

# Cortavance 0.584 mg/ml - Cutaneous spray, solution

Authorised

- Hydrocortisone aceponate

## Product identification

**Medicine name:**

Cortavance 0.584 mg/ml - Cutaneous spray, solution

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**Active substance:**

Hydrocortisone aceponate

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**Target species:**

Dog

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**Route of administration:**

Cutaneous use

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## Product details

**Active substance and strength:**

Hydrocortisone aceponate  
0.58 milligram(s) / 1.00 Bottle

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**Pharmaceutical form:**

Cutaneous spray, solution

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**Withdrawal period by route of administration:**

**Cutaneous use:**

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**Dog**

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QD07AC

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**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)

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**Available in:**

Austria , Cyprus , Denmark , Estonia , France , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Lithuania , Norway , Portugal , Romania , Sweden , United Kingdom (Northern Ireland)

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**Package description:**

Packaging: Bottle (HDPE), Package\_size: 1 bottle, Content: 31 ml

Packaging: Bottle (HDPE), Package\_size: 1 bottle, Content: 76 ml

Packaging: Bottle (PET), Package\_size: 1 bottle, Content: 31 ml

Packaging: Bottle (PET), Package\_size: 1 bottle, Content: 76 ml

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Complete application (stand-alone) - Annex of Council Regulation (EEC) No 2309/93

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**Marketing authorisation holder:**

Virbac S.A.

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**Marketing authorisation date:**

9/01/2007

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**Manufacturing sites for batch release:**

Virbac

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**Responsible authority:**

European Commission

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

13/05/2019

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

English (PDF)

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