File downloaded on 2026-11-16
Source URL: https://medicines.health.europa.eu/veterinary/en/600000001536

Cortavance 0.584 mg/ml - Cutaneous spray, solution

Authorised

- Hydrocortisone aceponate

Product identification

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

Package description:
Packaging:Bottle (PET), Package_size:1 bottle, Content:76 ml
Packaging:Bottle (PET), Package_size:1 bottle, Content:31 ml
Packaging:Bottle (HDPE), Package_size:1 bottle, Content:76 ml
Packaging:Bottle (HDPE), Package_size:1 bottle, Content:31 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Annex of Council Regulation (EEC) No 2309/93

Marketing authorisation holder:
Virbac

Marketing authorisation date:
9/01/2007

Manufacturing sites for batch release:
Virbac

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
9/01/2007

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

Documents

Combined File of all Documents

English (PDF)
Published on: 14/03/2026
Download

ema-puar-cortavance-v-110-par-en.pdf

ema-puar-cortavance-v-110-var-ii-0015-en.pdf