

Sporimune 50 mg/ml oral solution for cats and dogs

Authorised

- Ciclosporin

Product identification

Medicine name:

Sporimune 50 mg/ml oral solution for cats and dogs

Active substance:

Ciclosporin

Target species:

Dog

Cat

Route of administration:

Oral use

Product details

Active substance and strength:

Ciclosporin

50.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Oral solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QL04AD01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Available in:

France

Package description:

Brown glass bottles (type III) of 25 ml, closed with a child resistant closure (PP screw cap with a Teflon inlay). One bottle and a dispenser set (consisting of a child resistant HDPE screw cap and a 1 ml PP dosing syringe for cats and a 5 ml PP dosing syringe for dogs) packed in a cardboard box.

Brown glass bottles (type III) of 50 ml, closed with a child resistant closure (PP screw cap with a Teflon inlay). One bottle and a dispenser set (consisting of a child resistant HDPE screw cap and a 1 ml PP dosing syringe for cats and a 5 ml PP dosing syringe for dogs) packed in a cardboard box.

Brown glass bottles (type III) of 100 ml, closed with a child resistant closure (PP screw cap with a Teflon inlay). One bottle and a dispenser set (consisting of a child resistant HDPE screw cap and a 1 ml PP dosing syringe for cats and a 5 ml PP dosing syringe for dogs) packed in a cardboard box.

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:

Le Vet. Beheer B.V.

Marketing authorisation date:

27/05/2014

Manufacturing sites for batch release:

Produlab Pharma B.V.

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/2810441 6/2014

Date of authorisation status change:

9/10/2018

Reference member state:

Ireland

Procedure number:

IE/V/0302/001

Concerned member states:

Austria Belgium Croatia Cyprus Czechia Denmark Estonia Finland France
Germany Greece Hungary Iceland Italy Latvia Lithuania Luxembourg
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

Summary of Product Characteristics

English (PDF)

Published on: 16/01/2026

Download

Combined File of all Documents

Package Leaflet and Labelling

This document does not exist in this language (English). You can find it in another language below.