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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:
Denmark

Available in:
Denmark

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:
4/11/2013

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

Responsible authority:
Danish Medicines Agency

Authorisation number:
50798

Date of authorisation status change:
4/11/2013

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)

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Documents

Summary of Product Characteristics

English (PDF)

Published on: 16/01/2026
Updated on: 14/03/2026
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