SUNDAY, OCTOBER 23, 2011

The Peabody Hotel – Orlando
9801 International Drive
Orlando, Florida 32819

OPTIMIZING OUTCOMES WITH
TECNIS® MULTIFOCAL IOL

SUNDAY, OCTOBER 23, 2011
Registration and Breakfast: .......................... 6:30-7:00am
Presentations and Discussion: ..................... 7:00-8:30am

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Orlando, Florida 32819

6:30-7 am REGISTRATION AND BREAKFAST
7:00-7:30 SECTION 1: It's all About the Outcomes:
What's New with the TECNIS® Multifocal IOL?
Moderator: Elizabeth Davis, MD
Ehsan Sadri, MD
Mitch Jackson, MD
Larry Katzen, MD

7:30-8:00 SECTION 2: Practical Uses of the
TECNIS® Multifocal IOL
Moderator: William Trattler, MD
Daniel Chang, MD
D. Rex Hamilton, MD
John Hovanesian, MD
Ming Wang, MD

8:00-8:30 SECTION 3: Using Premium IOLs to Grow
Your Cataract Practice
Moderator: Robert Maloney, MD
William Trattler, MD
Farrell C. Tyson, MD
D. Rex Hamilton, MD

3 Ways to Register!
Call: 877-451-6511
Email: ReviewMeetings@Jobson.com

Important Safety Information for TECNIS® Multifocal IOL
Indications: TECNIS® multifocal intraocular lenses are indicated for visual correction of aphakia in adult patients with and without
presbyopia in whom a cataractous lens has been removed by phacoemulsification. The intraocular lenses are intended to be placed in the
capsular bag. Warnings: Physicians considering lens implantation under any of the conditions described in the Directions for Use labeling should
weigh the potential risk/benefit ratio prior to implanting a lens. Precautions: Do not resterilize or autoclave. Use only sterile irrigating solutions such
as balanced salt solution or sterile normal saline. Do not store in direct sunlight or over 45°C.
Adverse Events: The most frequently reported adverse
event that occurred during the clinical trial of the Tecnis® Multifocal lens was surgical reintervention, which occurred at a rate of 3.7% (lens-related:
0.6%; non-lens related: 3.2%) which included lens exchange, retinal repair, iris prolapse/wound repair, trabeculectomy, lens repositioning, and lens
removal due to patient dissatisfaction. Other events included macular edema (2.6%), hypopyon (0.3%) and endophthalmitis (0.3%).
Caution: Federal law restricts this device to sale by or on the order of a physician.

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