

Clomicalm 80 mg - Tablet

Authorised

- Clomipramine hydrochloride

Product identification

Medicine name:

Clomicalm 80 mg - Tablet

Active substance:

Clomipramine hydrochloride

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Clomipramine hydrochloride
80.00 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 , Ireland , Italy , Latvia , Lithuania , Luxembourg , Netherlands , Norway , Portugal , Sweden , United Kingdom (Northern Ireland)

Package description:

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 S.A.

Marketing authorisation date:

1/04/1998

Manufacturing sites for batch release:

Unither Developpement Bordeaux
Virbac S.A.

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)

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