

Tsefalen 50 mg/ml Powder for Oral Suspension for Dogs up to 20 kg and Cats

Authorised

- Cefalexin

Product identification

Medicine name:

Tsefalen 50 mg/ml Powder for Oral Suspension for Dogs up to 20 kg and Cats

Active substance:

Cefalexin

Target species:

Cat

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Cefalexin

50.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Powder for oral suspension

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01DB01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Package description:

Carton box with 1 bottle containing 66.6 g of powder providing 100 ml of suspension after reconstitution and 1 syringe of 5 ml

Carton box with 1 bottle containing 40.0 g of powder providing 60 ml of suspension after reconstitution and 1 syringe of 5 ml

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

Nextmune Italy S.r.l.

Marketing authorisation date:

4/12/2020

Manufacturing sites for batch release:

Acs Dobfar S.p.A.

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA23440/001/003

Date of authorisation status change:

4/12/2020

Reference member state:

Italy

Procedure number:

IT/V/0142/001

Concerned member states:

Austria Belgium Bulgaria Cyprus Czechia Denmark Estonia Finland France
Germany Greece Hungary Ireland Latvia Liechtenstein Lithuania
Luxembourg Malta Netherlands Norway Poland Portugal Romania Slovakia
Slovenia Spain Sweden United Kingdom (Northern Ireland)

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: 2/04/2026

[Download](#)