Gefriderm cutaneous spray solution for dogs

Authorised

- Marbofloxacin
- Ketoconazole
- Prednisolone

Product identification

Medicine name:
Gefriderm cutaneous spray solution for dogs
Gefriderm Spray zur Anwendung auf der Haut 1,025 mg/ml, 2,041 mg/ml, 0,926 mg/ml Lösung für Hunde

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:
Germany

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:
Alpha-Vet Kft.

Marketing authorisation date:
10/12/2020

Manufacturing sites for batch release:
Alpha-Vet Kft.

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
402696.00.00

Date of authorisation status change:
10/12/2020

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

Combined File of all Documents

English (PDF)
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