:

Apoquel 5.4 mg - Film-coated 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:

Film-coated 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 , Czechia , Denmark , Estonia , Finland , Hungary , Ireland , Italy , Latvia , Lithuania , Luxembourg , Netherlands , Poland , Romania , Slovakia , Spain , Sweden , United Kingdom (Northern Ireland)

Package description:

Packaging:Blister (PVC/PVDC/Alu), Package_size:100 tablets

Packaging:Blister (PVC/PVDC/Alu), Package_size:50 tablets

Packaging:Blister (PVC/PVDC/Alu), Package_size:20 tablets

Packaging:Bottle (HDPE), Package_size:50 tablets

Packaging:Bottle (HDPE), Package_size:100 tablets

Packaging:Bottle (HDPE), Package_size:20 tablets

Packaging:Blister (Alu/PVC/Aclar), Package_size:50 tablets

Packaging:Blister (Alu/PVC/Aclar), Package_size:100 tablets

Packaging:Blister (Alu/PVC/Aclar), Package_size:20 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:

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)

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