TRIDERM cutaneous spray solution for dogs

Authorised

- Marbofloxacin
- Ketoconazole
- Prednisolone

Product identification

Medicine name:
TRIDERM cutaneous spray solution for dogs

Active substance:
Marbofloxacin

Ketoconazole

Prednisolone

Target species:
Dog

Route of administration:
Cutaneous use

Product details

Active substance and strength:
Marbofloxacin

1.03 milligram(s) / 1.00 millilitre(s)

Ketoconazole
2.04 milligram(s) / 1.00 millilitre(s)

Prednisolone
0.93 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Cutaneous spray, solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QD06C

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Spain

Package description:
box containing 1 bottle of 30 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Organit Kft.

Marketing authorisation date:
11/04/2019

Manufacturing sites for batch release:

Alpha-Vet Kft.

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
3774 ESP

Date of authorisation status change:
20/03/2021

Reference member state:
Spain

Procedure number:
ES/V/0287/001

Concerned member states:
Croatia Cyprus Greece Latvia Lithuania Malta Poland Portugal

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

English (PDF)
Published on: 12/04/2023
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Package Leaflet

English (PDF)

Published on: 29/09/2025

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Labelling

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