

# Ototop Ear Drops and Cutaneous Suspension for Dogs, Cats and Guinea Pigs

Authorised

- POLYMYXIN B SULFATE
- Prednisolone acetate
- Miconazole nitrate

## Product identification

**Medicine name:**

Ototop Ear Drops and Cutaneous Suspension for Dogs, Cats and Guinea Pigs

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**Active substance:**

POLYMYXIN B SULFATE  
Prednisolone acetate  
Miconazole nitrate

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**Target species:**

Dog  
Cat  
Guinea pig

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**Route of administration:**

Cutaneous use  
Auricular use

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## Product details

**Active substance and strength:**

POLYMYXIN B SULFATE

5500.00 international unit(s) / 1.00 millilitre(s)

Prednisolone acetate

5.00 milligram(s) / 1.00 millilitre(s)

Miconazole nitrate

23.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Cutaneous/ear drops suspension

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

QS02CA01

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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:**

Lithuania

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**Available in:**

Lithuania

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**Package description:**

(ID1) 15 millilitre(s): Box (cardboard) with 1 Bottle (low-density polyethylene) with 15 millilitre(s), closed with Schraubdeckel (polypropylene)

(ID2) 30 millilitre(s): Box (cardboard) with 1 Bottle (low-density polyethylene) with 30 millilitre(s), closed with Schraubdeckel (polypropylene)

(ID3) 100 millilitre(s): Box (cardboard) with 1 Bottle (low-density polyethylene) with 100 millilitre(s), closed with Schraubdeckel (polypropylene)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Industrial Veterinaria S.A.

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**Marketing authorisation date:**

1/07/2020

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**Manufacturing sites for batch release:**

Industrial Veterinaria S.A.  
aniMedica Herstellungs GmbH  
aniMedica GmbH

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**Responsible authority:**

State Food And Veterinary Service

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**Authorisation number:**

LT/2/20/2602/001-003

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**Date of authorisation status change:**

3/11/2021

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0321/001

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**Concerned member states:**

Austria Belgium Cyprus Czechia Estonia Greece Hungary Italy Latvia  
Lithuania Malta Poland Portugal Romania Slovakia Slovenia Spain

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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)

Published on: 24/02/2026

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