

Reconcile 8 mg - Chewable tablet

Authorised

- Fluoxetine

Product identification

Medicine name:

Reconcile 8 mg - Chewable tablet

Active substance:

Fluoxetine

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Fluoxetine

8.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Chewable tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN06AB03

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 , Finland , France , Germany , Ireland , Italy , Luxembourg , Netherlands , Norway , Portugal , Spain , 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:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Forte Healthcare Limited

Marketing authorisation date:

8/07/2008

Manufacturing sites for batch release:

Forte Healthcare Limited

Responsible authority:

European Commission

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

8/07/2008

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: 19/09/2025

[Download](#)

ema-puar-reconcile-v-133-par-en.pdf