

**OTOMAX- gentamicin sulfate, betamethasone valerate, and clotrimazole ointment**  
**Merck Sharp & Dohme Corp.**

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**Otomax®**  
**(GENTAMICIN SULFATE, BETAMETHASONE VALERATE AND CLOTRIMAZOLE, OINTMENT)**

Approved by FDA under NADA # 140-896

**PRODUCT INFORMATION**

**VETERINARY**

**For Otic Use in Dogs Only**

**CAUTION: Federal law restricts this drug to use by or on the order of a licensed veterinarian. Keep this and all drugs out of the reach of children.**

**DESCRIPTION** Each gram of OTOMAX ointment contains gentamicin sulfate, equivalent to 3 mg gentamicin base; betamethasone valerate, equivalent to 1 mg betamethasone; and 10 mg clotrimazole, in a mineral oil-based system containing a plasticized hydrocarbon gel.

**PHARMACOLOGY**

**Gentamicin:** Gentamicin sulfate is an aminoglycoside antibiotic active against a wide variety of pathogenic gram-negative and gram-positive bacteria. *In vitro* tests have determined that gentamicin is bactericidal and acts by inhibiting normal protein synthesis in susceptible microorganisms. Specifically, gentamicin is active against the following organisms commonly isolated from canine ears: *Staphylococcus aureus*, other *Staphylococcus* spp., *Pseudomonas aeruginosa*, *Proteus* spp., and *Escherichia coli*.

**Betamethasone:** Betamethasone valerate is a synthetic adrenocorticoid for dermatologic use. Betamethasone, an analog of prednisolone, has a high degree of corticosteroid activity and a slight degree of mineralocorticosteroid activity. Betamethasone valerate, the 17-valerate ester of betamethasone, has been shown to provide anti-inflammatory and antipruritic activity in the topical management of corticosteroid-responsive otitis externa.

Topical corticosteroids can be absorbed from normal, intact skin. Inflammation can increase percutaneous absorption. Once absorbed through the skin, topical corticosteroids are handled through pharmacokinetic pathways similar to systemically administered corticosteroids.

**Clotrimazole:** Clotrimazole is a broad-spectrum antifungal agent that is used for the treatment of dermal infections caused by various species of pathogenic dermatophytes and yeasts. The primary action of clotrimazole is against dividing and growing organisms.

*In vitro*, clotrimazole exhibits fungistatic and fungicidal activity against isolates of *Trichophyton rubrum*, *Trichophyton mentagrophytes*, *Epidermophyton floccosum*, *Microsporum canis*, *Candida* spp., and *Malassezia pachydermatis* (*Pityrosporum canis*).

Resistance to clotrimazole is very rare among the fungi that cause superficial mycoses.

In an induced otitis externa infected with *Malassezia pachydermatis*, 1% clotrimazole in the OTOMAX vehicle was effective both microbiologically and clinically in terms of reduction of exudate, odor, and swelling.

In studies of the mechanism of action, the minimum fungicidal concentration of clotrimazole caused leakage of intra-cellular phosphorus compounds into the ambient medium with concomitant breakdown of cellular nucleic acids and accelerated potassium efflux. These events began rapidly and extensively after addition of the drug.

Clotrimazole is very poorly absorbed following dermal application.

**Gentamicin-Betamethasone-Clotrimazole:** By virtue of its three active ingredients, OTOMAX ointment has antibacterial, anti-inflammatory, and antifungal activity.

In component efficacy studies, the compatibility and additive effect of each of the components were demonstrated. In clinical field trials, OTOMAX ointment was effective in the treatment of otitis externa associated with bacteria and *Malassezia pachydermatis*. OTOMAX ointment reduced discomfort, redness, swelling, exudate, and odor, and exerted a strong antimicrobial effect.

**INDICATIONS** OTOMAX ointment is indicated for the treatment of canine acute and chronic otitis externa associated with yeast (*Malassezia pachydermatis*, formerly *Pityrosporum canis*) and/or bacteria susceptible to gentamicin.

**CONTRAINDICATIONS** If hypersensitivity to any of the components occurs, treatment should be discontinued and appropriate therapy instituted. Concomitant use of drugs known to induce ototoxicity should be avoided. Do not use in dogs with known perforation of eardrums.

**WARNINGS** The use of OTOMAX ointment has been associated with deafness or partial hearing loss in a small number of sensitive dogs (eg, geriatric). The hearing deficit is usually temporary. If hearing or vestibular dysfunction is noted during the course of treatment, discontinue use of OTOMAX ointment immediately and flush the ear canal thoroughly with a nonototoxic solution.

Corticosteroids administered to dogs, rabbits, and rodents during pregnancy have resulted in cleft palate in offspring. Other congenital anomalies including deformed forelegs, phocomelia, and anasarca have been reported in offspring of dogs which received corticosteroids during pregnancy.

Clinical and experimental data have demonstrated that corticosteroids administered orally or parenterally to animals may induce the first stage of parturition if used during the last trimester of pregnancy and may precipitate premature parturition followed by dystocia, fetal death, retained placenta, and metritis.

**PRECAUTIONS** Identification of infecting organisms should be made either by microscopic roll smear evaluation or by culture as appropriate. Antibiotic susceptibility of the pathogenic organism(s) should be determined prior to use of this preparation.

If overgrowth of nonsusceptible bacteria, fungi, or yeasts occur, or if hypersensitivity develops, treatment should be discontinued and appropriate therapy instituted.

Administration of recommended doses of OTOMAX ointment beyond 7 days may result in delayed wound healing. Avoid ingestion. Adverse systemic reactions have been

observed following the oral ingestion of some topical corticosteroid preparations. Patients should be closely observed for the usual signs of adrenocorticoid overdosage which include sodium retention, potassium loss, fluid retention, weight gain, polydipsia, and/or polyuria. Prolonged use or overdosage may produce adverse immunosuppressive effects.

Use of corticosteroids, depending on dose, duration, and specific steroid, may result in endogenous steroid production inhibition following drug withdrawal. In patients presently receiving or recently withdrawn from corticosteroid treatments, therapy with a rapidly-acting corticosteroid should be considered in especially stressful situations. Before instilling any medication into the ear, examine the external ear canal thoroughly to be certain the tympanic membrane is not ruptured in order to avoid the possibility of transmitting infection to the middle ear as well as damaging the cochlea or vestibular apparatus from prolonged contact.

**TOXICOLOGY** Clinical and safety studies with OTOMAX ointment have shown a wide safety margin at the recommended dose level in dogs (see **PRECAUTIONS/SIDE EFFECTS**).

### **SIDE EFFECTS**

**Gentamicin:** While aminoglycosides are absorbed poorly from skin, intoxication may occur when aminoglycosides are applied topically for prolonged periods of time to large wounds, burns, or any denuded skin, particularly if there is renal insufficiency. All aminoglycosides have the potential to produce reversible and irreversible vestibular, cochlear, and renal toxicity.

**Betamethasone:** Side effects such as SAP and SGPT enzyme elevations, weight loss, anorexia, polydipsia, and polyuria have occurred following the use of parenteral or systemic synthetic corticosteroids in dogs. Vomiting and diarrhea (occasionally bloody) have been observed in dogs and cats.

Cushing's syndrome in dogs has been reported in association with prolonged or repeated steroid therapy.

**Clotrimazole:** The following have been reported occasionally in humans in connection with the use of clotrimazole: erythema, stinging, blistering, peeling, edema, pruritus, urticaria, and general irritation of the skin not present before therapy.

**DOSAGE AND ADMINISTRATION** The external ear should be thoroughly cleaned and dried before treatment. Remove foreign material, debris, crusted exudates, etc., with suitable nonirritating solutions. Excessive hair should be clipped from the treatment area. After verifying that the eardrum is intact, instill 4 drops from the 7.5 g and 15 g tube, and 15 g and 30 g bottle (2 drops from the 215 g bottle) of OTOMAX ointment twice daily into the ear canal of dogs weighing less than 30 lbs. Instill 8 drops from the 7.5 g and 15 g tube, and 15 g and 30 g bottle (4 drops from the 215 g bottle) twice daily into the ear canal of dogs weighing 30 lbs or more. Massage external ear canal carefully after instillation to ensure appropriate distribution of medication. Therapy should continue for 7 consecutive days.

**HOW SUPPLIED** OTOMAX ointment is available in 7.5 g and 15 g tubes as well as in 215 g, 30 g, and 15 g plastic bottles.

**Store between 2° and 25°C (36° and 77°F). Shake well before use when using**

**the 215 g bottle, 30g bottle, and 15 g bottle.**

Made in Germany

Intervet Inc. d/b/a Merck Animal Health, Madison, NJ 07940

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Rev. 9/2022

397831 R1

**PRINCIPAL DISPLAY PANEL - 7.5 g Tube Carton**

NDC 0061-0387-02

7.5 g

Otomax®  
(GENTAMICIN SULFATE,  
BETAMETHASONE  
VALERATE, AND CLOTRIMAZOLE,  
OINTMENT)

Keep Out of Reach of Children

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# 140-896

MERCK  
Animal Health

18 929896



dXB  
LOT

Leave whole  
flap varnish  
free

†98018862900 (10)

**For otic use in dogs only.**

**Indications:** OTOMAX<sup>®</sup> (gentamicin sulfate, betamethasone valerate, and clotrimazole, ointment) is indicated for the treatment of canine acute and chronic otitis externa associated with yeast (*Malassezia pachydermatis*, formerly *Pityrosporum canis*) and/or bacteria susceptible to gentamicin.

**Each gram contains:** gentamicin sulfate, equivalent to 3 mg gentamicin base; betamethasone valerate equivalent to 1 mg betamethasone; and 10 mg clotrimazole, in a mineral oil-based system containing a plasticized hydrocarbon gel.

**Dosage and Administration:** The external ear should be properly cleaned and dried before treatment. Remove foreign material, debris, crusted exudates, etc., with suitable nonirritating solutions. Excessive hair should be clipped from the treatment area. After verifying that the eardrum is intact, instill 4 drops of OTOMAX ointment twice daily into the ear canal of dogs weighing less than 30 lbs. Instill 8 drops twice daily into the ear canal of dogs weighing 30 lbs. or more. Massage external ear canal carefully after instillation to ensure appropriate distribution of medication. Therapy should continue for 7 consecutive days.

**Read accompanying directions carefully.**

**Store between 2° and 25° C (36° and 77° F).**

# Otomax<sup>®</sup>

(GENTAMICIN SULFATE, BETAMETHASONE VALERATE, AND CLOTRIMAZOLE, OINTMENT)

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Made in Germany

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NDC 0061-0387-02

7.5 g

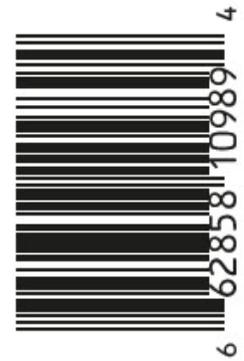
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Client \_\_\_\_\_

Patient \_\_\_\_\_

Directions: Apply \_\_\_\_\_ drops twice daily for 7 days.

PSA020042 01 R2  
32x25x125 mm  
CMYK

# Otomax<sup>®</sup>

(GENTAMICIN SULFATE, BETAMETHASONE VALERATE, AND CLOTRIMAZOLE, OINTMENT)



## OTOMAX

gentamicin sulfate, betamethasone valerate, and clotrimazole ointment

### Product Information

<b>Product Type</b>	PRESCRIPTION ANIMAL DRUG	<b>Item Code (Source)</b>	NDC:0061-0387
<b>Route of Administration</b>	AURICULAR (OTIC)		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>Gentamicin Sulfate</b> (UNII: 8X7386QRLV) (Gentamicin - UNII:T6Z9V48IKG)	Gentamicin	3 mg in 1 g
<b>Betamethasone valerate</b> (UNII: 9IFA5XM7R2) (Betamethasone - UNII:9842X06Q6M)	Betamethasone	1 mg in 1 g
<b>Clotrimazole</b> (UNII: G07GZ97H65) (Clotrimazole - UNII:G07GZ97H65)	Clotrimazole	10 mg in 1 g

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0061-0387-02	1 in 1 CARTON		
1		7.5 g in 1 TUBE		
2	NDC:0061-0387-03	1 in 1 CARTON		
2		15 g in 1 TUBE		
3	NDC:0061-0387-17	1 in 1 CARTON		
3		15 g in 1 BOTTLE, DISPENSING		
4	NDC:0061-0387-08	1 in 1 CARTON		
4		30 g in 1 BOTTLE, PLASTIC		
5	NDC:0061-0387-01	1 in 1 CARTON		
5		215 g in 1 BOTTLE, PLASTIC		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NADA	NADA140896	06/09/1993	

**Labeler** - Merck Sharp & Dohme Corp. (001317601)

