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Clomicalm 20 mg - Tablet

Authorised

- Clomipramine hydrochloride
- Clomipramine hydrochloride

Product identification

Medicine name:
Clomicalm 20 mg - Tablet

Active substance:
Clomipramine hydrochloride
Clomipramine hydrochloride

Target species:
Dog
Dog

Route of administration:
Oral use
Oral use

Product details

Active substance and strength:
Clomipramine hydrochloride
20.00 milligram(s) / 1.00 Tablet
Clomipramine hydrochloride

20.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN06AA04

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 , Denmark , Estonia , Finland , France , Germany , Italy , Latvia , Lithuania , Luxembourg , Netherlands , Norway , Portugal , Sweden ,
United Kingdom (Northern Ireland)

Package description:
Packaging:Bottle (HDPE), Package_size:30 tablets
Packaging:Bottle (HDPE), Package_size:30 tablets

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Council Directive 81/851/EEC

Marketing authorisation holder:
Virbac

Marketing authorisation date:

1/04/1998

Manufacturing sites for batch release:
Virbac
Unither Developpement Bordeaux

Responsible authority:
European Commission

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

Date of authorisation status change:
1/04/1998

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: 14/03/2026
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