Gefriderm cutaneous spray solution for dogs

Authorised

- Marbofloxacin
- Ketoconazole
- Prednisolone

Product identification

Medicine name:
Gefriderm 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:
QD07CA03

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

Authorisation status:
Valid

Authorised in:
Spain

Package description:
Child resistant polyethylene container with 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:
Alphavet Zrt.

Marketing authorisation date:
14/01/2022

Manufacturing sites for batch release:

Alpha-Vet Kft.

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
3978 ESP

Date of authorisation status change:
2/01/2023

Reference member state:
Spain

Procedure number:
ES/V/0371/001

Concerned member states:
Estonia Germany Ireland United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

Summary of Product Characteristics

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

Package Leaflet

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

Labelling

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

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