File downloaded on 2026-11-29
Source URL: https://medicines.health.europa.eu/veterinary/en/600000003465

Reconcile 32 mg - Chewable tablet

Authorised

- Fluoxetine

Product identification

Medicine name:
Reconcile 32 mg - Chewable tablet

Active substance:
Fluoxetine

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Fluoxetine
32.00 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: 1/09/2026
Download

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