

Atopica 100 mg/ml oral solution for cats and dogs

Authorised

- Ciclosporin

Product identification

Medicine name:

Atopica 100 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

100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Oral solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QL04A

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Finland

Available in:

Finland

Package description:

Multi-dose type III amber glass bottle containing 50 ml oral solution closed with a chlorobutyl rubber stopper and sealed with an aluminium tear-off cap. Each bottle is provided with two dispenser sets (consisting of a PE dip tube and a 1 ml or 4 ml polypropylene syringe) packed in a cardboard box. A polypropylene child-resistant screw cap is provided for closure of the bottle during the in-use period. Pack sizes 1 x 50 ml bottle and two dispenser sets

Multi-dose type III amber glass bottle containing 17 ml oral solution closed with a rubber stopper and sealed with a polypropylene child-resistant screw cap. One bottle and a dispenser set (consisting of a PE dip tube and a 1 ml polypropylene syringe) packed in a cardboard box. Pack sizes 1 x 17 ml and one dispenser set

Multi-dose type III amber glass bottle containing 5 ml oral solution closed with a rubber stopper and sealed with a polypropylene child-resistant screw cap. One bottle and a dispenser set (consisting of a PE dip tube and a 1 ml polypropylene syringe) packed in a cardboard box. Pack sizes 1 x 5 ml and one dispenser set

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:

Elanco GmbH

Marketing authorisation date:

15/09/2011

Manufacturing sites for batch release:

Elanco France S.A.S.

Responsible authority:

Finnish Medicines Agency

Authorisation number:

29061

Date of authorisation status change:

15/09/2011

Reference member state:

Ireland

Procedure number:

IE/V/0881/001

Concerned member states:

Austria Belgium Denmark Finland France Germany Iceland Italy
Luxembourg Netherlands Norway Portugal 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: 8/03/2023

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Package Leaflet

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