

# 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

Ototop ausu pilieni un uz ādas lietojama suspensija suņiem, kaķiem un jūrascūciņām

### Active substance:

POLYMYXIN B SULFATE

Prednisolone acetate

Miconazole nitrate

### Target species:

Dog

Cat

Guinea pig

### Route of administration:

Cutaneous use

Auricular use

## Product details

**Active substance and strength:**

POLYMYXIN B SULFATE

0.53 milligram(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:**

Latvia

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**Package description:**

(ID3) 100 millilitre(s): Box (Cardboard) with 1 Bottle (Low Density PolyEthylene) with 100 millilitre(s), closed with Screw cap (PolyPropylene)

(ID2) 30 millilitre(s): Box (Cardboard) with 1 Bottle (Low Density PolyEthylene) with 30 millilitre(s), closed with Screw cap (PolyPropylene)

(ID1) 15 millilitre(s): Box (Cardboard) with 1 Bottle (Low Density PolyEthylene) with 15 millilitre(s), closed with Screw cap (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:**

1/06/2020

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**Manufacturing sites for batch release:**

Industrial Veterinaria S.A.

aniMedica Herstellungs GmbH

aniMedica GmbH

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**Responsible authority:**

Food And Veterinary Service

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**Authorisation number:**

V/DCP/20/0030

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**Date of authorisation status change:**

1/06/2020

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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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[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

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