

Hydrocortisone aceponate Ecuphar 0.584 mg/ml - Cutaneous spray, solution

Authorised

- Hydrocortisone aceponate

Product identification

Medicine name:

Hydrocortisone aceponate Ecuphar 0.584 mg/ml - Cutaneous spray, solution

Active substance:

Hydrocortisone aceponate

Target species:

Dog

Route of administration:

Cutaneous use

Product details

Active substance and strength:

Hydrocortisone aceponate
0.58 milligram(s) / 1.00 Bottle

Pharmaceutical form:

Cutaneous spray, solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QD07AC

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Available in:

Belgium , Denmark , Finland , Germany , Greece , Hungary , Italy , Netherlands , Norway , Poland , Portugal , Spain , Sweden , United Kingdom (Northern Ireland)

Package description:

Packaging: Bottle (PET), Package_size: 1 bottle, Content: 76 ml

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Ecuphar

Marketing authorisation date:

27/08/2018

Manufacturing sites for batch release:

Divasa Farmavic S.A.

Responsible authority:

European Commission

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

27/08/2018

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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