File downloaded on 2026-11-04
Source URL: https://medicines.health.europa.eu/veterinary/en/600000047222

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:
Sweden

Available in:
Sweden

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:
5/06/2014

Manufacturing sites for batch release:
Produlab Pharma B.V.

Responsible authority:
Swedish Medical Products Agency

Authorisation number:
50702

Date of authorisation status change:
5/06/2014

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

Package Leaflet

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

Summary of Product Characteristics

English (PDF)
Published on: 16/01/2026
Updated on: 14/03/2026
Download

Combined File of all Documents

Labelling

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