IDENTIFIED PRACTICE GAPS/EDUCATIONAL NEEDS
As medicine moves increasingly toward evidence-based care, treatment practices and adherence to stan-
dard protocols is not optimal in Otolaryngology. There is a lack of knowledge of current best evidence and,
even when evidence is known, there is a lack of consistent implementation in practice of the best practices
or standard protocols. This course is an update course on best practices, best evidence and new technolo-
gies and their application to patient care.

TARGET AUDIENCE
This course is designed for practicing otolaryngologists, fellows and residents in training.

EDUCATIONAL OBJECTIVES
The objective of any CME program is to improve patient care. Upon completion of this course, participants
will be able to:
• Identify current evaluation and treatment for sleep apnea.
• Analyze the current evaluation, medical and surgical treatments for chronic sinusitis.
• Evaluate the latest treatment for tonsil and adenoid disease, including surgical techniques.
• Analyze the diagnostic evaluation of patients with dizziness.
• Identify the current role of surgery in the management of patients with head and neck cancer.

ACCREDITATION AND CREDIT DESIGNATION STATEMENTS
Weill Cornell Medical College is accredited by the Accreditation Council for Continuing Medical Education
to provide continuing medical education for physicians.

Weill Cornell Medical College designates this live activity for a maximum of 13.75 AMA PRA Category 1
Credit(s)™. Physicians should claim only the credit commensurate with the extent of their participation in
the activity.

DISCLOSURE OF RELATIONSHIPS/CONTENT VALIDITY
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scientific rigor in all its sponsored activities. All speakers, Course Directors, Co-Course Directors,
planners, reviewers, and staff members participating in sponsored activities are expected to disclose
relevant financial relationships pertaining to their contribution to the activity. Relationship information is
analyzed to determine whether conflicts of interest exist. All conflicts of interest are resolved prior to
participation in the planning or implementation of this activity. Presenters and authors are also expected
to disclose any discussion of (1) off-label or investigational uses of FDA approved commercial products or
devices or (2) products or devices not yet approved in the United States.