

Gefriderm cutaneous spray solution for dogs

Authorised

- Marbofloxacin
- Ketoconazole
- Prednisolone

Product identification

Medicine name:

Gefriderm cutaneous spray solution for dogs

Gefriderm, nahasprei, lahus koertele

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:

Estonia

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:

30/11/2020

Manufacturing sites for batch release:

Alpha-Vet Kft.

Responsible authority:

State Agency Of Medicines

Authorisation number:

2263

Date of authorisation status change:

30/11/2020

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

Summary of Product Characteristics

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

Combined File of all Documents

eu-PUAR-gefriderm-cutaneous-spray-solution-for-dogs-en.pdf