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# Betafuse 1 mg/g + 5 mg/g gel for dogs

Authorised

- Betamethasone valerate
- Fusidic acid hemihydrate

## Product identification

**Medicine name:**
Betafuse 1 mg/g + 5 mg/g gel for dogs

**Active substance:**
Betamethasone valerate

Fusidic acid hemihydrate

**Target species:**
Dog

**Route of administration:**
Cutaneous use

## Product details

**Active substance and strength:**
Betamethasone valerate
1.21 milligram(s) / 1.00 gram(s)

Fusidic acid hemihydrate

10.17 milligram(s) / 1.00 gram(s)

**Pharmaceutical form:**
Gel

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QD07CC01

**Legal status of supply:**
Veterinary medicinal product subject to veterinary prescription

**Authorisation status:**
Valid

**Authorised in:**
Latvia

**Package description:**
White polyethylene coated aluminium tubes of 15 g closed with a polypropylene cap.
White polyethylene coated aluminium tubes of 30 g closed with a polypropylene cap.

## 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:**
Norbrook Laboratories (Ireland) Limited

**Marketing authorisation date:**
2/08/2016

**Manufacturing sites for batch release:**
Norbrook Manufacturing Limited
Norbrook Laboratories Limited

**Responsible authority:**

Food And Veterinary Service

**Authorisation number:**
V/DCP/16/0028

**Date of authorisation status change:**
2/08/2016

**Reference member state:**
Ireland

**Procedure number:**
IE/V/0558/001

**Concerned member states:**
Austria Belgium Denmark Estonia Finland Germany Greece Italy Latvia Lithuania Netherlands Portugal Spain Sweden

United Kingdom (Northern Ireland)

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

## Documents

Summary of Product Characteristics

Combined File of all Documents

English (PDF)
Published on: 18/10/2026
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