

Guide to Academy Exhibitors



AMERICAN ACADEMY
OF OPHTHALMOLOGY

The Eye M.D. Association



AAO-APAO
CHICAGO
2012

NOVEMBER 10-13

IMPORTANT MEETING DATES, HOURS AND LOCATIONS

EXHIBITION HOURS

Location: South Building, Level 3, Hall A
 Sat. Nov. 10 9 a.m. to 5 p.m.
 Sun. Nov. 11 9 a.m. to 5 p.m.
 Mon. Nov. 12 9 a.m. to 5 p.m.
 Tues. Nov. 13 9 a.m. to 3 p.m.

REGISTRATION LOCATION AND HOURS

Location: South Building, Level 3, Hall A
 Thurs. Nov. 8 4 to 6 p.m.
 Fri. Nov. 9 7 a.m. to 5 p.m.
 Sat. Nov. 10 7 a.m. to 5 p.m.
 Sun. Nov. 11 8 a.m. to 5 p.m.
 Mon. Nov. 12 8 a.m. to 5 p.m.
 Tues. Nov. 13 8 a.m. to 3 p.m.

SHUTTLE HOURS

Fri. Nov. 9 6:30 to 9:30 a.m.
 and 3 to 6 p.m.
 Sat. Nov. 10 6:30 a.m. to 6:30 p.m.
 Sun. Nov. 11 6:30 a.m. to 6:30 p.m.
 Mon. Nov. 12 6:30 a.m. to 6:30 p.m.
 Tues. Nov. 13 6:30 a.m. to 6 p.m.

During peak hours shuttles will run every 10 to 15 minutes.

During non-peak hours shuttles will run every 20 to 25 minutes.

More details are available on pages 23 and 24 of the Pocket Guide.

SUBSPECIALTY DAY LOCATIONS AND HOURS

Cornea

Grand Ballroom, S100ab
 Sat. Nov. 10 8 a.m. to 5:30 p.m.

Glaucoma

E354
 Sat. Nov. 10 8 a.m. to 5:30 p.m.

Oculofacial Plastic Surgery

Vista Room S406a
 Sat. Nov. 10 8 a.m. to 5:17 p.m.

Pediatric Ophthalmology

Grand Ballroom S100c
 Sat. Nov. 10 8 a.m. to 5:15 p.m.

Refractive Surgery

North Hall B (Sections I-X, Fri. and Sat.)
 Grand Ballroom S100ab (Free Papers, Fri. Only)
 Fri. Nov. 9 8 a.m. to 7:30 p.m.
 Sat. Nov. 10 8 a.m. to 5:24 p.m.

Retina

Arie Crown Theater
 Fri. Nov. 9 8 a.m. to 5:04 p.m.
 Sat. Nov. 10 8 a.m. to 5:22 p.m.

IMPORTANT MEETING LOCATIONS

All rooms are in the McCormick Place, North Building (N), South Building (S), and Lakeside Center (E). The Exhibition is on Level 3, Hall A, in the South Building.

AAOasis (Wi-Fi HotSpot) Booth 2981
 AAOE Coding Sessions Room S105
 AAOE Member Lounge South, Level 5
 AAOE Practice Management Center Academy Resource Center (Booth 508)
 AAOE Program South, Level 5
 Academy Café Room S406b
 Academy Resource Center Booth 508
 Badge Replacement Registration Services (Hall A)
 Bags and Programs Hall A
 Bistro AAO Booth 2490
 Breakfast With the Experts Roundtables Hall A
 Business Center South, Level 2.5
 New Orleans 2013 Grand Concourse, Level 2.5
 CME Reporting/Proof-of-Attendance Grand Concourse Level 2.5 & Academy Resource Center (Booth 508)
 Coat & Bag Check South, Level 1 Lobby
 E-mail & Internet Access Grand Concourse & Booth 2987
 Executive Office Room S401
 Exhibitor Locator Booth 3500
 Exhibitor Lounge (Wi-Fi available) Booth 987
 Exhibitor Service Center/Exhibitions Office Hall A
 ExpoCard Replacement Registration Services (Hall A)
 First Aid South, Level 2.5
 Foundation of the Academy Academy Resource Center (Booth 508)
 Global Alliances Office Room S403a
 Hotel Assistance Grand Concourse, Level 2.5
 Informational Exhibits & Posters Hall A
 International Center and Outreach Booth 4509
 Learning Lounge Booth 107
 Lost & Found Meetings Office (Room S402)
 Meditation/Prayer Room Room SA1a
 Meeting Information Grand Concourse & South, Level 1 Lobby
 Meetings Office Room S402
 Mobile Meeting Guide AAO Connect (Booth 2987)
 Museum of Vision Booth 704
 Newsroom Room N426a
 Ophthalmic Mutual Insurance Company (OMIC) Booth 1104
 Ophthalmology Job Center Room N426c
 Proof-of-Attendance/CME Reporting Grand Concourse, Level 2.5 & Academy Resource Center (Booth 508)
 Publishers' Row Hall A
 Reconnect Booth 2981
 Registration (Attendees) Hall A
 Registration (Exhibitors) Hall A
 Ribbons Bags & Programs (Hall A)
 Scientific Posters Hall A
 Scientific Poster Tours Meeting Point, Hall A
 Senior Ophthalmologist (SO) Lounge Grand Concourse Lobby
 Shuttle Buses South, Level 1 Lobby and North, Level 1, Gate 26
 Speaker Ready Room Grand Concourse Lobby
 Technology Pavilion Booth 880
 The Electronic Office (IHE) Booth 114
 Ticketed Events and Tour Sales Hall A
 Tour Program Departures South, Level 1 Lobby
 Videos on Demand/
 Scientific Posters Online Booth 165
 Wi-Fi Access Booth 2981
 Young Ophthalmologist (YO) Lounge Grand Concourse Lobby

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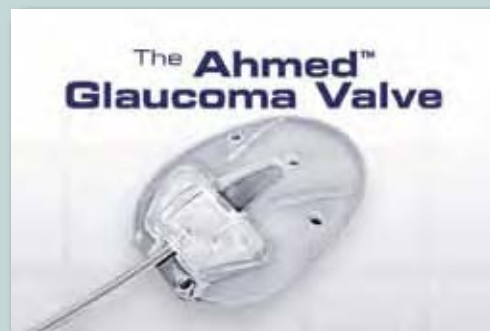
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Letter From the Publisher

Dear Meeting Attendee,

Welcome to Chicago! With so much to see and do during the 2012 Joint Meeting, I am pleased to present you with *EyeNet's Guide to Academy Exhibitors*. Designed to maximize your time on the show floor, the *Guide's* alphabetical business listings will help you quickly identify the companies you wish to visit and in many cases see at a glance their new products and services. Its foldout map will aid navigation of the exhibit floor, and note pages at the back give you space to plot your course through the hall or take notes during vendor meetings.

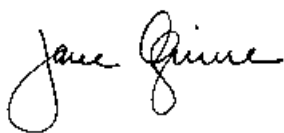
EyeNet is the Academy's monthly member magazine that delivers trusted clinical information that you can implement in your practice immediately. Look out for other *EyeNet* publications that you will find similarly useful during the meeting and throughout the year:

- **Academy News**—convention hall tabloid available Friday and Sunday, Nov. 9 and 11
- **Academy Live**—e-mail blasts bringing you up-to-the-minute Annual Meeting coverage
- **EyeNet Extras**—special topic-focused supplements

And, as always, all of *EyeNet's* articles are online at www.eyenet.org, where you can post questions and leave feedback using the Comments feature.

On behalf of all of us at *EyeNet*, we hope your time at the Joint Meeting is productive and enjoyable.

Sincerely,



Jane Aguirre, CAE
Publisher
EyeNet Magazine

Disclaimer

The product information and claims provided in the profiles included in *EyeNet's Guide to Academy Exhibitors* are those of the manufacturers and have not been verified, nor does the appearance of a product constitute an endorsement by *EyeNet* or the American Academy of Ophthalmology. This guide went to press on Sept. 20, 2012. Please check www.aao.org for updates to Exhibit Hall booths.

David W. Parke II, MD

Editor-in-Chief

Richard P. Mills, MD, MPH

Chief Medical Editor

Jane Aguirre, CAE

Publisher

Patty Ames

Executive Editor

Carey S. Ballard

Art Director / Production Manager

Peggy Denny

Chris McDonagh

Senior Editors

Contributing Writers

Barbara Boughton
Leslie Burling-Phillips
Marianne Doran
Miriam Karmel
Linda Roach
Lori Baker Schena, EdD
Jean Shaw
Annie Stuart

Business Manager/ Assistant Editor

Catherine Morris

Advertising Sales

Kelly Miller
Mark Mrvica

M.J. Mrvica Associates, Inc.

2 West Taunton Ave., Berlin, NJ 08009

Phone 856-768-9360; Fax 856-753-0064

kmiller@mrsvica.com

markmrsvica@mrsvica.com



**AMERICAN ACADEMY
OF OPHTHALMOLOGY**

The Eye M.D. Association

655 Beach Street
San Francisco, CA 94109

P.O. Box 7424
San Francisco, CA 94120

Phone: 866-561-8558 or 415-561-8500

Fax: 415-561-8575

E-mail: eyenet@aao.org

Online: www.aao.org and www.eyenet.org

Governmental Affairs Division

20 F Street NW, Suite 400,
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800-729-1959	www.aitindustries.com

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AKtive Srl	Booth 2575
011-39-0686-2155-40	www.aktive-corp.com

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011-39-4989-6207-4	www.alchimiasrl.com

Alcon Laboratories, Inc.	Booth 2808
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Ambler Surgical	Booth 2337
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What Resources Exist for International Attendees?

International attendees are encouraged to visit the **International Center (Booth 4509)**, an ideal place to learn more about Academy programs, check e-mail, meet or leave messages for colleagues, or just relax. An interpreter will be available and light refreshments will be served. Admittance is by attendee badge.

American Academy of Ophthalmic Executives (AAOE) Booth 508

The **American Academy of Ophthalmic Executives (AAOE)** is the practice management arm of the Academy.

Visit the AAOE Practice Management Center in the Resource Center for help with critical business issues like coding, PQRS, e-prescribing, EHR, and more.

The Center is also the place where you can meet with a practice management consultant for a free 20-minute consultation and preview practice management products and services.



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American Academy of Ophthalmology (AAO) Booth 508**See What's New at the Academy Resource Center**

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- Demos of the ONE[®] Network, and EyeSmart[®] patient education website
- Academy, ISRS, and AAOE membership
- Video Production Services studio
- Free consultations with practice management experts
- Advice on coding, e-prescribing, and PQRS
- Information on EyeCare America and the Academy Foundation
- CME credit reporting stations
- Enter to win an iPad loaded with Academy products in one of four daily raffles!
- Get your copy of *Cutting for Stone* signed by Abraham Verghese, MD, MACP, after Sunday's Opening Session.



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American Academy of Ophthalmology—Ophthalmology Journal Booth 3572

Ophthalmology, the official journal of the American Academy of Ophthalmology, publishes original, peer-reviewed reports of research in ophthalmology, including basic science investigations and clinical studies. Topics include new diagnostic and surgical techniques, treatment methods, instrument updates, the latest drug findings, results of clinical trials, and research findings.

Ophthalmology also publishes major reviews of specific topics by acknowledged authorities.

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American Express OPEN Booth 1574

212-640-2000

www.open.com

American Optisurgical, Inc. Booth 3100

949-580-1266

www.optisurgical.com

American Society of Cataract & Refractive Surgery (ASCRS) Booth 3262

703-591-2220

www.ascrs.org

American Society of Ophthalmic Registered Nurses (ASORN) Booth 2782

415-561-8513

www.asorn.org

Anadolu Tip A.S. Booth 1779

011-90-346-218-1418

www.zarcomplenses.com

Angie's List Booth 2576

866-543-5478

www.angieslist.com

Angiotech Booth 1151

800-523-3332

www.angiotech.com

Anodyne Surgical Booth 2550

800-428-5628

www.anodynesurgical.com

AOLab - American Ophthalmic Lab Booth 2700

410-647-5140

www.aolab.com

Appasamy Associates Booth 557

718-656-7390

www.appasamy.com

Apramed Medical Devices Booth 4337

011-55-16-3306-1122

www.apramed.com.br

ARCADOPHTA Booth 1949

011-33-5-6140-5235

www.arcadophta.com

ArcticDx, Inc. Booth 2076

866-964-5182

www.macularisk.com

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800-847-5786

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Army Medical Recruiting Booth 2276

502-626-1981

www.goarmy.com

Art Optical Contact Lens, Inc. Booth 2075

616-453-1888

www.artoptical.com

Asia-Pacific Academy of Ophthalmology Booth 1200

011-852-3943-5927

www.apaophth.org

ASICO, LLC	Booth 3300
630-986-8032	www.asico.com

Association for Research in Vision and Ophthalmology (ARVO)	Booth 2046
240-221-2900	www.arvo.org

Audio Digest	Booth 245
818-240-7500	www.audiodigest.org

Aumed Group Corp.	Booth 433
011-86-1082-8841-73	www.aumedgroup.com

Aurolab	Booth 3276
011-91-4523-0961-00	www.aurolab.com

Aurora Surgical, LLC	Booth 3170
727-821-3303	www.aurorasurgical.com

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Bausch + Lomb	Booth 3126
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Bayer Healthcare Pharmaceuticals	Booth 153
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BJB Medical Associates	Booth 159
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Bruder Healthcare Company	Booth 1774
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Bryn Mawr Communications LLC	Booth 662
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Cambrian Medical, Inc.	Booth 2450
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Canon U.S.A., Inc.	Booth 2009
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Chace and Associates	Booth 4071
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For patients with elevated intraocular pressure (IOP)
in open-angle glaucoma (OAG) or ocular hypertension (OHT)

START WITH ZIOPTAN

6–8 mmHg
at month 3

5–8 mmHg
at month 6

POWERFUL IOP REDUCTIONS

> Based on clinical studies of up to 24 months in 905 patients with a baseline pressure of 23–26 mmHg.

Once-daily, single-use containers

Preservative-free formulation

ZIOPTAN is indicated for reducing elevated IOP in patients with OAG or OHT.

SELECT IMPORTANT SAFETY INFORMATION

ZIOPTAN has been reported to cause changes to pigmented tissues. The most frequently reported changes have been to the iris, periorbital tissue (eyelid), and eyelashes. Pigmentation is expected to increase as long as ZIOPTAN is administered. Pigmentation of the iris is likely to be permanent and may not be noticeable for several months to years, while pigmentation of the periorbital tissue and eyelash changes may be reversible in some patients. The long-term effects of increased pigmentation are not known.

ZIOPTAN may gradually change eyelashes and vellus hair in the treated eye. These changes include increased length, color, thickness, shape, and number of lashes. Eyelash changes are usually reversible on discontinuation of treatment.

ZIOPTAN should be used with caution in patients with active intraocular inflammation (eg, iritis/uveitis) because the inflammation may be exacerbated.

Macular edema, including cystoid macular edema, has been reported during treatment with prostaglandin F_{2α} analogs. ZIOPTAN should be used with caution in aphakic patients, in pseudophakic patients with a torn posterior lens capsule, or in patients with known risk factors for macular edema.

In clinical trials of patients receiving either preservative-containing or preservative-free ZIOPTAN, the most common pooled adverse reaction observed was conjunctival hyperemia, which was reported in a range of 4% to 20% of patients.

Please see the adjacent Brief Summary of the Prescribing Information.

Brief Summary of the Prescribing Information for ZIOPTAN.

INDICATIONS AND USAGE

ZIOPTAN is indicated for reducing elevated intraocular pressure in patients with open-angle glaucoma or ocular hypertension.

DOSAGE AND ADMINISTRATION

The recommended dose is 1 drop of ZIOPTAN in the conjunctival sac of the affected eye(s) once daily in the evening.

The dose should not exceed once daily since it has been shown that more frequent administration of prostaglandin analogs may lessen the intraocular pressure-lowering effect.

Reduction of the intraocular pressure starts approximately 2 to 4 hours after the first administration with the maximum effect reached after 12 hours.

ZIOPTAN may be used concomitantly with other topical ophthalmic drug products to lower intraocular pressure. If more than 1 topical ophthalmic product is being used, each 1 should be administered at least 5 minutes apart.

The solution from 1 individual unit is to be used immediately after opening for administration to 1 or both eyes. Since sterility cannot be maintained after the individual unit is opened, the remaining contents should be discarded immediately after administration.

CONTRAINDICATIONS

None.

WARNINGS AND PRECAUTIONS

Pigmentation

Tafluprost ophthalmic solution has been reported to cause changes to pigmented tissues. The most frequently reported changes have been increased pigmentation of the iris, periorbital tissue (eyelid), and eyelashes. Pigmentation is expected to increase as long as tafluprost is administered. The pigmentation change is due to increased melanin content in the melanocytes rather than to an increase in the number of melanocytes. After discontinuation of tafluprost, pigmentation of the iris is likely to be permanent, while pigmentation of the periorbital tissue and eyelash changes have been reported to be reversible in some patients. Patients who receive treatment should be informed of the possibility of increased pigmentation. The long-term effects of increased pigmentation are not known.

Iris color change may not be noticeable for several months to years. Typically, the brown pigmentation around the pupil spreads concentrically towards the periphery of the iris and the entire iris or parts of the iris become more brownish. Neither nevi nor freckles of the iris appear to be affected by treatment. While treatment with ZIOPTAN can be continued in patients who develop noticeably increased iris pigmentation, these patients should be examined regularly. [See Patient Counseling Information.]

Eyelash Changes

ZIOPTAN may gradually change eyelashes and vellus hair in the treated eye. These changes include increased length, color, thickness, shape, and number of lashes. Eyelash changes are usually reversible upon discontinuation of treatment.

Intraocular Inflammation

ZIOPTAN should be used with caution in patients with active intraocular inflammation (eg, iritis/uveitis) because the inflammation may be exacerbated.

Macular Edema

Macular edema, including cystoid macular edema, has been reported during treatment with prostaglandin F_{2α} analogs. ZIOPTAN should be used with caution in aphakic patients, in pseudophakic patients with a torn posterior lens capsule, or in patients with known risk factors for macular edema.

ADVERSE REACTIONS

Clinical Studies Experience

Because clinical studies are conducted under widely varying conditions, adverse reaction rates observed in the clinical studies of a drug cannot be directly compared to rates in the clinical studies of another drug and may not reflect the rates observed in practice.

Preservative-containing or preservative-free tafluprost 0.0015% was evaluated in 905 patients in 5 controlled clinical studies of up to 24-months' duration. The most common adverse reaction observed in patients treated with tafluprost was conjunctival hyperemia which was reported in a range of 4% to 20% of patients. Approximately 1% of patients discontinued therapy due to ocular adverse reactions.

Ocular adverse reactions reported at an incidence of ≥2% in these clinical studies included ocular stinging/irritation (7%), ocular pruritus including allergic conjunctivitis (5%), cataract (3%), dry eye (3%), ocular pain (3%), eyelash darkening (2%), growth of eyelashes (2%), and blurred vision (2%).

Nonocular adverse reactions reported at an incidence of 2% to 6% in these clinical studies in patients treated with tafluprost 0.0015% were headache (6%), common cold (4%), cough (3%), and urinary tract infection (2%).

Postmarketing Experience

The following adverse reactions have been identified during postapproval use of tafluprost. Because postapproval adverse reactions are reported voluntarily from a population of uncertain size, it is not always possible to reliably estimate their frequency or establish a causal relationship to drug exposure.

In postmarketing use with prostaglandin analogs, periorbital and lid changes, including deepening of the eyelid sulcus, have been observed.

USE IN SPECIFIC POPULATIONS

Pregnancy

Pregnancy Category C.

Teratogenic effects: In embryo-fetal development studies in rats and rabbits, tafluprost administered intravenously was teratogenic. Tafluprost caused increases in post-implantation losses in rats and rabbits and reductions in fetal body weights in rats. Tafluprost also increased the incidence of vertebral skeletal abnormalities in rats and the incidence of skull, brain, and spine malformations in rabbits. In rats, there were no adverse effects on embryo-fetal development at a dose of 3 µg/kg/day corresponding to maternal plasma levels of tafluprost acid that were 343 times the maximum clinical exposure based on C_{max}. In rabbits, effects were seen at a tafluprost dose of 0.03 µg/kg/day corresponding to maternal plasma levels of tafluprost acid during organogenesis that were approximately 5 times higher than the clinical exposure based on C_{max}. At the no-effect dose in rabbits (0.01 µg/kg/day), maternal plasma levels of tafluprost acid were below the lower level of quantification (20 pg/mL).

In a pre- and postnatal development study in rats, increased mortality of newborns, decreased body weights, and delayed pinna unfolding were observed in offspring. The no observed adverse effect level was at a tafluprost intravenous dose of 0.3 µg/kg/day, which is greater than 3 times the maximum recommended clinical dose based on body surface area comparison.

There are no adequate and well-controlled studies in pregnant women. Although animal reproduction studies are not always predictive of human response, ZIOPTAN should not be used during pregnancy unless the potential benefit justifies the potential risk to the fetus.

Women of childbearing age/potential should have adequate contraceptive measures in place.

Nursing Mothers

A study in lactating rats demonstrated that radio-labeled tafluprost and/or its metabolites were excreted in milk. It is not known whether this drug or its metabolites are excreted in human milk. Because many drugs are excreted in human milk, caution should be exercised when ZIOPTAN is administered to a nursing woman.

Pediatric Use

Use in pediatric patients is not recommended because of potential safety concerns related to increased pigmentation following long-term chronic use.

ZIOPTAN™ (tafluprost ophthalmic solution) 0.0015%

Geriatric Use

No overall clinical differences in safety or effectiveness have been observed between elderly and other adult patients.

PATIENT COUNSELING INFORMATION

See FDA-Approved Patient Labeling (Patient Information).

Nightly Application

Patients should be advised to not exceed once-daily dosing since more frequent administration may decrease the intraocular pressure-lowering effect of ZIOPTAN.

Handling the Single-Use Container

Patients should be advised that ZIOPTAN is a sterile solution that does not contain a preservative. The solution from 1 individual unit is to be used immediately after opening for administration to 1 or both eyes. Since sterility cannot be maintained after the individual unit is opened, the remaining contents should be discarded immediately after administration.

Potential for Pigmentation

Patients should be advised about the potential for increased brown pigmentation of the iris, which may be permanent. Patients should also be informed about the possibility of eyelid skin darkening, which may be reversible after discontinuation of ZIOPTAN.

Potential for Eyelash Changes

Patients should also be informed of the possibility of eyelash and vellus hair changes in the treated eye during treatment with ZIOPTAN. These changes may result in a disparity between eyes in length, thickness, pigmentation, number of eyelashes or vellus hairs, and/or direction of eyelash growth. Eyelash changes are usually reversible upon discontinuation of treatment.

When to Seek Physician Advice

Patients should be advised that if they develop a new ocular condition (eg, trauma or infection), experience a sudden decrease in visual acuity, have ocular surgery, or develop any ocular reactions, particularly conjunctivitis and eyelid reactions, they should immediately seek their physician's advice concerning the continued use of ZIOPTAN.

Use with Other Ophthalmic Drugs

If more than 1 topical ophthalmic drug is being used, the drugs should be administered at least five (5) minutes between applications.

Storage Information

Patients should be instructed on proper storage of cartons, unopened foil pouches, and opened foil pouches [see How Supplied/Storage and Handling]. Recommended storage for cartons and unopened foil pouches is to store refrigerated at 2-8°C (36-46°F). After the pouch is opened, the single-use containers may be stored in the opened foil pouch for up to 28 days at room temperature: 20-25°C (68-77°F). Protect from moisture.

For more detailed information, please read the Prescribing Information.

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Meet the Author — Abraham Verghese, MD, MACP



After presenting the Keynote Address at Sunday's Opening Session (8:30 to 10 a.m.), Dr. Verghese will stroll over to the Academy Resource Center (Booth 508), where he will be signing copies of *Cutting for Stone*, which spent more than two years on the *The New York Times* best sellers list.

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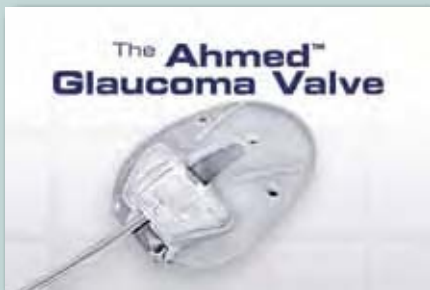
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ACTION: The antibiotic, Bacitracin, exerts a profound action against many gram positive pathogens, including the common Streptococci and Staphylococci. It is also destructive for certain gram negative organisms. It is ineffective against fungi.

INDICATIONS: For the treatment of superficial ocular infections involving the conjunctiva and/or cornea caused by Bacitracin susceptible organisms.

CONTRAINDICATIONS: This product should not be used in patients with a history of hypersensitivity to Bacitracin.

PRECAUTIONS: Bacitracin ophthalmic ointment should not be used in deep-seated ocular infections or in those that are likely to become systemic. The prolonged use of antibiotic containing preparations may result in overgrowth of nonsusceptible organisms particularly fungi. If new infections develop during treatment appropriate antibiotic or chemotherapy should be instituted.

ADVERSE REACTIONS: Bacitracin has such a low incidence of allergenicity that for all practical purposes side reactions are practically non-existent. However, if such reaction should occur, therapy should be discontinued.

DOSAGE AND ADMINISTRATION: The ointment should be applied directly into the conjunctival sac 1 to 3 times daily. In blepharitis all scales and crusts should be carefully removed and the ointment then spread uniformly over the lid margins. Patients should be instructed to take appropriate measures to avoid gross contamination of the ointment when applying the ointment directly to the infected eye.

HOW SUPPLIED: 3.5 g (1/8 Oz) sterile tamper proof tubes, NDC 48102-007-35.



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Bacitracin ophthalmic ointment should not be used in deep-seated ocular infections or in those that are likely to become systemic.

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