Carprofen Orion 100 mg - Chewable tablet

Authorised

- Carprofen

Product identification

Medicine name:
Carprofen Orion 100 mg - Chewable tablet

Active substance:
Carprofen

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Carprofen
100.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Chewable tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QM01AE91

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

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) , 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)

Package description:

Packaging:Bottle (HDPE), Package_size:180 tablets

Packaging:Bottle (HDPE), Package_size:60 tablets

Packaging:Bottle (HDPE), Package_size:20 tablets

Packaging:Bottle (HDPE), Package_size:10 tablets

Additional information

Entitlement type:

Marketing Authorisation

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 18 of Regulation (EU) 2019/6)

Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:

Orion Corporation

Orion Corporation

Marketing authorisation date:
19/12/2024
19/12/2024

Manufacturing sites for batch release:
Orion Corporation
Orion Corporation
Orion Corporation

Responsible authority:
European Commission
European Commission

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

Date of authorisation status change:
19/12/2024
19/12/2024

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: 29/04/2025
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