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# DuOtic 1 mg/ml + 10 mg/ml - Ear gel

Authorised

- Betamethasone acetate
- Terbinafine

## Product identification

**Medicine name:**
DuOtic 1 mg/ml + 10 mg/ml - Ear gel

**Active substance:**
Betamethasone acetate

Terbinafine

**Target species:**
Dog

**Route of administration:**
Auricular use

## Product details

**Active substance and strength:**
Betamethasone acetate
1.00 milligram(s) / 1.00 Tube

Terbinafine

10.00 milligram(s) / 1.00 Tube

**Pharmaceutical form:**
Ear gel

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QS02CA90

**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 , Croatia , Cyprus , Czechia , Denmark , Finland , France , Germany , Greece , Hungary , Ireland , Italy , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

**Package description:**
Packaging:Tube (Alu/HDPE), Package_size:2 tubes, Content:1.2 g
Packaging:Tube (Alu/HDPE), Package_size:40 tubes, Content:1.2 g
Packaging:Tube (Alu/HDPE), Package_size:20 tubes, Content:1.2 g

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Full application - known active substance (Article 8 of Regulation (EU) 2019/6)

**Marketing authorisation holder:**
Dechra Regulatory B.V.

**Marketing authorisation date:**
22/11/2024

**Manufacturing sites for batch release:**
Genera d.d.

**Responsible authority:**
European Commission

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

**Date of authorisation status change:**
22/11/2024

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
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