

# bulletin

American Academy of Otolaryngology—Head and Neck Surgery

March 2013—Vol.32 No.03

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See the World Voice Day Poster  
included in this mailing



**New**

# **DYMISTA™**

(azelastine hydrochloride and fluticasone propionate) Nasal Spray  
137 mcg / 50 mcg per Spray

# for rapid and

## **Indication**

Dymista Nasal Spray, containing an H<sub>1</sub>-receptor antagonist and a corticosteroid, is indicated for the relief of symptoms of seasonal allergic rhinitis in patients 12 years of age and older who require treatment with both azelastine hydrochloride and fluticasone propionate for symptomatic relief.

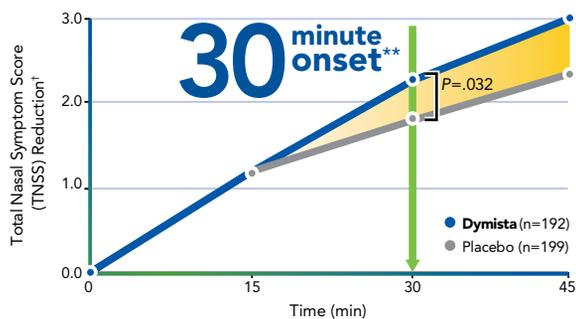
## **Important Risk Information**

- Patients may experience somnolence. Caution patients against engaging in hazardous occupations requiring complete mental alertness, such as driving or operating machinery
- Patients should avoid concurrent use of alcohol or other central nervous system (CNS) depressants because additional reductions in alertness and additional impairment of CNS performance may occur
- Because of the inhibitory effect of corticosteroids on wound healing, avoid use in patients with recent nasal ulcers, nasal surgery, or nasal trauma until healed
- Glaucoma, cataracts, and increased intraocular pressure may be associated with nasal corticosteroid use; therefore, close monitoring is warranted in patients with a change in vision and/or with a history of increased intraocular pressure, glaucoma, and/or cataracts
- Patients using corticosteroids may be susceptible to infections and may experience a more serious or even fatal course of chicken pox or measles. Dymista should be used with caution in patients with active or quiescent tuberculosis; fungal, bacterial, viral, or parasitic infections; or ocular herpes simplex
- Systemic corticosteroid effects, such as hypercorticism and adrenal suppression, may occur with very high dosages or at the regular dosage in susceptible individuals. If such changes occur, discontinue Dymista gradually, under medical supervision
- Potent inhibitors of cytochrome P450 (CYP) 3A4 may increase blood levels of fluticasone propionate
- Ritonavir: coadministration is not recommended
- Other potent CYP3A4 inhibitors, such as ketoconazole: use caution with coadministration
- Intranasal corticosteroids may cause a reduction in growth velocity when administered to pediatric patients. Monitor the growth routinely of pediatric patients receiving Dymista
- In clinical trials, the most common adverse reactions that occurred with Dymista Nasal Spray, azelastine hydrochloride nasal spray, fluticasone nasal spray, and vehicle placebo groups, respectively, were dysgeusia (4%, 5%, 1%, <1%), epistaxis (2% for each group), and headache (2%, 2%, 2%, and 1%)
- Pregnancy Category C: based on animal data; may cause fetal harm

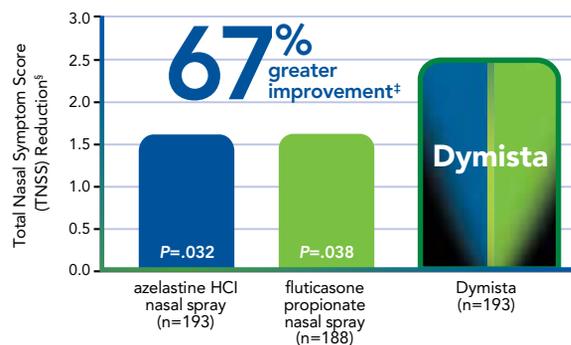
# more complete relief

## from seasonal allergy symptoms

### Nasal Symptom Reduction: Statistically Superior at 30 Minutes<sup>\*1,2</sup>



### Magnitude of Nasal Symptom Relief Relative to azelastine HCl and to fluticasone propionate<sup>\*1,2</sup>



Data shown are from study MP 4004. Across the 3 pivotal clinical trials, the improvement with Dymista ranged from 40% to 67% greater relative to the improvement achieved with either comparator.<sup>1,2</sup>

\*As listed in the Full Prescribing Information, in 3 pivotal trials, symptom relief was measured by change from baseline in Total Nasal Symptom Score (TNSS) averaged over the 14-day study period. Dymista provided a statistically significant improvement in TNSS compared with both azelastine hydrochloride (HCl) and fluticasone propionate. The azelastine HCl and fluticasone propionate comparators used the same device and vehicle as Dymista and are not commercially marketed. Additionally, Dymista provided a statistically significant, rapid improvement in TNSS as early as 30 minutes after administration when compared with placebo.<sup>1</sup>

\*\*Data shown are from study MP 4004. Onset of action was defined as the first timepoint at which Dymista was statistically superior to placebo in the mean change from baseline in instantaneous TNSS and was sustained thereafter.<sup>1</sup>

†Change from baseline in instantaneous TNSS following administration.<sup>2</sup>

‡Percent difference represents the improvement in TNSS with Dymista relative to azelastine HCl or fluticasone propionate comparator.<sup>2</sup>

§Change from baseline in the placebo-subtracted mean TNSS for each day (maximum score 24), averaged over the 14-day study period.<sup>2</sup>

References: 1. Dymista [package insert]. Somerset, NJ: Meda Pharmaceuticals Inc; 2012.  
2. Data on File. Meda Pharmaceuticals Inc.

Please see Brief Summary of Full Prescribing Information on the following pages.

**DYMISTA™**  
(azelastine hydrochloride and  
fluticasone propionate) Nasal Spray  
137 mcg / 50 mcg per Spray

[www.Dymista.com](http://www.Dymista.com)

# DYMISTA (AZELASTINE HYDROCHLORIDE 137 MCG / FLUTICASONE PROPIONATE 50 MCG) NASAL SPRAY

## Brief Summary (for Full Prescribing Information, see package insert)

### 1 INDICATIONS AND USAGE

Dymista Nasal Spray is indicated for the relief of symptoms of seasonal allergic rhinitis in patients 12 years of age and older who require treatment with both azelastine hydrochloride and fluticasone propionate for symptomatic relief.

### 5 WARNINGS AND PRECAUTIONS

#### 5.1 Somnolence

In clinical trials, the occurrence of somnolence has been reported in some patients (6 of 853 patients) taking Dymista Nasal Spray [see *Adverse Reactions* (6.1)]. Patients should be cautioned against engaging in hazardous occupations requiring complete mental alertness and motor coordination such as operating machinery or driving a motor vehicle after administration of Dymista Nasal Spray. Concurrent use of Dymista Nasal Spray with alcohol or other central nervous system depressants should be avoided because additional reductions in alertness and additional impairment of central nervous system performance may occur [see *Drug Interactions* (7.1)].

#### 5.2 Local Nasal Effects

In clinical trials of 2 to 52 weeks' duration, epistaxis was observed more frequently in patients 38 treated with Dymista Nasal Spray than those who received placebo [see *Adverse Reactions* (6)].

Instances of nasal ulceration and nasal septal perforation have been reported in patients following the intranasal application of corticosteroids. There were no instances of nasal ulceration or nasal septal perforation observed in clinical trials with Dymista Nasal Spray. Because of the inhibitory effect of corticosteroids on wound healing, patients who have experienced recent nasal ulcers, nasal surgery, or nasal trauma should not use Dymista Nasal Spray until healing has occurred. In clinical trials with fluticasone propionate administered intranasally, the development of localized infections of the nose and pharynx with *Candida albicans* has occurred. When such an infection develops, it may require treatment with appropriate local therapy and discontinuation of treatment with Dymista Nasal Spray. Patients using Dymista Nasal Spray over several months or longer should be examined periodically for evidence of *Candida* infection or other signs of adverse effects on the nasal mucosa.

#### 5.3 Glaucoma and Cataracts

Nasal and inhaled corticosteroids may result in the development of glaucoma and/or cataracts. Therefore, close monitoring is warranted in patients with a change in vision or with a history of increased intraocular pressure, glaucoma, and/or cataracts.

Glaucoma and cataract formation were evaluated with intraocular pressure measurements and slit 56 lamp examinations in a controlled 12-month study in 612 adolescent and adult patients aged 12 years and older with perennial allergic or vasomotor rhinitis (VMR). Of the 612 patients enrolled in the study, 405 were randomized to receive Dymista Nasal Spray (1 spray per nostril twice daily) and 207 were randomized to receive fluticasone propionate nasal spray (2 sprays per nostril once daily). In the Dymista Nasal Spray group, one patient had increased intraocular pressure at month 6. In addition, three patients had evidence of posterior subcapsular cataract at month 6 and one at month 12 (end of treatment). In the fluticasone propionate group, three patients had evidence of posterior subcapsular cataract at month 12 (end of treatment).

#### 5.4 Immunosuppression

Persons who are using drugs, such as corticosteroids, that suppress the immune system are more susceptible to infections than healthy individuals. Chickenpox and measles, for example, can have a more serious or even fatal course in susceptible children or adults using corticosteroids. In children or adults who have not had these diseases or been properly immunized, particular care should be taken to avoid exposure. How the dose, route, and duration of corticosteroid administration affect the risk of developing a disseminated infection is not known. The contribution of the underlying disease and/or prior corticosteroid treatment to the risk is also not known. If exposed to chickenpox, prophylaxis with varicella zoster immune globulin (VZIG) may be indicated. If exposed to measles, prophylaxis with pooled intramuscular immunoglobulin 74 (IG) may be indicated. (See the respective package inserts for complete VZIG and IG prescribing information.) If chickenpox develops, treatment with antiviral agents may be considered.

Corticosteroids should be used with caution, if at all, in patients with active or quiescent tuberculous infections of the respiratory tract; untreated local or systemic fungal or bacterial infections; systemic viral or parasitic infections; or ocular herpes simplex because of the potential for worsening of these infections.

#### 5.5 Hypothalamic-Pituitary-Adrenal (HPA) Axis Effects

When intranasal steroids are used at higher than recommended dosages or in susceptible individuals at recommended dosages, systemic corticosteroid effects such as hypercorticism and adrenal suppression may appear. If such changes occur, the dosage of Dymista Nasal Spray should be discontinued slowly, consistent with accepted procedures for discontinuing oral corticosteroid therapy. The concomitant use of intranasal corticosteroids with other inhaled corticosteroids could increase the risk of signs or symptoms of hypercorticism and/or suppression of the HPA axis. The replacement of a systemic corticosteroid with a topical corticosteroid can be accompanied by signs of adrenal insufficiency, and in addition some patients may experience symptoms of withdrawal, e.g., joint and/or muscular pain, lassitude, and depression. Patients previously treated for prolonged periods with systemic corticosteroids and transferred to topical corticosteroids should be carefully monitored for acute adrenal insufficiency in response to stress. In those patients who have asthma or

other clinical conditions requiring long-term systemic corticosteroid treatment, too rapid a decrease in systemic corticosteroids may cause a severe exacerbation of their symptoms.

#### 5.6 Use of Cytochrome P450 3A4 Inhibitors

Ritonavir and other strong cytochrome P450 3A4 (CYP3A4) inhibitors can significantly increase plasma fluticasone propionate exposure, resulting in significantly reduced serum cortisol concentrations [see *Drug Interactions* (7.2) and *Clinical Pharmacology* (12.3)]. During postmarketing use, there have been reports of clinically significant drug interactions in patients receiving fluticasone propionate and ritonavir, resulting in systemic corticosteroid effects including Cushing syndrome and adrenal suppression. Therefore, coadministration of Dymista Nasal Spray and ritonavir is not recommended unless the potential benefit to the patient outweighs the risk of systemic corticosteroid side effects.

Use caution with the coadministration of Dymista Nasal Spray and other potent CYP3A4 inhibitors, such as ketoconazole [see *Drug Interactions* (7.2) and *Clinical Pharmacology* (12.3)].

#### 5.7 Effect on Growth

Corticosteroids may cause a reduction in growth velocity when administered to pediatric patients. Monitor the growth routinely of pediatric patients receiving Dymista Nasal Spray [see *Use in Specific Populations* (8.4)].

### 6 ADVERSE REACTIONS

Systemic and local corticosteroid use may result in the following:

- Somnolence [see *Warnings and Precautions* (5.1)]
- Local nasal effects, including epistaxis, nasal ulceration, nasal septal perforation, impaired wound healing, and *Candida albicans* infection [see *Warnings and Precautions* (5.2)]
- Cataracts and glaucoma [see *Warnings and Precautions* (5.3)]
- Immunosuppression [see *Warnings and Precautions* (5.4)]
- Hypothalamic-pituitary-adrenal (HPA) axis effects, including growth reduction [see *Warnings and Precautions* (5.5 and 5.7), *Use in Specific Populations* (8.4)]

#### 6.1 Clinical Trials Experience

Because clinical trials are conducted under widely varying conditions, adverse reaction rates observed in clinical trials of a drug cannot be directly compared to rates in the clinical trials of another drug and may not reflect rates observed in practice. The safety data described below reflect exposure to Dymista Nasal Spray in 853 patients (12 years of age and older; 36% male and 64% female) with seasonal allergic rhinitis in 3 double-blind, placebo-controlled clinical trials of 2-week duration. The racial distribution for the 3 clinical trials was 80% white, 16% black, 2% Asian, and 1% other. In the 12-month open-label, active-controlled clinical trial, 404 Asian patients (240 males and 164 females) with perennial allergic rhinitis or vasomotor rhinitis were treated with Dymista Nasal Spray, 1 spray per nostril twice daily.

##### Adults and Adolescents 12 Years of Age and Older

In the 3 placebo-controlled clinical trials of 2-week duration, 3411 patients with seasonal allergic rhinitis were treated with 1 spray per nostril of Dymista Nasal Spray, azelastine hydrochloride nasal spray, fluticasone propionate nasal spray, or placebo, twice daily. The azelastine hydrochloride and fluticasone propionate comparators use the same vehicle and device as Dymista Nasal Spray and are not commercially marketed. Overall, adverse reactions were 16% in the Dymista Nasal Spray treatment groups, 15% in the azelastine hydrochloride nasal spray groups, 13% in the fluticasone propionate nasal spray groups, and 12% in the placebo groups. Overall, 1% of patients in both the Dymista Nasal Spray and placebo groups discontinued due to adverse reactions.

Table 1 contains adverse reactions reported with frequencies greater than or equal to 2% and more frequently than placebo in patients treated with Dymista Nasal Spray in the seasonal allergic rhinitis controlled clinical trials.

	1 spray per nostril twice daily			
	Dymista Nasal Spray (N=853)*	Azelastine Hydrochloride Nasal Spray† (N=851)	Fluticasone Propionate Nasal Spray† (N=846)	Vehicle Placebo (N=861)
Dysgeusia	30 (4%)	44 (5%)	4 (1%)	2 (<1%)
Headache	18 (2%)	20 (2%)	20 (2%)	10 (1%)
Epistaxis	16 (2%)	14 (2%)	14 (2%)	15 (2%)

\*Safety population N=853, intent-to-treat population N=848

†Not commercially marketed

In the above trials, somnolence was reported in <1% of patients treated with Dymista Nasal Spray (6 of 853) or vehicle placebo (1 of 861) [see *Warnings and Precautions* (5.1)].

##### Long-Term (12-Month) Safety Trial:

In the 12-month, open-label, active-controlled, long-term safety trial, 404 patients (12 years of age and older) with perennial allergic rhinitis or vasomotor rhinitis were treated with Dymista Nasal Spray 1 spray per nostril twice daily and 207 patients were treated with fluticasone propionate nasal spray, 2 sprays per nostril once daily. Overall, adverse reactions were 47% in the Dymista Nasal Spray treatment group and 44% in the fluticasone propionate nasal spray group. The most frequently reported adverse reactions (≥ 2%) with Dymista Nasal Spray were headache, pyrexia, cough, nasal congestion, rhinitis, dysgeusia, viral infection, upper respiratory tract infection, pharyngitis, pain, diarrhea, and epistaxis. In the Dymista Nasal Spray treatment

group, 7 patients (2%) had mild epistaxis and 1 patient (<1%) had moderate epistaxis. In the fluticasone propionate nasal spray treatment group 1 patient (<1%) had mild epistaxis. No patients had reports of severe epistaxis. Focused nasal examinations were performed and no nasal ulcerations or septal perforations were observed. Eleven of 404 patients (3%) treated with Dymista Nasal Spray and 6 of 207 patients (3%) treated with fluticasone propionate nasal spray discontinued from the trial due to adverse events.

## 7 DRUG INTERACTIONS

No formal drug interaction studies have been performed with Dymista Nasal Spray. The drug interactions of the combination are expected to reflect those of the individual components.

### 7.1 Central Nervous System Depressants

Concurrent use of Dymista Nasal Spray with alcohol or other central nervous system depressants should be avoided because somnolence and impairment of central nervous system performance may occur [see *Warnings and Precautions (5.1)*].

### 7.2 Cytochrome P450 3A4

Ritonavir (a strong CYP3A4 inhibitor) significantly increased plasma fluticasone propionate exposure following administration of fluticasone propionate aqueous nasal spray, resulting in significantly reduced serum cortisol concentrations [see *Clinical Pharmacology (12.3)*]. During postmarketing use, there have been reports of clinically significant drug interactions in patients receiving fluticasone propionate and ritonavir, resulting in systemic corticosteroid effects including Cushing syndrome and adrenal suppression. Therefore, coadministration of fluticasone propionate and ritonavir is not recommended unless the potential benefit to the patient outweighs the risk of systemic corticosteroid side effects.

Ketoconazole (also a strong CYP3A4 inhibitor), administered in multiple 200 mg doses to steady-state, increased plasma exposure of fluticasone propionate, reduced plasma cortisol AUC, but had no effect on urinary excretion of cortisol, following administration of a single 1000 mcg dose of fluticasone propionate by oral inhalation route.

Caution should be exercised when Dymista Nasal Spray is coadministered with ketoconazole and other known strong CYP3A4 inhibitors.

## 8 USE IN SPECIFIC POPULATIONS

### 8.1 Pregnancy

#### **Dymista Nasal Spray: Teratogenic Effects: Pregnancy Category C:**

There are no adequate and well-controlled clinical trials of Dymista Nasal Spray, azelastine hydrochloride only, or fluticasone propionate only in pregnant women. Animal reproductive studies of azelastine hydrochloride and fluticasone propionate in mice, rats, and/or rabbits revealed evidence of teratogenicity as well as other developmental toxic effects. Because animal reproduction studies are not always predictive of human response, Dymista Nasal Spray should be used during pregnancy only if the potential benefit justifies the potential risk to the fetus.

**Azelastine hydrochloride: Teratogenic Effects:** In mice, azelastine hydrochloride caused embryo-fetal death, malformations (cleft palate; short or absent tail; fused, absent or branched ribs), delayed ossification, and decreased fetal weight at an oral dose approximately 610 times the maximum recommended human daily intranasal dose (MRHDID) in adults (on a mg/m<sup>2</sup> basis at a maternal dose of 68.6 mg/kg). This dose also caused maternal toxicity as evidenced by decreased body weight. Neither fetal nor maternal effects occurred at a dose that was approximately 26 times the MRHDID (on a mg/m<sup>2</sup> basis at a maternal dose of 3 mg/kg).

In rats, azelastine hydrochloride caused malformations (oligo- and brachydactylia), delayed ossification and skeletal variations, in the absence of maternal toxicity, at an oral dose approximately 530 times the MRHDID in adults (on a mg/m<sup>2</sup> basis at a maternal dose of 30 mg/kg). At a dose approximately 1200 times the MRHDID (on a mg/m<sup>2</sup> basis at a maternal dose of 68.6 mg/kg), azelastine hydrochloride also caused embryo-fetal death and decreased fetal weight; however, this dose caused severe maternal toxicity. Neither fetal nor maternal effects occurred at a dose approximately 53 times the MRHDID (on a mg/m<sup>2</sup> basis at a maternal dose of 3 mg/kg).

In rabbits, azelastine hydrochloride caused abortion, delayed ossification, and decreased fetal weight at oral doses approximately 1100 times the MRHDID in adults (on a mg/m<sup>2</sup> basis at a maternal dose of 30 mg/kg); however, these doses also resulted in severe maternal toxicity. Neither fetal nor maternal effects occurred at a dose approximately 11 times the MRHDID (on a mg/m<sup>2</sup> basis at a maternal dose of 0.3 mg/kg).

**Fluticasone propionate: Teratogenic Effects:** Corticosteroids have been shown to be teratogenic in laboratory animals when administered systemically at relatively low dosage levels. Subcutaneous studies in the mouse and rat at doses approximately equivalent to and 4 times, respectively, the MRHDID in adults (on a mcg/m<sup>2</sup> basis at maternal doses of 45 and 100 mcg/kg respectively), revealed fetal toxicity characteristic of potent corticosteroid compounds, including embryonic growth retardation, omphalocele, cleft palate, and retarded cranial ossification.

In the rabbit, fetal weight reduction and cleft palate were observed at a subcutaneous dose less than the MRHDID in adults (on a mcg/m<sup>2</sup> basis at a maternal dose of 4 mcg/kg). However, no teratogenic effects were reported at oral doses up to approximately 25 times the MRHDID in adults (on a mcg/m<sup>2</sup> basis at a maternal dose of 300 mcg/kg) of fluticasone propionate to the rabbit. No fluticasone propionate was detected in the plasma in this study, consistent with the established low bioavailability following oral administration [see *Clinical Pharmacology (12.3)*].

Experience with oral corticosteroids since their introduction in pharmacologic, as opposed to physiologic, doses suggests that rodents are more prone to teratogenic effects from corticosteroids than humans. In addition, because there is a natural increase in corticosteroid production during pregnancy, most women will require a lower exogenous corticosteroid dose and many will not need corticosteroid treatment during pregnancy.

**Nonteratogenic Effects:** Fluticasone propionate crossed the placenta following oral administration of approximately 4 and 25 times the MRHDID in adults (on a mcg/m<sup>2</sup> basis at maternal doses of 100 mcg/kg and 300 mcg/kg to rats and rabbits, respectively).

### 8.3 Nursing Mothers

**Dymista Nasal Spray:** It is not known whether Dymista Nasal Spray is excreted in human breast milk. Because many drugs are excreted in human milk, caution should be exercised when Dymista Nasal Spray is administered to a nursing woman. Since there are no data from well-controlled human studies on the use of Dymista Nasal Spray by nursing mothers, based on data from the individual components, a decision should be made whether to discontinue nursing or to discontinue Dymista Nasal Spray, taking into account the importance of Dymista Nasal Spray to the mother.

**Azelastine hydrochloride:** It is not known if azelastine hydrochloride is excreted in human milk.

**Fluticasone propionate:** It is not known if fluticasone propionate is excreted in human milk. However, other corticosteroids are excreted in human milk. Subcutaneous administration to lactating rats of 10 mcg/kg of tritiated fluticasone propionate (less than the maximum recommended daily intranasal dose in adults on a mcg/m<sup>2</sup> basis) resulted in measurable radioactivity in the milk.

### 8.4 Pediatric Use

Safety and effectiveness of Dymista Nasal Spray in pediatric patients below the age of 12 years have not been established.

Controlled clinical studies have shown that intranasal corticosteroids may cause a reduction in growth velocity in pediatric patients. This effect has been observed in the absence of laboratory evidence of HPA axis suppression, suggesting that growth velocity is a more sensitive indicator of systemic corticosteroid exposure in pediatric patients than some commonly used tests of HPA axis function. The long-term effects of this reduction in growth velocity associated with intranasal corticosteroids, including the impact on final adult height, are unknown. The potential for "catch-up" growth following discontinuation of treatment with intranasal corticosteroids has not been adequately studied. The growth of pediatric patients receiving intranasal corticosteroids, including Dymista Nasal Spray, should be monitored routinely (e.g., via stadiometry). The potential growth effects of prolonged treatment should be weighed against the clinical benefits obtained and the risks/benefits of treatment alternatives.

### 8.5 Geriatric Use

Clinical trials of Dymista Nasal Spray did not include sufficient numbers of patients 65 years of age and older to determine whether they respond differently from younger patients. Other reported clinical experience has not identified differences in responses between the elderly and younger patients. In general, dose selection for an elderly patient should be cautious, usually starting at the low end of the dosing range, reflecting the greater frequency of decreased hepatic, renal, or cardiac function, and of concomitant disease or other drug therapy.

## 10 OVERDOSAGE

**Dymista Nasal Spray:** Dymista Nasal Spray contains both azelastine hydrochloride and fluticasone propionate; therefore, the risks associated with overdosage for the individual components described below apply to Dymista Nasal Spray.

**Azelastine hydrochloride:** There have been no reported overdosages with azelastine hydrochloride. Acute azelastine hydrochloride overdosage by adults with this dosage form is unlikely to result in clinically significant adverse events, other than increased somnolence, since one (1) 23 g bottle of Dymista Nasal Spray contains approximately 23 mg of azelastine hydrochloride. Clinical trials in adults with single doses of the oral formulation of azelastine hydrochloride (up to 16 mg) have not resulted in increased incidence of serious adverse events. General supportive measures should be employed if overdosage occurs. There is no known antidote to Dymista Nasal Spray. Oral ingestion of antihistamines has the potential to cause serious adverse effects in children. Accordingly, Dymista Nasal Spray should be kept out of the reach of children.

**Fluticasone propionate:** Chronic fluticasone propionate overdosage may result in signs/symptoms of hypercorticism [see *Warnings and Precautions (5.2)*]. Intranasal administration of 2 mg (10 times the recommended dose) of fluticasone propionate twice daily for 7 days to healthy human volunteers was well tolerated. Single oral fluticasone propionate doses up to 16 mg have been studied in human volunteers with no acute toxic effects reported. Repeat oral doses up to 80 mg daily for 10 days in volunteers and repeat oral doses up to 10 mg daily for 14 days in patients were well tolerated. Adverse reactions were of mild or moderate severity, and incidences were similar in active and placebo treatment groups. Acute overdosage with this dosage form is unlikely since one (1) 23 g bottle of Dymista Nasal Spray contains approximately 8.5 mg of fluticasone propionate.

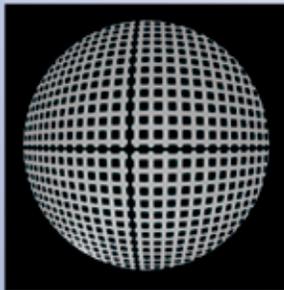
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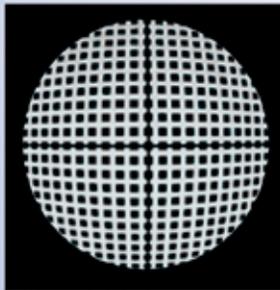
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# bulletin

American Academy of Otolaryngology—Head and Neck Surgery

March 2013—Vol.32 No.03



## World Voice Day 2013: Connect with Your Voice

When you really need to connect with people, there is no substitute for the human voice. Connecting is about bringing people or things together and establishing relationships. Pause for a moment and think about how you personally connect with people.

# 20



AMERICAN ACADEMY OF  
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David R. Nielsen, MD  
Executive Vice President, CEO, and Editor,  
the *Bulletin*

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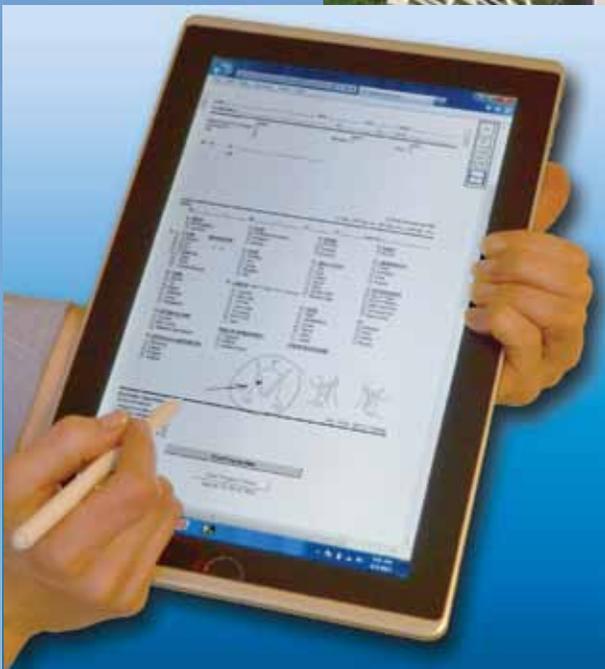
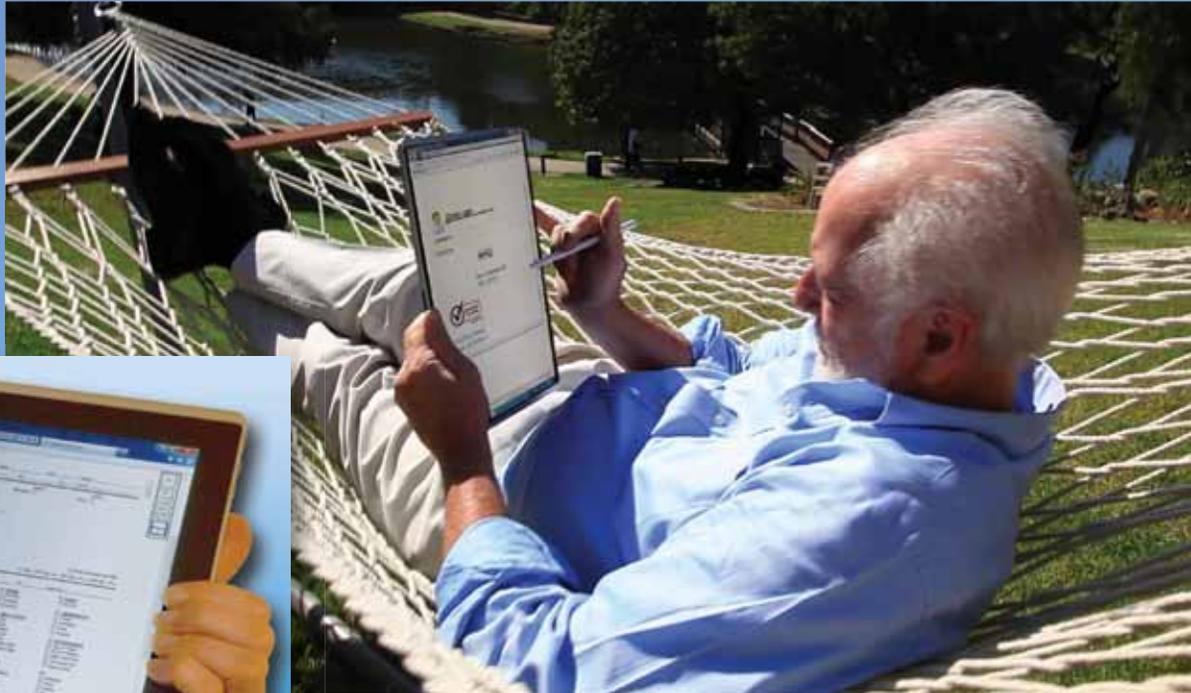
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## World Voice Day: Our Efforts Connect Us to the Future

April 16 is coming. Aside from the fact that it is the day after our taxes are due, otolaryngologists should better know the day as World Voice Day. In a Google search of “World Voice Day,” the Academy’s 2012 campaign material comes up first and second, and our images for the campaign come up third. Following these entries are more than 20 pages of World Voice Day listings from organizations such as the University of Utah, Cleveland Clinic, WBAL News, Johns Hopkins, and the University of Cincinnati with the Cincinnati Opera, as well as international listings from the UK, Brazil, and other international observances. This day is no small deal.

Our Voice Committee, chaired by **Clark A. Rosen, MD**, with the help of **Michael M.E. Johns III, MD**, and **Norman D. Hogikyan, MD**, developed a task force to lead this international observation, and it met during our AAO-HNSF Annual Meeting 2012 & OTO EXPO<sup>SM</sup> last September to plan the 2013 and 2014 campaigns. The task force heard about the highly successful observances in France and Belgium, (from **Marc J. Remacle, MD, PhD**) and the country where the observance originated (from **Mario Andrea, MD, PhD**). While it was noted that it is difficult to engage a renowned celebrity for our domestic campaigns due partially to privacy issues, attendees noted that a campaign featuring the importance of voice to all professions would be the approach to take and to build on. The committee then chose “Connect with Your Voice” as its theme this year.

In following that lead, I came upon a blog by Katie Peters, (<http://katepeters.com/blog/>) a professional speaker. Katie agreed to support World Voice Day 2013 in her blog as she did last year, and she offered this thought on our 2013 theme:

“As humans, we are passionately driven to communicate. We want to be heard. We want to be understood. But to be heard above all the noise of our culture, you must have a voice that others will listen to. There has never been a better time to develop and

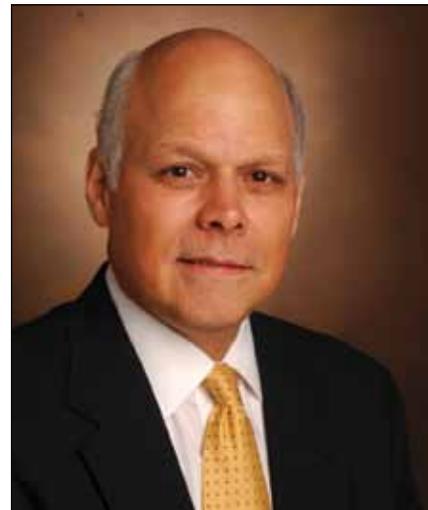
care for that voice. Resources for study and practice are abundant. Expertise is at an all-time high and instead of taking on less importance as our technology advances, the human voice is more important than ever, adding warmth and humanity to a digital world.”

As humans, we are passionately driven to communicate. We want to be heard. We want to be understood. But to be heard above all the noise of our culture, you must have a voice that others will listen to.

- Katie Peters

To help members support this observance and promote the special role otolaryngologists play in the treatment of voice disorders, this *Bulletin* offers you some starting tools. You’ll find a message about the importance of a healthy voice from committee member **Norm D. Hogikyan, MD**, that may be offered to your patients and referral base prior to the observation. Also included in this issue is a fold-out poster to display in your offices that offers an Academy link ([www.entnet.org/HealthInformation/WorldVoiceDay.cfm](http://www.entnet.org/HealthInformation/WorldVoiceDay.cfm)) for additional information about the campaign. Other materials for outreach have been developed by the Voice Committee and the Media and Public Relations Committee (**Wendy B. Stern, MD**, and **Ramon A. Franco Jr., MD**) and include a template letter to send to local media when you login as a Member.

In conclusion, I offer an example of the power of the voice to move human endeavor.



*James L. Netterville, M.D.*

**James L. Netterville, MD**  
AAO-HNS/F President

The following “interplanetary voice-mail” by NASA administrator Charles Bolden was returned to Earth via the Mars rover, Curiosity. The message, which had been sent to Mars and back, was played on Aug. 27, 2012, becoming the first voice transmission from Mars.

“Hello. This is Charlie Bolden, NASA administrator, speaking to you via the broadcast capabilities of the Curiosity rover, which is now on the surface of Mars. Since the beginning of time, humankind’s curiosity has led us to constantly seek new life...new possibilities just beyond the horizon.” (Hear the actual message at [http://www.nasa.gov/mision\\_pages/msl/news/bolden20120827.html](http://www.nasa.gov/mision_pages/msl/news/bolden20120827.html).)

While the message itself is simple, I was struck by the significance of the broadcast as explained by the NASA Curiosity program executive, Dave Lavery, “With this voice, another small step is taken in extending human presence beyond Earth...we hope these words will be an inspiration to someone alive today who will become the first to stand upon the surface of Mars. And like the great Neil Armstrong, they will speak aloud of that next giant leap in human exploration.”

Surely, this is a “connection” to the future powered by the human voice. **5**



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## How Comparatively Effective Are We?

As everyone knows by now, embedded in the Patient Protection and Affordable Care Act of 2010 (ACA) is language designed to address the unsustainable cost of healthcare in the United States by reducing waste, eliminating unnecessary care, and dealing with the unwanted and unexplained variations in care. One specific method the ACA employs is support for comparative effectiveness research (CER)—defined by the Agency for Health Research and Quality (AHRQ) as research methods “designed to inform healthcare decisions by providing evidence on the effectiveness, benefits, and harms of different treatment options. The evidence is generated from research studies that compare drugs, medical devices, tests, surgeries, or ways to deliver healthcare.” [<http://effectivehealthcare.ahrq.gov/index.cfm/what-is-comparative-effectiveness-research1/>]

The Patient-Centered Outcomes Research Institute (PCORI) was founded within the ACA language specifically for the purpose of providing direction and oversight for an entire spectrum of envisioned comparative effectiveness research that could dramatically and positively influence the decision making of professionals, bring consensus around the most efficient ways of providing high quality care for those conditions and interventions for which there is enough data to support a conclusion, and achieve the three aims of the National Quality Strategy: better individual health outcomes, better population health, and reduced cost of healthcare.

While using this approach to improve quality and resource use is a laudable goal and one that every physician and surgeon

can support, the challenge of prioritizing clinical topics, designing relevant and meaningful studies, and acting on what is learned can be complex and daunting. The concept of CER is not new. The medical profession has many years of experience in this approach, but, to date, limited benefit to show from what we have learned. While there are many examples of how such research has improved quality and reduced cost, the promise of CER as envisioned by those who crafted the ACA language remains largely unfulfilled. What are the reasons for this?

A recent article in *Health Affairs* (October 30, 2012) is instructive. After careful study of the literature on many types of CER, the authors conclude that five root causes appear to be responsible for the failure of CER to be translated into positive changes in clinical practice. Misalignment of incentives, ambiguity of results, cognitive biases in interpreting the new information, failure to take into consideration the needs of end users of the data, and limited use of clinical decision support tools all impair the goal of changing clinical behavior. The cognitive biases alone reveal that physicians are not exempt from the powerful effect of traditional behavior and thought processes. As clinicians, the paper discovers, we demonstrate confirmation bias (the effect of believing and acting on that data that supports our pre-conceived notions of what is true); pro-intervention bias (that is, we tend to want to act, rather than to observe or wait, even when the evidence clearly shows that intervention has little or no benefit or may be harmful); and a pro-technology bias



*David R. Nielsen MD*

**David R. Nielsen, MD**  
AAO-HNS/F EVP/CEO

(more recent technological advances are superior to existing modalities).

The article concludes that PCORI has learned that multi-stakeholder involvement in CER from design to implementation is essential to minimize the negative effects of these five barriers and three biases to changing clinical practice for the better. The AAO-HNS/F agrees that collaboration is essential, and has made multi-disciplinary engagement in our Guidelines Task Force a hallmark of our published evidence-based guideline development process. Now in its third edition, if you have not read it, please take the time to review the supplement to the January issue of *Otolaryngology–Head and Neck Surgery*. Since learning to eliminate bias, carefully searching for and critically examining data, and being willing to change our clinical practice to achieve better results are all essential to improving quality, we each need to become familiar with relevant health services research and CER and master the ability to implement what we learn. 

**Source:**

Timbie JW, Fox DS, Van Busum K, Schneider EC. *Health Aff (Millwood)*. Five reasons that many comparative effectiveness studies fail to change patient care and clinical practice. 2012 Oct;31(10):2168-75.

### The particular approach championed by ARHQ and the ACA includes seven distinct steps for optimal implementation:

1. Identify new and emerging clinical interventions.
2. Review and synthesize current medical research.
3. Identify gaps between existing medical research and the needs of clinical practice.
4. Promote and generate new scientific evidence and analytic tools.
5. Train and develop clinical researchers.
6. Translate and disseminate research findings to diverse stakeholders.
7. Reach out to stakeholders via a citizens' forum.

## Using Our Voices to Connect on Grassroot Initiatives

*Denis C. Lafreniere, MD  
Chair, Board of Governors*

To promote World Voice Day, April 16, many of our BOG state and local societies have successfully petitioned their state legislators to formally recognize the day. The voice is vitally important to our patients and ourselves as practitioners, allowing us all the ability to communicate, educate, and entertain.

The evolution of our ability to diagnose and treat voice disorders has certainly accelerated during the last several decades. Technological innovations have continually improved our ability to visualize the larynx and measure physiologic functions involved in voice production. Endoscope images with stroboscopic capabilities can now be seen in high definition, making the diagnosis of even the most subtle mucosal abnormality easier than before. Our ability to perform in-office diagnostic and therapeutic procedures has also improved with these technologic advances, and we routinely perform biopsies, laryngeal EMGs, medialization laryngoplasty injections, and laser treatment of laryngeal lesions under local anesthesia.

The evolution of our ability to diagnose and treat voice disorders has certainly accelerated during the last several decades.

Perhaps the most congenial development has been that the comprehensive care of the voice-disordered patient has led to a significant partnership between the otolaryngologist, the speech pathologist, and vocal pedagogues. Many voice centers can provide expert evaluation of the functional issues involved in many voice disorders

and develop a team approach to the resolution of each patient's voice disorder. This successful collaboration between the otolaryngologist and the allied healthcare provider has required a complete understanding of the roles of each member of the voice care team, which has resulted in the best patient care experience. The ability to work together toward the goal of optimum vocal care ensures that voices will continue to be heard. This metaphor is one that now needs to be applied to the House of Medicine.

### Affordable Care Act

As this column is being written, we have just postponed the "fiscal cliff," but still have no definitive answer to the many fiscal questions that desperately need answers such as the upcoming debt ceiling, underfunded entitlement programs, etc. The Affordable Care Act (ACA) is now in full swing with many states still trying to find their way in this new healthcare world. The meaningful use incentives are now in effect and many, if not most, of our practices have implemented electronic medical records as we work to meet the criteria for incentive payments. During the next several years, we will see these incentives turn into penalties for those not on board. The near future will also mean the introduction of quality metrics that will also result in penalties if these parameters are not measured and met. There are many aspects of the ACA that we as your BOG of the AAO-HNS are working to amend to allow us to maintain our ability to take outstanding care of our patients. The legislative arm of the AAO-HNS has been working with our colleagues from the House of Medicine to repeal the Independent Payment Advisory Board (IPAB) from the ACA as it allows payment decisions for medical expenditures to be influenced by non-elected officials. Many fellows of the AAO-HNS signed a "Declaration of Independence" from the flawed Sustainable Growth Rate Formula this past September in Washington, DC. In this document we collectively raised our voices on this particular issue, and



Denis C. Lafreniere, MD

we plan to continue this fight during this current legislative session. This is a battle that requires as many voices as we can muster. We, the BOG, are asking for your help!

### Spring Meeting and Advocacy Summit

The 2013 BOG Spring Meeting & OTO Advocacy Summit of the AAO-HNS will be taking place May 5-7 in Alexandria, VA. This meeting will immediately follow the Academy's Boards of Directors meeting on May 4. Visits with our individual Members of Congress' offices will take place nearby on Capitol Hill on May 7. I strongly encourage all Academy members, especially our new members and resident members in-training, to invest your time in these meetings. The BOG Spring Meeting will include useful practice information with talks on quality measures, and the changes scheduled to occur with ICD-10. The OTO Advocacy Summit will educate you on current issues being considered on Capitol Hill as we hear from several legislators. We will also discuss talking points for our meetings with our representatives. The stakes for our patients and us as practitioners has never been so high. We need our "voices," both as individuals and as a collective, to be heard loud and clear. 

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## Choosing Wisely®: Our List of Five Things Physicians and Patients Should Question

### A Campaign to Improve the Nation's Healthcare Quality and Safety

On February 21, the American Academy of Otolaryngology—Head and Neck Surgery Foundation (AAO-HNSF) released its list of five things physicians and patients should question as part of the American Board of Internal Medicine (ABIM) Foundation's *Choosing Wisely*® campaign. To date, 25 specialty societies have developed and released lists as part of the initiative.

This month we highlight our five items and provide a set of questions and answers to stimulate discussion of the campaign in your practice and with your patients. Further information about the campaign is available at <http://www.entnet.org/choosingwisely> and <http://www.choosingwisely.org/>.

In releasing the list, the AAO-HNSF would like to thank everyone who provided leadership and input during the list's development. In particular, we would like to highlight the role of the Patient Safety and Quality Improvement Committee that spearheaded the AAO-HNSF list development process.

### What Is the Choosing Wisely Campaign?

The campaign is an initiative of the ABIM Foundation to help physicians and patients engage in conversations about the overuse of tests and procedures and support physician efforts to help patients make smart and effective care choices. Recognizing the importance of physicians and patients working together, leading specialty societies, along with *Consumer Reports*, have joined the campaign to help improve the quality and safety of healthcare in America.

### What Issues Stimulated the Campaign?

As the nation continues to tackle the rising costs of healthcare, it is important for physicians to take a leading role in ensuring patients receive the safest and highest quality of care. According to the ABIM Foundation, "The Congressional Budget Office estimates up to 30 percent of care delivered in the United States goes toward unnecessary tests, procedures, hospital stays, and other services that may not improve people's health—and

Recognizing the importance of physicians and patients working together, leading specialty societies, along with *Consumer Reports*, have joined the campaign to help improve the quality and safety of healthcare in America.

in fact may actually cause harm." The campaign promotes physicians and patients working together and having conversations about wise treatment decisions.

### How Was the List Developed?

The AAO-HNSF's list was developed during a six-month period beginning in May 2012. The Academy's Patient Safety and Quality Improvement Committee spearheaded the list development process.



*An initiative of the ABIM Foundation*

## Five Things Physicians and Patients Should Question

1

### Don't order computed tomography (CT) scan of the head/brain for sudden hearing loss.

Computed tomography scanning is expensive, exposes the patient to radiation and offers no useful information that would improve initial management. CT scanning may be appropriate in patients with focal neurologic findings, a history of trauma or chronic ear disease.

2

### Don't prescribe oral antibiotics for uncomplicated acute tympanostomy tube otorrhea.

Oral antibiotics have significant adverse effects and do not provide adequate coverage of the bacteria that cause most episodes; in contrast, topically administered products do provide coverage for these organisms. Avoidance of oral antibiotics can reduce the spread of antibiotic resistance and the risk of opportunistic infections.

3

### Don't prescribe oral antibiotics for uncomplicated acute external otitis.

Oral antibiotics have significant adverse effects and do not provide adequate coverage of the bacteria that cause most episodes; in contrast, topically administered products do provide coverage for these organisms. Avoidance of oral antibiotics can reduce the spread of antibiotic resistance and the risk of opportunistic infections.

4

### Don't routinely obtain radiographic imaging for patients who meet diagnostic criteria for uncomplicated acute rhinosinusitis.

Imaging of the paranasal sinuses, including plain film radiography, computed tomography (CT) and magnetic resonance imaging (MRI) is unnecessary in patients who meet the clinical diagnostic criteria for uncomplicated acute rhinosinusitis. Acute rhinosinusitis is defined as up to four weeks of purulent nasal drainage (anterior, posterior or both) accompanied by nasal obstruction, facial pain-pressure-fullness or both. Imaging is costly and exposes patients to radiation. Imaging may be appropriate in patients with a complication of acute rhinosinusitis, patients with comorbidities that predispose them to complications and patients in whom an alternative diagnosis is suspected.

5

### Don't obtain computed tomography (CT) or magnetic resonance imaging (MRI) in patients with a primary complaint of hoarseness prior to examining the larynx.

Examination of the larynx with mirror or fiberoptic scope is the primary method for evaluating patients with hoarseness. Imaging is unnecessary in most patients and is both costly and has potential for radiation exposure. After laryngoscopy, evidence supports the use of imaging to further evaluate 1) vocal fold paralysis, or 2) a mass or lesion of the larynx.

Input was sought from Academy and Foundation Committees, the Specialty Society Advisory Council (SSAC), and the Guidelines Task Force (GTF), previously known as the Guidelines Development Task Force. The AAO-HNSF's final list was based on support of the above groups, evidence supporting each of the items (such as clinical practice guidelines), and the current frequency/use of the test or treatment. A more detailed description of the list development process can be found in a commentary in April's edition of *Otolaryngology–Head and Neck Surgery*.

### What Resources Are Available?

The ABIM Foundation has made each participating society's list available publicly. In addition, *Consumer Reports* has begun translating the lists into patient education materials. The AAO-HNSF plans to have patient materials available in the coming months. All AAO-HNSF resources related to the campaign can be found at <http://www.entnet.org/choosingwisely>.

### Will the AAO-HNSF Develop Further lists?

Yes, the AAO-HNSF will continue to participate in the campaign and we hope to develop several more iterations of the list. A third phase of specialty societies have agreed to join the campaign and their lists will be released later this year.

### Which Specialty Societies Have Participated?

Twenty five specialty societies have participated in the campaign and released lists of five items. The first phase included nine societies that released lists in April 2012, they included:

- American Academy of Allergy, Asthma, & Immunology
- American Academy of Family Physicians\*
- American College of Cardiology
- American College of Physicians

- American College of Radiology
- American Gastroenterological Association
- American Society of Clinical Oncology
- American Society of Nephrology
- American Society of Nuclear Cardiology

*\* Released its second list on February 21, 2013.*

The second phase included 16 new societies, with the following societies releasing their lists alongside the AAO-HNSF:

- American Academy of Hospice and Palliative Medicine
- American Academy of Neurology
- American Academy of Ophthalmology
- American Academy of Pediatrics
- American College of Obstetricians and Gynecologists
- American College of Rheumatology
- American Geriatrics Society
- American Society for Clinical Pathology
- American Society of Echocardiography
- American Urological Association
- Society for Vascular Medicine
- Society of Cardiovascular Computed Tomography
- Society of Hospital Medicine
- Society of Nuclear Medicine and Molecular Imaging
- Society of Thoracic Surgeons

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## Continue the *Choosing Wisely*® Conversation: Find AAO-HNSF Updates and Resources 24/7

To help continue to spread the word about *Choosing Wisely*®, visit [www.entnet.org/choosingwisely](http://www.entnet.org/choosingwisely).

**cfm.** Here you can find the announcement and formal list of “Five Things Physicians and Patients Should Question.” In addition to the list, you will find direct access to the sources from which the list was developed

You can also watch a brief video with AAO-HNSF EVP/CEO David R. Nielsen, MD, that explains the scope and purpose of the campaign and the AAO-HNSF commitment to the effort.

Follow the conversation about the campaign through the Academy’s social media sites on Twitter, Facebook, and LinkedIn right from the page. Using the Twitter hashtag #ChoosingWisely will allow you to be part of what the medical community and consumers are saying about the campaign. 



Continue the conversation with resources from [www.entnet.org/choosingwisely.cfm](http://www.entnet.org/choosingwisely.cfm). Additional resources can be found via the ABIM *Choosing Wisely*® campaign website at [www.choosingwisely.org](http://www.choosingwisely.org).

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# Laryngectomy and Laryngeal Cancer: A Fascinating and Inspiring Chapter in the Expansion of Otolaryngology

*Valerie A. Fritsch, MD  
Medical University of South  
Carolina, Charleston*

Once a universally fatal diagnosis, “epithelioid” carcinoma of the larynx has evolved during the past 150 years into one of the most curable cancers. A historical overview of the diagnosis and surgical management of laryngeal carcinoma shows how this disease became the cornerstone of otolaryngology cancer care.

Highlights include important contributions, such as Billroth’s first total laryngectomy in 1873, and Gluck and Cohen’s modified version (1884), which involved completely separating the trachea and pharynx to reduce the risk of post-operative aspiration. Still, at the turn of the 19th century, operative and peri-operative mortality rates were reportedly as high as 50 percent and the procedure was nearly abandoned. Fortunately, rapid biomedical and technologic advances during that time eventually lifted many of the initial limitations. The evolution of laryngoscopy, tracheostomy, neck dissection, and reconstructive surgery, as well as the availability of antibiotics, endotracheal anesthesia, intravenous access, and blood replacement led to a resurgence of radical surgical extirpation in the 1940s.

The role of otolaryngologists in improving diagnostic techniques and surgical approaches and reconstruction was key. Since the latter half of the last century, efforts have focused on refining more advanced techniques to improve voice and swallowing outcomes, while maintaining or improving oncologic outcomes.

“Once a universally fatal diagnosis, “epithelioid” carcinoma of the larynx has evolved during the past 150 years into one of the most curable cancers.”

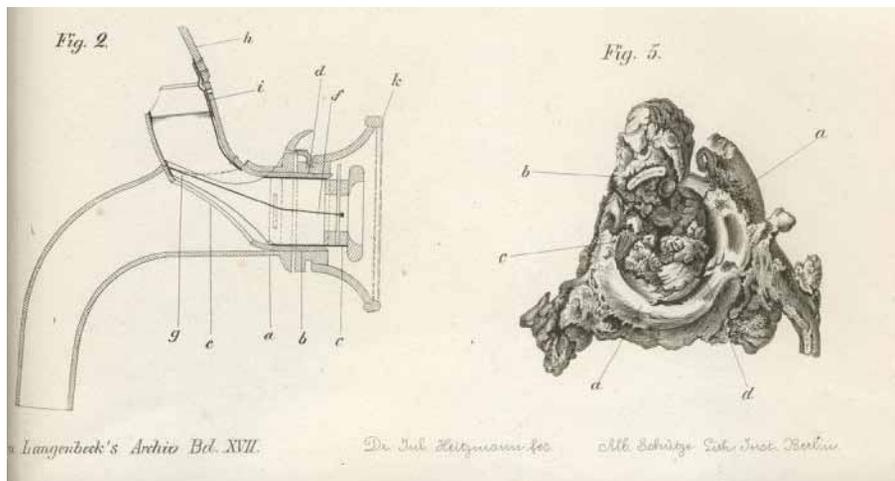
In addition, an increased understanding of the molecular basis of cancer has catalyzed a significant interest in individualized, targeted therapies.

Although the prognosis of laryngeal carcinoma remains far from “favorable” by today’s standards, the astoundingly rapid advances of knowledge and technology by our innovative predecessors illustrates the unbounded potential for future discoveries and improvements in our understanding and management of this complex disease. 



THE INTRODUCTION OF THE POORANG.

See additional photos at [www.entnet.org/bulletin](http://www.entnet.org/bulletin).



## Otolaryngology Historical Society Call for Papers

If you are interested in presenting at the next OHS meeting, which will take place Sept. 30 in Vancouver, BC, Canada, email [museum@entnet.org](mailto:museum@entnet.org).

To join the society or renew your membership, please check the box on your Academy dues invoice or email Catherine R. Lincoln, CAE, MA (Oxon) at [clincoln@entnet.org](mailto:clincoln@entnet.org) or call 1-703-535-3738.

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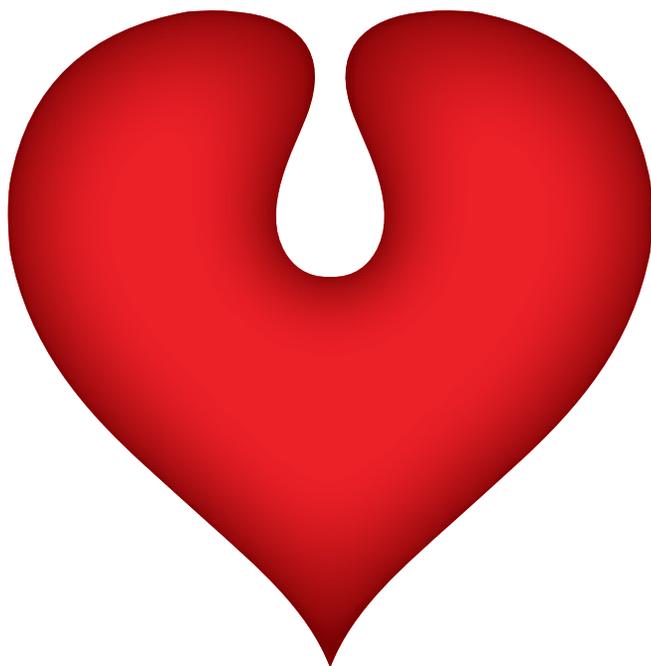
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***Stops symptoms, not patients***

# World Voice Day 2013: Connect with Your Voice



**WORLD VOICE DAY**

**APRIL 16th**

*Norman D. Hogikyon, MD*

**W**hen you really need to connect with people, there is no substitute for the human voice. Connecting is about bringing people or things together and establishing relationships. Pause for a moment and think about how you personally connect with people. Sure, you can send emails, texts, tweets, and photos in order to transfer information or data, but are you really making a connection? How often do these methods lead to misunderstandings or misinformation? Have you ever had the experience of needing to speak with someone in order to clarify what was sent in an email or to soothe angry emotions or hurt feelings from a charged message? For most of us, the answer to this question is a definite yes.

The voice conveys a rainbow of emotion and provides a window into an individual's personality and intentions. It is both the choice of words and how the voice sounds that convey

Clip and copy these two pages to share in your community.







(voice-specialists) available to treat people with complex voice issues.

**Dr. Smith:** Real-time imaging of vocal fold physiology is an area of constant improvement. With better visualization, and therefore improved characterization of what is occurring, the otolaryngologist becomes a better diagnostician.

### What Would You Like the Healthcare Consumer With a Voice Problem to Know?

**Dr. Altman:** I would like to tell those with concerns not to delay proper evaluation, and to be aware of risk factors such as tobacco smoking, reflux, and HPV.

**Dr. Cohen:** Voice problems are common and may be caused by a variety of conditions from benign to neurologic to malignant. They can have a significant impact on patients' ability to communicate, work, function socially, and are treatable. Patients should seek evaluation if symptoms persist more than three weeks.

**Dr. Heman-Ackah:** As industry and technology require a greater use of the voice on a daily basis for working and communicating, more individuals are experiencing difficulties with their voices. Devices as commonplace as a Bluetooth headset present a unique demand on the vocal folds that most people never had to accommodate before, and many individuals are experiencing vocal problems in numbers that did not previously exist. Voice problems today are the 21st century version of carpal tunnel syndrome from the 1980s and 1990s. Many voice problems are the direct result of repetitive

use of the vocal folds and occur from repetitive vocal fold injury.

**Dr. Hogikyan:** I feel that there are two key items here: 1. Hoarseness or voice change can be a sign of a serious problem and should be evaluated by an otolaryngologist if it is persistent (longer than about two weeks can be considered persistent), and 2. Most voice problems can be helped so don't just accept hoarseness without pursuing treatment by a specialist.

**Dr. Akst:**

- Voice quality is an important part of how we present ourselves to others and how we are perceived by others.
- Voice problems are a very common source of work-related difficulties, especially as more jobs depend on verbal communication.
- Voice disorders need not be "accepted as normal"—very often, there are things that can be done to diagnose and treat voice problems.
- Diagnosis and treatment for voice disorders should involve an otolaryngologist with experience in working with voice patients.

**Dr. Young:** Many voice disorders can be improved with proper treatment. Evaluation by an otolaryngologist (or laryngologist) is invaluable. Hoarseness does not need to be simply "tolerated." The most important message is that persistent hoarseness is not normal, and laryngeal examination should be performed to rule out more serious underlying causes, such as cancer.

**Dr. Smith:** There are dedicated otolaryngologists/laryngologists and voice therapists who want to help. We realize that voice is often a reflection of self, especially for the professional

voice user (singer, preacher, teacher, etc). But, the quality of life related to voice is just as important for non-professional voice users, as it allows us to communicate with friends and family, conduct our jobs, and sing.

Each year, aside from working to ensure that quality and appropriate patient care is available to all who need it, the 22 people who make up the Academy Voice Committee champion awareness of the importance of caring for the voice through the World Voice Day campaign on April 16.

### How Does the Observation of World Voice Day Benefit the Public?

**Dr. Altman:** It raises awareness for the working public, but also helps expand the limits of care we provide for voice professionals.

**Dr. Cohen:** By promoting awareness of how vital the voice is to our everyday life, awareness of resources if problems do arise, and discussing prevention in order to keep the voice healthy. The voice is often taken for granted until its function becomes compromised.



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### AcademyU® Connects You to Online Learning Options for Voice Disorders

The Foundation's comprehensive otolaryngology education source, contains hundreds of learning options presented in a variety of formats to complement different learning styles. Developed by leading expert volunteers, the materials are designed to deliver relevant education that is meaningful to your practice.

In recognition of World Voice Day, a list of online lectures addressing issues related to voice disorders is included below:

- Phonomicrosurgery for Benign Pathology  
**Steven M. Zeitels, MD**
- Phonosurgery: What to Do and What to Avoid, Laser and Cold Instruments  
**Jean Abitbol, MD**
- Professional Singers: The Science and Art of Clinical Care  
**Robert Thayer Sataloff, MD**
- A Diagnostic Model for Voice Disorders  
**Robert W. Bastian, MD**

- Vocal Fold Injection: Fact, Fiction, and Material Selection  
**Clark A. Rosen, MD**
- New Lasers: Office and OR, Fiber Cutting and Pulsed Angiolsysis  
**Steven M. Zeitels, MD**
- Vocal Nodules, Polyps, and Cysts: Diagnosis and Current Treatment  
**Clark A. Rosen, MD**
- Presbyphonia  
**Michael M. E. Johns III, MD**  
**Edie Renee Hapner, PhD**

Online Lectures are based on annual meeting instruction courses of the same name. Each lecture provides the highlights of these key sessions.



The Foundation's complete library of online courses and lectures can be found at [www.entnet.org/onlinecourses](http://www.entnet.org/onlinecourses).



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April 10–14, Orlando, FL

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# DRY NASAL ALLERGY SPRAYS

## Return as a treatment option

Into the 1990's dry-formula intranasal steroid (INS) sprays comprised nearly 30% of the nasal allergy market.<sup>1</sup> And then, poof, they were gone. What happened? Many physicians had come to rely on them, including Eli Meltzer, MD, of the Allergy & Asthma Medical Group & Research Center in San Diego, California. "Many clinicians prescribed those corticosteroid nasal aerosol sprays more often than some other medications we had at that time."

### *The Montreal Protocol brought an end to the dry spray*

The problem with the dry sprays was their propellant—chlorofluorocarbon (CFC). CFCs are known to be an ozone-depleting substance (ODS) and harmful to the environment.<sup>2</sup> The "Montreal Protocol on Substances that Deplete the Ozone Layer" is an important international environmental treaty under which the US agreed to phase out the production and importation of ODSs. An exception to this rule was medical products that were determined to be "medically essential."<sup>3</sup> Many asthma and chronic obstructive pulmonary disease (COPD) products fell into this category but nasal allergy sprays did not—and as of January 1, 1996, non-medically essential products could no longer be manufactured.<sup>2,4</sup>



### *Wet sprays attempted to fill the treatment void*

With dry formula sprays no longer an option, doctors sought other solutions for their patients. "You can only use what you have available," said Dr. Meltzer. "On a personal level I preferred the aerosols, but they became less and less available, so we switched to the aqueous corticosteroid sprays, and they were

effective." To this day, aqueous nasal sprays are a valuable treatment option for many patients. However, they are not without their issues.

### *NASAL study revealed patient dissatisfaction*

In 2010, a landmark survey of allergic rhinitis patients and their physicians was conducted to assess how well patients were being managed.<sup>5</sup> The National Allergy Survey Assessing Limitations (NASAL) revealed that many patients were dissatisfied with their current medication. Over 60% of surveyed patients who had used an INS spray in the past year reported that they experienced "medication drip back down the throat." Additionally, just over 18% of patients reported that they experienced "discomfort from spray." Nearly 1 in 5 nasal allergy sufferers asked their doctor to change their INS spray. Of those patients, 28% cited "bothersome side effects" as the cause of their dissatisfaction.

### *Dry sprays make a welcome return*

In time, researchers developed a new, environmentally friendly aerosol propellant.<sup>2</sup> This was welcome news for physicians like Dr. Meltzer: "We were very pleased when HFA (hydrofluoroalkane) asthma inhalers became available and we encouraged the pharmaceutical companies to develop them for nasal allergy treatment. It's nice to say that we now have a couple of dry spray options. I liked them when they were first available, I preferred them when I had access to both the aqueous and the aerosol, and I still prefer them today." Many patients may also agree. "There are patients who prefer one over the other, and it's important to individualize treatment. I consider the dry sprays for patients who have a great amount of nasal drainage or blockage, or for patients who prefer something that doesn't have sensory attributes," said Dr. Meltzer.



"On a personal level I preferred the (dry) aerosols, but they became less and less available..."

ELI  
MELTZER,  
MD

**References:** 1. IMS Health Incorporated. Danbury CT. Monthly Rx data; 1996-2011. 2. US Environmental Protection Agency. Metered dose inhalers: the transition to ozone-safe propellants. <http://www.epa.gov/ozone/title6/downloads/mditransition.pdf>. Updated September 28, 2011. Accessed December 20, 2012. 3. US Environmental Protection Agency. Montreal Protocol: regulatory summary. [http://www.epa.gov/ozone/downloads/MP20\\_Reg\\_Summary.pdf](http://www.epa.gov/ozone/downloads/MP20_Reg_Summary.pdf). Updated September 28, 2011. Accessed December 20, 2012. 4. US Department of Health and Human Services, US Food and Drug Administration. Use of ozone-depleting substances; essential use determinations. In: *Federal Register*. 1999;64(169). Washington, DC: US Department of Health and Human Services. 5. Fromer LM, Ortiz G, Ryan SF, Stoloff SW. Insights on allergic rhinitis from the patient perspective. *J Fam Pract*. 61(2)(suppl 1):S16-S22.

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### Agenda (tentative):

#### BOARD OF GOVERNORS SPRING MEETING

##### Sunday, May 5, 2013

- |  |                     |
|--|---------------------|
| Registration Opens   | 9:00 AM – 4:30 PM   |
| <input type="checkbox"/> Ice-Breaker Event                                 | 10:00 AM – 10:55 AM |
| <input type="checkbox"/> BOG Rules & Regulations Committee Meeting         | 11:00 AM – 12:00 PM |
| <input type="checkbox"/> Luncheon/Media Training                           | 12:05 PM – 1:00 PM  |
| <input type="checkbox"/> BOG Socioeconomic & Grassroots Committee Meeting  | 1:05 PM – 2:55 PM   |
| <input type="checkbox"/> BOG Legislative Representatives Committee Meeting | 3:05 PM – 4:20 PM   |
| <input type="checkbox"/> BOG Executive Committee (by invitation only)      | 4:25 PM – 5:20 PM   |

##### Monday, May 6, 2013

- |   |                    |
|---|--------------------|
| Registration Open   | 7:00 AM – 12:00 PM |
| <input type="checkbox"/> SRF Governing Council (by invitation only) | 7:00 AM – 8:00 AM  |
| <input type="checkbox"/> Society Information Sharing                | 7:30 AM – 8:25 AM  |
| <input type="checkbox"/> Keynote Speaker, Dr. Rahul K. Shah         | 8:30 AM – 9:30 AM  |
| <input type="checkbox"/> BOG General Assembly                       | 9:40 AM – 12:00 PM |
| <input type="checkbox"/> BOG Luncheon Speaker, Wendy L. Kroll, JD   | 12:00 PM – 1:00 PM |
| <input type="checkbox"/> WIO Governing Council (by invitation only) | 5:40 PM – 6:30 PM  |

#### OTO ADVOCACY SUMMIT

##### Sunday, May 5, 2013

- |  |                   |
|--|-------------------|
| <input type="checkbox"/> ENT PAC Reception (Ticket Required) | 6:00 PM – 8:00 PM |
|--|-------------------|

##### Monday, May 6, 2013

- |   |                   |
|---|-------------------|
| <input type="checkbox"/> Advocacy Briefing      | 1:15 PM – 2:30 PM |
| <input type="checkbox"/> Congressional Speakers | 2:45 PM – 5:00 PM |

##### Tuesday, May 7, 2013

- |   |                   |
|---|-------------------|
| <input type="checkbox"/> Hill Visit Q&A                     | 7:00 AM – 8:00 AM |
| <input type="checkbox"/> Pre-Scheduled Congressional Visits | 9:30 AM – 2:30 PM |

##### De-Briefing/Lunch\*

\*AAO-HNS Capitol Hill Office, Washington DC

#### IMPORTANT NOTES:

- \* Attendance is a FREE AAO-HNS Member Benefit!
- \* AAO-HNS staff will schedule your Capitol Hill visits.
- \* Transportation provided to Capitol Hill.
- \* Please plan flight departures NO EARLIER than 4:00 PM on Tuesday, May 7, 2013.

#### Hotel Accommodations:

Hotel Reservations can be made separately at the Embassy Suites Alexandria—Old Town Hotel in the group block named **AAO BOG Spring Meeting & OTO Advocacy Summit**. Our group has rooms available from May 3 - May 7, 2013. Reservations made in this block will receive a special rate of \$189++single/double per night. Housing deadline is April 8, 2013. To make reservations, please visit [www.entnet.org/bog&summit](http://www.entnet.org/bog&summit).

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**Long-Term 52-Week Safety Trial:** In a 52-week placebo-controlled long-term safety trial in patients with PAR, 415 patients (128 males and 287 females, aged 12 to 74 years) were treated with QNASL Nasal Aerosol at a dose of 320 mcg once daily and 111 patients (44 males and 67 females, aged 12 to 67 years) were treated with placebo. Of the 415 patients treated with QNASL Nasal Aerosol, 219 patients were treated for 52 weeks and 196 patients were treated for 30 weeks. While most adverse events were similar in type and rate between the treatment groups, epistaxis occurred more frequently in patients who received QNASL Nasal Aerosol (45 out of 415, 11%) than in patients who received placebo (2 out of 111, 2%). Epistaxis also tended to be more severe in patients treated with QNASL Nasal Aerosol. In 45 reports of epistaxis in patients who received QNASL Nasal Aerosol, 27, 13, and 5 cases were of mild, moderate, and severe intensity, respectively, while the reports of epistaxis in patients who received placebo were of mild (1) and moderate (1) intensity. Seventeen patients treated with QNASL Nasal Aerosol experienced adverse reactions that led to withdrawal from the trial compared to 3 patients treated with placebo. There were 4 nasal erosions and 1 nasal septum ulceration which occurred in patients who received QNASL Nasal Aerosol, and no erosions or ulcerations noted in patients who received placebo. No patient experienced a nasal septum perforation during the trial.

### 6.2 Postmarketing Experience

In addition to adverse reactions reported from clinical trials for QNASL Nasal Aerosol, the following adverse events have been reported during use of other intranasal and inhaled formulations of beclomethasone dipropionate. Because these 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. These events have been chosen for inclusion due to either their seriousness, frequency of reporting, or causal connection to beclomethasone dipropionate or a combination of these factors.

**Intranasal beclomethasone dipropionate:** Nasal septal perforation, glaucoma, cataracts, loss of taste and smell, and hypersensitivity reactions including anaphylaxis, angioedema, rash, and urticaria have been reported following intranasal administration of beclomethasone dipropionate.

**Inhaled beclomethasone dipropionate:** Hypersensitivity reactions, including anaphylaxis, angioedema, rash, urticaria, and bronchospasm have been reported following the oral inhalation of beclomethasone dipropionate.

## 7 DRUG INTERACTIONS

No drug interaction studies have been performed with QNASL Nasal Aerosol.

## 8 USE IN SPECIFIC POPULATIONS

### 8.1 Pregnancy

**Teratogenic Effects:** Pregnancy Category C

There are no adequate and well-controlled clinical trials in pregnant women treated with QNASL Nasal Aerosol. Beclomethasone dipropionate was teratogenic and embryocidal in the mouse and rabbit although these effects were not observed in rats. QNASL Nasal Aerosol should be used during pregnancy only if the potential benefit justifies the potential risk to the fetus. Experience with oral corticosteroids since their introduction in pharmacologic, as opposed to physiologic, doses suggests that rodents are more prone to teratogenic effects from corticosteroids than humans.

Beclomethasone dipropionate administered subcutaneously was teratogenic and embryocidal in the mouse and rabbit at doses approximately twice the maximum recommended human daily intranasal dose (MRHDID) in adults (on a mg/m<sup>2</sup> basis at maternal doses of 0.1 and 0.025 mg/kg/day in mice and rabbits, respectively). No teratogenicity or embryocidal effects were seen in rats at approximately 460 times MRHDID (in adults on a mg/m<sup>2</sup> basis at a maternal inhalation dose of 15 mg/kg/day).

**Non-teratogenic Effects:** Hypoadrenalism may occur in infants born of mothers receiving corticosteroids during pregnancy. Such infants should be carefully monitored.

### 8.3 Nursing Mothers

It is not known whether beclomethasone dipropionate is excreted in human breast milk. However, other corticosteroids have been detected in human breast milk and thus caution should be exercised when QNASL Nasal Aerosol is administered to a nursing mother.

## 8.4 Pediatric Use

The safety and effectiveness for seasonal and perennial allergic rhinitis in children 12 years of age and older have been established. Controlled clinical trials with QNASL Nasal Aerosol included 188 adolescent patients 12 to 17 years of age [see *Clinical Studies* (14)]. The safety and effectiveness of QNASL Nasal Aerosol in children younger than 12 years of age have not been established.

Controlled clinical trials have shown that intranasal corticosteroids may cause a reduction in growth velocity in pediatric patients. This effect has been observed in the absence of laboratory evidence of hypothalamic-pituitary-adrenal (HPA) axis suppression, suggesting that growth velocity is a more sensitive indicator of systemic corticosteroid exposure in pediatric patients than some commonly used tests of HPA-axis function. The long-term effects of reduction in growth velocity associated with intranasal corticosteroids, including the impact on final adult height, are unknown. The potential for "catch-up" growth following discontinuation of treatment with intranasal corticosteroids has not been adequately studied. The growth of pediatric patients receiving intranasal corticosteroids, including QNASL Nasal Aerosol, should be monitored routinely (e.g., via stadiometry).

A 12-month, randomized, controlled clinical trial evaluated the effects of QVAR<sup>®</sup>, an orally inhaled HFA beclomethasone dipropionate product, without spacer versus chlorofluorocarbon-propelled (CFC) beclomethasone dipropionate with large volume spacer on growth in children with asthma ages 5 to 11 years. A total of 520 patients were enrolled, of whom 394 received HFA-beclomethasone dipropionate (100 to 400 mcg/day ex-valve) and 126 received CFC-beclomethasone dipropionate (200 to 800 mcg/day ex-valve). When comparing results at month 12 to baseline, the mean growth velocity in children treated with HFA-beclomethasone dipropionate was approximately 0.5 cm/year less than that noted with children treated with CFC-beclomethasone dipropionate via large volume spacer. The potential growth effects of prolonged treatment should be weighed against the clinical benefits obtained and the risks/benefits of treatment alternatives.

The potential for QNASL Nasal Aerosol to cause reduction in growth velocity in susceptible patients or when given at higher than recommended dosages cannot be ruled out.

## 8.5 Geriatric Use

Clinical trials of QNASL Nasal Aerosol did not include sufficient numbers of subjects aged 65 years and older to determine whether they responded differently than younger subjects. Other reported clinical experience has not identified differences in responses between the elderly and younger patients. In general, administration to elderly patients should be cautious, reflecting the greater frequency of decreased hepatic, renal, or cardiac function, and of concomitant disease or other drug therapy.

## 10 OVERDOSAGE

Chronic overdosage may result in signs/symptoms of hypercorticism [see *Warnings and Precautions* (5.5)]. There are no data available on the effects of acute or chronic overdosage with QNASL Nasal Aerosol.



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# What Are RUC Surveys and Why Should They Matter to Me?

**A**s an Academy member, you've probably seen frequent requests distributed in "The News" asking for volunteers for upcoming AMA Relative Value Scale Update Committee (RUC) surveys of physician services. Many of you may have asked yourself, "what the RUC is and why are these surveys important?" During the last several years, the Academy has provided members with background on the RUC in an effort to educate and engage members in the annual RUC process. This year, we'd like to address the common questions that arise during the RUC survey process in hopes of outlining why member participation in these surveys is so critical.

## What is the RUC and Who Participates?

The AMA RUC was developed in response to the transition to a physician payment system based on a Resource-Based Relative Value Scale (RBRVS). The RUC is a multispecialty committee that provides clinical expertise and input on the resources required to provide physician services. The RUC submits recommendations annually to the Centers for Medicare and Medicaid Services (CMS), which uses them to develop relative values for physician services provided to Medicare beneficiaries. The RUC, in conjunction with the Current Procedural Terminology (CPT®) Editorial Panel, has created a process where specialty societies can develop relative value recommendations for new and revised codes, and the RUC carefully reviews survey data presented by specialty societies to develop recommendations for consideration by CMS. CMS then issues final payment policies and values in the final Medicare Physician Fee Schedule rule, which is typically released around the first of November each year.

The RUC is intended to represent the entire medical profession and includes the following medical specialties: anesthesiology, cardiology, dermatology, emergency medicine, family medicine, general surgery, geriatrics, internal medicine, neurology, neurosurgery, obstetrics/gynecology, ophthalmology, orthopedic surgery, otolaryngology, pathology, pediatrics, plastic surgery, primary care (rotating seat), pulmonary medicine (rotating seat), psychiatry, radiology, rheumatology (rotating seat), thoracic surgery, urology, and vascular surgery (rotating seat). Four seats rotate on a two-year basis, with two reserved for an internal medicine subspecialty: one for a primary care representative, and one for any other specialty. The RUC chair, the co-chair of the RUC Health Care Professionals Advisory Committee Review Board, and representatives of the AMA, American Osteopathic Association, the chair of the Practice Expense Review Committee and CPT Editorial Panel hold the remaining six seats. The AMA Board of Trustees selects the RUC chair and the AMA representative to the RUC. The individual RUC members are nominated by the specialty societies and are approved by the AMA.

## Who Represents the Academy at the RUC?

The Academy actively participates in the RUC process and surveys codes for nearly every RUC meeting. Meetings take place every winter, spring, and fall. The Academy's current RUC representatives are RUC panel member **Charles F. Koopmann Jr., MD, MSHA**, and panel member alternate, **Jane T. Dillon, MD**, as well as our RUC advisors **Wayne M. Koch, MD**, and advisor alternate **John T. Lanza, MD**. It is important to recognize that the RUC panel member representatives for each specialty are not advocates for their specialties, rather,

they participate in an individual capacity and represent their own views and independent judgment while serving on the panel. In contrast, AAO-HNS' RUC advisors are responsible for working with the Physician Payment Policy Workgroup (3P) and Academy staff to develop relative value recommendations and practice expense direct inputs for otolaryngology services that are presented to the RUC on behalf of the Academy.

## Why are RUC Surveys Conducted?

Surveys are used by the AMA RUC to allow medical specialty societies to have an active role in ensuring that relative values assigned to medical procedures and services are accurately and fairly presented to CMS. These surveys are critical because the values derived by member survey responses are used by our RUC advisors to make valuation recommendations to the AMA RUC. The goal of the surveys is to obtain time and complexity estimates required when performing a specific medical procedure. This information is then used to estimate a recommended physician work value.

## How Does the Survey Generate a Recommended Value?

The surveys will ask physician members to compare the time, complexity, and work required to perform the procedure being surveyed as compared to another existing medical procedure. A list of possible comparator, or reference, procedures is provided to survey respondents as part of the survey.

## What are the Key Components of the RUC Survey?

First, it is critical that members carefully review the code descriptor and vignette. This is critical because code

descriptors may have been modified and survey respondents will be asked if the descriptor and vignette match their typical (i.e., more than 50 percent of the time) patient. If the descriptor and vignette do not match the respondent's typical patient, the respondent will be asked to write a brief rationale for how their typical patient differs from the survey descriptor or vignette.

Next, surveyees will be asked to review and provide their basic contact information. They will then be asked to identify a reference procedure from the list of potential reference codes. Respondents should select the code from the list that is most similar in physician time and work to the new/revised CPT code descriptor and typical patient. The reference service does not have to be clinically similar to the procedure being surveyed, but must be similar in work required to perform

Surveys are used by the AMA RUC to allow medical specialty societies to have an active role in ensuring that relative values assigned to medical procedures and services are accurately and fairly presented to CMS.

the procedure. It is also important that respondents consider the global period of the service being reviewed. For CPT codes with 000, 010, or 090 day globals, physician services or visits provided within 24 hours prior are included and should be considered by respondents in their recommended value for the service. Likewise, for 010 and 090 globals, the post care following the procedure should be included in the estimate of physician work for a given procedure.

Another key component to the RUC survey is estimating physician time. Respondents should base their

recommendations of the time it takes them to perform the procedure under review on their own personal experience. It is important to note that time estimates provided should be based on the typical patient and not the most straightforward or most complex case the physician respondent has encountered.

There are three components to time estimates. First, the pre-service time, which begins the day prior to the procedure and lasts until the time of the operative procedure. Pre-service time is divided into three activities: evaluation; positioning; and scrub, dress, and wait time. Second, the intra-service time, which includes all "skin to skin" work that is a necessary part of the procedure. And last, the post-service time, which includes the physician services provided on the day of the procedure after the procedure has been performed.

One common source of confusion is the component of moderate sedation. Moderate sedation is a service provided by the operating physician or under the direct supervision of the physician performing the procedure. If anesthesia is provided separately by an anesthesiologist who is not performing the primary procedure, this work should not be included in the valuation of the procedure for the purposes of the RUC survey.

Finally, survey respondents will be asked to evaluate physician work and assign a recommended relative value unit for the work required to perform the procedure. Physician work includes the time it takes the physician to perform the procedure. Physician work should also include the mental effort and judgment necessary, as well as the technical skill required to perform the procedure. Note, time and work valuation should not include any work or service provided by clinical staff that are employed by the physician's practice and cannot bill separately. It is important to keep in mind that the survey methodology aims to set the work RVU for the procedure under review "relative" to the comparable reference procedure selected at the outset of the survey, and respondents may want to

print out the reference service list to refresh them on the value of the comparator code selected.

### What About the Practice Expense Portion of My Payment?

As part of its role in the RUC process, the Academy RUC team is asked to provide the AMA RUC and CMS with information regarding the direct practice expense inputs for all procedures that undergo RUC review. This includes recommendations on clinical staff time needed during the procedure, as well as equipment and supplies required for the procedure. These recommendations are reviewed by the Practice Expense Advisory Committee (PEAC) of the RUC and approved or modified prior to being submitted to CMS for acceptance in the final CY MPFS.

### What About the Malpractice Portion of My Payment?

The AMA RUC sends recommendations to CMS on practice liability crosswalks for each procedure reviewed by the AMA RUC. This occurs in May of each year and, similar to the practice expense and physician work recommendations submitted by the AMA RUC, are approved or modified by CMS in the MPFS for that calendar year. All values finalized in the final rule then take effect the following January.

### Still Have Questions?

For more background on the RUC survey process, members can access the following PowerPoint presentation on the Academy website: <http://www.entnet.org/Practice/upload/2012-ruc-survey-presentation.pdf>. Members can also email any questions to Jenna Minton at [Jminton@entnet.org](mailto:Jminton@entnet.org). We hope this information will assist members in better understanding the composition of the RUC surveys as well as the importance of your participation in future surveys and the valuation of otolaryngology-head and neck surgery procedures. 

## Physician Compare: What Is CMS Posting about Me?

You've probably heard about CMS' Physician Compare Website. You may even know that the development of this public website is a statutory mandate from the Affordable Care Act. Unfortunately, what many physicians and members are not aware of is the information that's available about them, and their practices, on this website.

The first thing members should know is that this program is updated and modified on an annual basis through federal rulemaking. This means that CMS will announce proposed revisions or additions to the Physician Compare Website in the notice of proposed rulemaking (NPRM) of the Medicare Physician Fee Schedule (MPFS) each calendar year. They then accept feedback on their proposals from the public during a 60-day comment period and finalize their policies in the final MPFS for that year, typically published on or around November 1 each year. The

Website can be accessed at: [http://www.medicare.gov/find-a-doctor/\(X\(1\)S\(0grwmc55y5poo245uxj5ifv5\)\)/provider-search.aspx?AspxAutoDetectCookieSupport=1](http://www.medicare.gov/find-a-doctor/(X(1)S(0grwmc55y5poo245uxj5ifv5))/provider-search.aspx?AspxAutoDetectCookieSupport=1).

### Background

The Physician Compare Website was launched in December 2010 and originally included data on those eligible professionals (EPs) who successfully reported on the Physician Quality Reporting System (PQRS) measures in CY 2009.

Today, the Website includes the following information on providers:

- The provider's primary, and any applicable secondary, specialty(ies);
- The provider's practice locations;
- The providers group practice or hospital affiliations, where applicable;
- The provider's education information, language skills, and gender;
- The names EPs who have successfully reported on quality programs, specifically e-prescribing and PQRS for CY 2011.

The first thing members should know is that this program is updated and modified on an annual basis through federal rulemaking. This means that CMS will announce proposed revisions or additions to the Physician Compare Website in the notice of proposed rulemaking (NPRM) of the Medicare Physician Fee Schedule (MPFS) each calendar year.

### What to Expect in 2013

By January 2013, CMS is required to outline a plan for posting information on provider's quality performance, as well as patient experience data, on the Physician Compare Website. CMS is presently undertaking a full Website redesign project aimed at improving the usage and function of the site. In addition, CMS has finalized the following information for release in CY 2013:

- 2012 data on PQRS Group Practice Reporting Option (GPRO) measures for practices that meet the minimum sample size of 20 patients;
- Whether providers accept Medicare patients;
- Board certification information; and
- Improved information on language skills and hospital affiliations.

As part of the Website redesign project, CMS allowed the Academy



to view and comment on the proposed redesign. In the letter, the Academy addressed several issues included in the proposed redesign. Visit the Academy's "What's New" page at: <http://www.entnet.org/Practice/CMS-News.cfm> to access the full letter. Areas addressed by the Academy included:

- Data accuracy is paramount in the physician compare site and CMS proposes including claims based verification for physician information rather than just relying on PECOS;
- Concern regarding the small sample size used for posting information such as participation in PQRS GPRO;
- The posting of GPRO performance rates and the need for a review period to ensure the data posted is accurate; and
- The inclusion of CG-CAHPS survey data, as well as the S-CAHPS data, into the Physician Compare website.

### What to Expect in 2014 and Beyond

For 2014, CMS anticipates posting information on provider performance rates on measures reported by physician groups or Accountable Care Organizations (ACOs) via the GRPO web interface system. These groups will have 30 days to review their information for accuracy before it becomes publicly available. CMS also hopes to post patient experience survey data gathers using the CG-CAHPS survey method for groups of 100 or more providers. Finally, CMS will post information on providers who obtain PQRS maintenance of certification incentives during CY 2014.

Other information CMS is considering for inclusion on the website in the future are performance on quality measures developed by specialty societies, continued efforts to align the PQRS and value based payment modifier (VBP) program measures, the release of provider performance in these programs, as well

as individual EP performance measure data.

### Data Accuracy

Currently, the website pulls physician information from the PECOS enrollment system. CMS plans to verify the accuracy of PECOS information via a claims based verification, which the Academy supports. Until these changes are implemented, the Academy encourages members to check their PECOS enrollment information, as well as what information is currently available about them on the Physician Compare website, and to contact CMS if they find the information is incorrect.

The Academy's health policy team will continue tracking the development of this important public website and will alert members to any key changes to the program in the future. Should you have any questions or concerns about the website, or your publicly available information, please contact us at [healthpolicy@entnet.org](mailto:healthpolicy@entnet.org). 

# Bulletin Content AT YOUR FINGERTIPS



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**Contact us** any time Toll-free 1-877-722-6467 (U.S. and Canada); 1-703-836-4444 (international); or [memberservices@entnet.org](mailto:memberservices@entnet.org).



With membership comes many rewarding ways to engage with your colleagues through the Academy and its Foundation. Members can select opportunities based on schedules, interests, and priorities. ▶

## Involvement Levels: You Choose!



As an otolaryngologist—head and neck surgeon, you want to provide the best care to your patients. To help you succeed, the AAO-HNS/F has created opportunities for you to continue improving as a physician and as a leader.

We know your time is limited, so we have made it easy for you to Maximize Your Membership by getting involved in Academy and Foundation activities.

We encourage you to visit [www.entnet.org/getinvolved](http://www.entnet.org/getinvolved) to see which activities spark your interest and fit your schedule.

These activities are designed to fit any level of participation, from face-to-face networking opportunities to activities that do not require you to go any farther than your computer.

These opportunities are not only member benefits, but they also provide a valuable service to the specialty. Participation in these volunteer activities is based on your own personal schedule and interests.



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## AcademyU® Online Education Offers Hundreds of Learning Opportunities

AcademyU®, the Foundation's otolaryngology education source, offers five types of learning formats that include knowledge resources, subscriptions, live events, eBooks, and online education. Each one contains elements that make up the breadth of the education opportunities available through the Foundation. In this second article in the series, we explore the variety of activities that make up the online education component of AcademyU®; these include online courses and lectures and COOL cases.

AcademyU® Online Education ([www.entnet.org/onlinecourse](http://www.entnet.org/onlinecourse)) is organized by the eight subspecialties within otolaryngology-head and neck surgery to make it easy for any otolaryngology specialist to find the courses that best fit his or her education needs. In addition, the online platform makes it easy for learners to take advantage of these education opportunities on their own schedules and at their own pace.



### Online Courses and Lectures

Online Courses are learning activities developed by the Foundation education committees. These peer-reviewed courses provide in-depth

study of otolaryngology head and neck surgery topics determined by an expert-driven analysis of learner education needs. These high-quality courses offer 45 to 60 minutes of detailed instruction on a particular topic. Each contains rich media elements such as detailed images and short video clips.

### The online courses are:

- Preventing Operating Room Fires
- Optimal Safety in Otolaryngic Allergy Practice
- Chin Augmentation: Sliding Osteotomy and Alloplastic Implants

- Nasal Trauma
- Graves' Disease
- Alternative Medicine: Perioperative Management Issues of Herbal Supplements and Vitamins
- Basic Head and Neck Pathology
- Laser Safety
- Evaluation of an Adult Patient with a Benign Neck Mass
- Evaluation of an Adult Patient with a Malignant Neck Mass
- Loco-regional Recurrence in Head and Neck Squamous Cell Carcinoma
- Introducing the AAO-HNS Expert Witness Guidelines
- English-to-Spanish Ear Examination Phrases
- Gender Equity in the Workplace
- Hearing Assessment
- Understanding Stereotactic Radiation for Skull Base Tumors
- The Ten Minute Exam of the Dizzy Patient
- Office Otoscopy I: Normal Examination, Spectrum of Otitis Media, and Characteristic Appearances of Abnormal Pathologies
- Office Otoscopy II: Case Studies
- Office Otoscopy III: Clinical Case Studies Featuring Long-term Serial Examination and Anatomic Cross Section
- Risks of Steroids for Sudden Sensorineural Hearing Loss
- Cleft Lip and Palate Overview
- Introduction to Velopharyngeal Dysfunction
- Management of Sinonasal Cerebrospinal Fluid Leaks

Online Lectures are based on the Annual Meeting & OTO EXPO<sup>SM</sup> instruction courses of the same name. They are selected from the top abstracts submitted to the Annual Meeting; faculty are invited to record a condensed version of their presentation for publication to the AcademyU® website. Each lecture provides highlights of key sessions in short 20- to 40-minute segments using the speakers' slides and audio

recordings. There are online lectures available, including more than 100 from the 2012 Annual Meeting & OTO EXPO.

### The 2013 Online Lectures are:

- Worldwide Otolaryngology Humanitarian Missions
- Developing a Quality Control Program for Surgeons
- Rhinoplasty: Arming Novices for Success
- Facial Aesthetic Enhancements: Chemodenervation and Tissue Augmentation
- Current Management of Oropharyngeal Cancer
- The Management of Glottic Cancer in 2012
- Endoscopic and Robotic Thyroid Surgery
- Minimally Invasive Salivary Endoscopy
- Chronic Cough: Hacking Up a Treatment Algorithm
- Endoscopic Microsurgical Techniques for Laryngeal Disease
- Laryngopharyngeal Reflux (second edition)
- Tympanoplasty/Ossicular Reconstruction—Some Novel Ideas?
- Balance Problems in the Elderly
- Tinnitus: New Frontiers in Radiology and Brain Imaging
- Meniere's or Migraine: Similarities, Differences, Treatments
- Surgical Management of Eustachian Tube Disorders
- Pediatric Obstructive Sleep Apnea What to do after T and A?
- Chronic Rhinosinusitis in Children (second edition)
- Stertor, Stridor, and Babies that Squeak: A Practical Approach
- Up-to-Date Management of Recalcitrant Sinonasal Polyposis
- Five New Landmarks to Make You a Better Sinus Surgeon

Target audiences for both the online courses and online lectures are practicing otolaryngology-head and neck



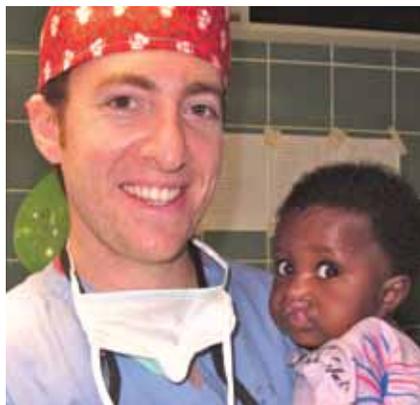


## 2013 Humanitarian Travel Grants: Congratulations to the 15 Residents, Fellows-in-Training Awarded

Thanks to the generous support of Academy members who donated to our humanitarian efforts projects, 15 residents and fellows-in-training received grants of \$1,000 each toward medical missions from January through July 2013.

For more than a decade, our AAO-HNS Foundation's Humanitarian Efforts Committee has selected senior residents and fellows-in-training for travel grants to accompany mission teams. While the grants of \$1,000 each cannot cover the travel costs, they are an inspiration to the grantees, who return profoundly changed by their experiences. Feedback from returning residents has demonstrated how invaluable these encounters are for both their personal and professional development. Overwhelmingly, the awardees commit themselves to continuing to volunteer for missions throughout their lives.

The awardees will be recognized during the AAO-HNSF 2013 Annual Meeting and OTO EXPO<sup>SM</sup>, Vancouver Convention Centre in Vancouver, BC, during the Humanitarian Forum. Please join us in congratulating these dedicated residents and fellows-in-training.



1. **Sarah N. Bowe, MD**, Ohio State University Medical Center, Project EAR, Inc., Dominican Republic, Los Alcarrizos, April 13-21, 2013.
2. **Do-Yeon Cho, MD**, Stanford University, Myungsung Christian Foundation, Ethiopia, Addis Ababa, May 20-31, 2013.
3. **David J. Crockett, MD**, University of Utah, division of otolaryngology, Operation Restore Hope, Philippines, Cebu, February 15-24, 2013.
4. **Ethan B. Handler, MD**, Kaiser Permanente Oakland, Faces of Tomorrow, Ecuador, Quito, June 8-17, 2013.

5. **Andrew C. Heaford, MD**, University of Iowa Hospitals and Clinics; department of otolaryngology: head and neck surgery, Miles of Smiles in Guatemala; Iowa MOST mission, Guatemala, Huehuetenango, February 14-24, 2013.
6. **Evan R. McBeath, MD**, University Hospitals Case Medical Center, Case Western Reserve University, Concern for Children, El Salvador, San Salvador, January 9-19, 2013.
7. **Bryan R. McRae, MD**, Indiana University School of Medicine, department of otolaryngology-head & neck

### Join KJ Lee, MD, for the 2013 China Tour

KJ Lee, MD, invites you to experience China, June 5-16, after the IFOS World Congress, Seoul, South Korea, and ending at the World Chinese ENT Academy Congress, Hong Kong.

Exchange ideas with Chinese otolaryngology leaders and enjoy Chinese cultural heritage, with such famous sights as:

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- Xi'an's terra cotta warriors and the World Heritage Site, Fujian Tulou
- Hong Kong

To reserve, call 1-203-772-0060, 1-800-243-1806 or email [donna.dalnekoff@atpi.com](mailto:donna.dalnekoff@atpi.com). Questions? Contact Dr. Lee, Academy past president, by calling 1-203-777-4005 or emailing [kjleemd@aol.com](mailto:kjleemd@aol.com).





- surgery, IU-Kenya Program, Kenya, Eldoret, January 19-February 3, 2013.
8. **Sachin S. Pawar, MD**, Oregon Health & Science University, FACES Foundation, Peru, Lambayeque, January 25-February 3, 2013.
  9. **Angela S. Peng, MD**, University of Minnesota, department of otolaryngology-head & neck surgery, Mayflower Medical Outreach, Nicaragua, Managua, February 16-24, 2013.
  10. **Maria de Lourdes Quintanilla-Dieck, MD**, Oregon Health & Science

- University, FACES Foundation, Peru, Lambayeque, January 25-February 3, 2013.
11. **Joseph W. Rohrer, MD**, San Antonio Uniformed Services Health Education Consortium, Face the Future Mission Rwanda, Rwanda, Kigali, February 1-10, 2013.
  12. **Dhava Setabutr, MD**, Penn State Hershey Medical Center, Faces of Tomorrow, Ecuador, Quito, June 7-16, 2013.

13. **Laura L. Shively, MD**, Dartmouth-Hitchcock Medical Center, Mayflower Medical Outreach, Nicaragua, Jinotega, Managua, February 17-25, 2013.
14. **Yi-Hsuan E. Wu, MD**, Tufts Medical Center, Medical Missions for Children, Rwanda, Gitwe, March 7-17, 2013.
15. **Estelle S. Yoo, MD**, Alfred I. DuPont Hospital for Children, department of surgery/division of otolaryngology, World Hearing Foundation, Honduras, Tegucigalpa, March 23-29, 2013.

Visit the Humanitarian Efforts Member Engagement Portal to help facilitate matching critical needs with medical specialty expertise: [www.entnet.org/humanitarianportal](http://www.entnet.org/humanitarianportal).

To learn more about Humanitarian Resident Travel Grants visit <http://www.entnet.org/HumanitarianTravel>. May 31, 2013, is the deadline for grant applications for mission trips during July 1 through December 31, 2013.

| [www.karenzupko.com/workshops/otolaryngology](http://www.karenzupko.com/workshops/otolaryngology)

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<p><b>Sept 13-14</b> Minneapolis, MN Radisson Blu Mall of America</p>	<p><b>Oct 25-26</b> Las Vegas, NV The Westin Las Vegas</p>	<p><b>Nov 8-9</b> Chicago, IL The Hyatt Chicago Mag Mile</p>

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**Send letter of inquiry & curriculum vitae to:**

Gerald S. Berke, M.D., Professor and Chair  
UCLA Department of Head and Neck Surgery  
10833 Le Conte Avenue, CHS 62-132  
Los Angeles, CA 90095-1624



**The Department of Otolaryngology at West Virginia University** is seeking a fellowship-trained head and neck surgeon to join a well established head and neck oncology service in the summer of 2013. Expertise with both ablative and reconstructive procedures is desired. Responsibilities include education of residents and medical students and patient care. Opportunities are available for those interested in clinical/basic research.

The department currently has ten physician faculty members and fifteen residents and has an active NIH-funded research division with three PhD members.

West Virginia University is located in beautiful Morgantown, which is rated one of the best small towns in America in regard to quality of life. Located 80 miles south of Pittsburgh and three hours from Washington, DC, Morgantown has an excellent public school system and offers culturally diverse, large-city amenities in a safe, family setting.

**The position will remain opened until filled. Please send a CV with three professional references to:**

Laura Blake  
Director, Physician Recruitment  
Fax: 304-293-0230  
blakel@wvuhealthcare.com  
<http://www.hsc.wvu.edu/som/otolaryngology/>

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 Current Florida license  
 Bilingual (English/Spanish) preferred  
 Excellent communication and interpersonal skills.  
 ENT Experience a must  
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**Contact Information**

Contact name: Stacey Citrin, CEO  
 Phone: (305)558-3724  
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 Cellular: (954)816-1087

**Dade Location:**

Horacio Groisman, MD  
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West Virginia University

**The Department of Otolaryngology/Head & Neck Surgery at West Virginia University** is seeking a general otolaryngologist to join a thriving academic practice in the summer of 2013. Applicants must be board certified/eligible by the American Board of Otolaryngology. Responsibilities include teaching of residents and medical students, patient care and clinical/basic research.

The department currently has ten physician faculty members and fifteen residents and has an active NIH-funded research division with three PhD members.

With a metro area population of over 115,000, Morgantown, WV, is consistently rated as one of the best small cities in the U.S., with affordable housing, excellent schools, a picturesque countryside, many outdoor recreational activities, and close proximity to major cities, such as Pittsburgh, PA, and Washington, DC.

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The University of Kansas Department of Otolaryngology-Head & Neck Surgery is seeking a laryngologist, a head and neck surgeon, and a rhinologist/skull base surgeon who are interested in full-time academic positions.

The successful candidate will have fellowship training with expertise in their specialty and is BC/BE. The candidate will join as an Assistant or Associate Professor and will be involved with resident and medical student education while developing a strong clinical practice and research interests.

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Join a busy voice and swallow team with a state-of-the-art laryngeal lab and experienced speech pathology support.

#### Head and Neck Surgeon

Position Number M0203642

Join a division of four head and neck surgeons. Fellowship in microvascular surgery, surgical oncology and an interest in oncologic research preferred.

#### Veterans Affairs Clinician/Scientist

The Department is looking for a full-time VA position with potential for VA research funding. Ideally this position will allow 50% protected time for research.

#### Head and Neck Fellowship

Clinical Focus: Head and Neck Surgical Oncology, Skull Base Surgery, Endoscopic Laser Surgery, Minimally Invasive Endocrine Surgery, Microvascular Reconstructive Surgery and Robotic Surgery.

Applications are accepted through the American Head and Neck Society: [www.ahns.info](http://www.ahns.info).



To view position online, go to <http://jobs.kumc.edu>  
(Search by position number.)

**Letters of inquiry and CV may be mailed to:**  
Douglas Girod, MD, FACS, Professor and Chairman  
The University of Kansas School of Medicine  
Department of Otolaryngology-Head & Neck Surgery  
3901 Rainbow Blvd. MS 3010, Kansas City, KS 66160

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# Bulletin Content AT YOUR FINGERTIPS



AMERICAN ACADEMY OF  
OTOLARYNGOLOGY-  
HEAD AND NECK SURGERY

Read the *Bulletin*  
online or on your  
mobile device at:



[www.entnet.org/bulletin](http://www.entnet.org/bulletin)



### Pediatric Otolaryngology - Academic Position

The Department of Otorhinolaryngology is recruiting a third Pediatric Otolaryngologist to join a busy, tertiary Pediatric Otolaryngology practice. This is a unique opportunity to join a rapidly growing Department at a major University Children's Hospital with a large Level III NICU and a Level I Trauma Center. Excellent compensation and benefits. Academic appointment commensurate with experience. Strong interest in resident and medical student teaching and research is encouraged.

*Applicants should forward a CV and statement of interest to:*

Soham Roy, MD, FACS, FAAP  
 Director of Pediatric Otolaryngology  
 The University of Texas Medical School at Houston  
 Department of Otorhinolaryngology-Head & Neck Surgery  
 713-383-3727 (fax)  
 Soham.Roy@uth.tmc.edu  
 http://www.ut-ent.org



*UTMSH is an equal opportunity employer.*



Purchase ENT, a well-established regional ENT practice in Western Kentucky, is actively searching for physicians to join our practice. Paducah, Kentucky offers everything from Broadway shows at the Carson Center to festivals like the annual BBQ on the River. Located a short 20 minute drive from Kentucky and Barkley Lakes, and the Land Between the Lakes National Recreation Area, where great recreational opportunities are offered, and also just a short two to three hour drive to larger cities like Nashville, Memphis, St. Louis and Louisville, Paducah is a great community. There is something for everyone in the region.

Purchase ENT is well respected in the medical community and is a very progressive practice offering in-office Allergy, Audiology, CT Scan, and Ultrasound. Additionally, there is an investment opportunity in a successful ambulatory surgery center. Although we are seeking candidates to practice general ENT, a fellowship in Otolaryngology or Head and Neck would be welcomed. For the successful candidate, the opportunity is a one year partnership track. For additional information about this great opportunity, contact Jim Wring, Practice Administrator at [jwring@purchaseent.com](mailto:jwring@purchaseent.com) or 270-408-3208.

## OTOLARYNGOLOGIST OPPORTUNITY

### Geisinger Wyoming Valley (GWV) Medical Center, Wilkes-Barre, Pa., is seeking a BC/BE Otolaryngologist.

Geisinger's otolaryngology specialists treat a wide range of conditions of the head and neck by providing the latest technologies in diagnostic, medical, surgical and rehabilitative techniques. We have board-certified and fellowship-trained specialists who collaborate to ensure the most comprehensive care.

#### About the Position

- Take part in the growth of this dynamic department
- Pursue research in your area of interest

Geisinger Wyoming Valley (GWV) Medical Center, Wilkes-Barre, Pa., is an acute care hospital that is licensed for 243 beds and houses the only Level II Trauma center in Luzerne County. The campus includes the Frank M. and Dorothea Henry Cancer Center, The Richard and Marion Pearsall Heart Hospital, the Janet Weis Children's Hospital Pediatric Unit, a transplant program and the Brain & Spine Tumor Institute. Geisinger South Wilkes-Barre (GSWB) is GWV's ambulatory campus.

**Discover for yourself why Geisinger has been nationally recognized as a visionary model of integrated healthcare.**

For more information or to apply for this position, please contact:

**Autum Ellis,**  
 Department of  
 Professional Staffing,  
 at 1-800-845-7112 or  
[amellis1@geisinger.edu](mailto:amellis1@geisinger.edu)



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Children's Mercy Hospitals and Clinics – Kansas City is seeking fellowship trained Pediatric Otolaryngologists to join our professional staff at the assistant or associate professor level. The position would entail clinical care, research, and teaching of medical students, and pediatric and otolaryngology residents.

Our active Pediatric Otolaryngology Section provides comprehensive tertiary patient care in a family-centered environment. There are currently 7 pediatric otolaryngologists on staff, as well as 3 neurotologists. In addition, our ACGME-accredited pediatric otolaryngology fellowship welcomed our 4th fellow this July, 2012. Children's Mercy Hospitals & Clinics is a large pediatric health care system that is affiliated with the University of Missouri-Kansas City School of Medicine. The main hospital is growing to nearly 400 beds this year with plans to expand to 41 PICU beds and 80 NICU beds.

Kansas City is a bi-state community with close to 2 million residents who enjoy an excellent quality of life. There is a robust offering of arts and entertainment, with a number of new venues having just opened within the past few years. The Kansas City metroplex contains a wide selection of highly rated public and private schools. We are also the regional home to several major colleges and universities. Salary and academic range are commensurate with experience. EOE/AAP

Robert A. Weatherly, MD  
 Section Chief, Ear, Nose, and Throat  
 rweatherly@cmh.edu  
 Phone: 866-CMH-IN-KC/866-264-4652  
 www.childrensmercy.org

## University of Missouri

Department of Otolaryngology—  
 Head and Neck Surgery



Seeks clinicians, teachers, and researchers who are personable, energetic and innovative to join a rapidly growing and collaborative group of physicians. A Faculty opportunity at all academic levels (Assistant/Associate Professor or Professor or Clinical Assistant/Associate Professor or Clinical Professor) is available in **Head and Neck Surgical Oncology with microvascular experience**. Title, track, and salary are commensurate with experience.

- Competitive production incentive
- Research interests encouraged and supported
- New outpatient clinic with state-of-the-art equipment and ancillary services
- Well established and expanding hospital system
- Live and work in Columbia, ranked by *Money* magazine and *Outside* magazine as one of the best cities in the U.S.

For additional information about the position, please contact:

*Robert P. Zitsch III, M.D.*  
*William E. Davis Professor and Chair*  
 Department of Otolaryngology—Head and Neck Surgery  
 University of Missouri—School of Medicine  
 One Hospital Dr MA314 DC027.00  
 Columbia, MO 65212  
 zitschr@health.missouri.edu

To apply for this position, please visit the MU web site at  
[hrs.missouri.edu/find-a-job/academic/](http://hrs.missouri.edu/find-a-job/academic/)

The University of Missouri is an Equal Opportunity/Affirmative Action Employer and complies with the guidelines of the Americans with Disabilities Act of 1990. To request ADA accommodations, please contact (573) 884-7282 (V/TTY). Diversity applicants are encouraged to apply.

# Southern States Rhinology Course

*An Evidence Based, Interactive, Hands On Learning Experience*  
 May 2 - 4, 2013 • Kiawah Island Golf Resort • Kiawah Island, SC



### Course Topics:

Evidence Based Panels on Medical and Surgical Management of CRS; Pediatric CRS, and ESS Failures, Balloon Sinuplasty; Consult the Experts Panel; and much more!

### Cadaver Lab

A hands on laboratory dissection is also available with this course. Participants will have the opportunity to dissect cadaver specimens featuring state-of-the-art endoscopic instrumentation, video, and image guidance system

### Faculty:

- John DelGaudio, MD - Emory School of Medicine
- Stil Kountakis, MD - Georgia Regents University
- Frederick Kuhn, MD - GA Nasal & Sinus Institute
- Rodney Schlossler, MD - Medical University of SC
- Brent Senior, MD - UNC School of Medicine
- Michael Sillers, MD - Alabama Nasal & Sinus Center



Scan the QR Code directly from your smartphone for full meeting details and Registration or visit [www.southernstatesrhinology.org](http://www.southernstatesrhinology.org)

13TH ANNUAL  
**Charleston Magnolia  
Conference**  
MAY 31 - JUNE 1, 2013

— GUEST SPEAKERS —

**George T. Hashisaki, M.D.**  
Associate Professor  
Department of Otolaryngology – HNS  
University of Virginia Health Systems – Charlottesville, VA

**Robert C. Kern, M.D.**  
Professor and Chairman  
Department of Otolaryngology – HNS  
Northwestern Feinberg School of Medicine – Chicago, IL

**Gregory Postma, M.D.**  
Professor, Department of Otolaryngology  
Director, MCG Center for Voice & Swallowing Disorders  
Georgia Health Science Medical Center – Augusta, GA

— TOPICS —

- Endoscopic Sinus Surgery • Pediatric Sleep Apnea
- Endoscopic and Robotic Head and Neck Cancer Surgery
- Management of Cholesteatoma
- Management of Salivary Gland Disease
- Facial Reconstructive Surgery • Endocrine Surgery
- Vascular Anomalies in Children

— RECREATION —

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Otolaryngology – Head & Neck Surgery  
Contact: 843-876-0493 • Email: mansfee@musc.edu  
http://ENT.musc.edu



Mayo School of Continuous Professional Development

## Endoscopic Sinus and Skull Base Surgery 2013

**Mayo Clinic • Scottsdale, Arizona • April 3-6, 2013**

**Guests of Honor:**  
Prof. Heinz Stammberger (Austria)  
Prof. Piero Nicolai (Italy)

**Honored National Faculty:**  
Anne E. Getz, MD  
Peter H. Hwang, MD  
Juan Fernandez-Miranda, MD  
Carl H. Snyderman, MD  
James A. Stankiewicz, MD

**Course Director:** Devyani Lal, MD

**Mayo Clinic and Arizona Faculty:**  
Stephen F. Bansberg, MD  
Timothy W. Haegan, MD  
Joseph M. Hoxworth, MD  
Erin K. O'Brien, MD  
John F. Pallanch, MD  
Naresh P. Patel, MD  
Ryan M. Rehl, MD

*Endoscopic Sinus and Skull Base Surgery 2013* is our second state-of-the-art course designed for otolaryngologists and endoscopic skull base surgeons. The curriculum will focus on inflammatory sinus disease on April 3-4, highlighting advanced, salvage and novel treatment strategies. Endoscopic skull base surgery will be the focus April 5-6. The curriculum is designed to introduce the novice surgeon to basic techniques, and provide advanced training for the more experienced surgeon. Hands-on dissection sessions will be conducted in our world-class laboratory with fresh frozen cadavers, powered instrumentation and image guidance.

**Accommodations:** Westin Kierland Resort • [www.kierlandresort.com](http://www.kierlandresort.com)  
• (480) 924-1202 • Residence Inn Phoenix Desert View at Mayo Clinic  
• [www.marriott.com/phxmh](http://www.marriott.com/phxmh) • (800) 331-3131

**Meeting Location:** Mayo Clinic's Phoenix and Scottsdale campuses  
**To Register, Contact MSCPD:** [www.mayo.edu/cme/otorhinolaryngology](http://www.mayo.edu/cme/otorhinolaryngology)  
• email: [mca.cme@mayo.edu](mailto:mca.cme@mayo.edu) • (480) 301-4580

**Featuring: Hands-on dissection, live prosection and endoscopic 3D anatomy**



# 2013 Lone Star Rhinology Course

April 26-28, 2013

UT Southwestern Medical Center  
Dallas, Texas

Cadaver Lab  
with Image Guidance

Program details and registration  
available at: [sinuscourse.com](http://sinuscourse.com)

**GUEST OF HONOR**  
**Brent A. Senior, M.D., F.A.C.S., F.A.R.S.**

**COURSE DIRECTORS**  
**Pete S. Batra, M.D., F.A.C.S.**  
**Samer Fakhri, M.D., F.A.C.S.**

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