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LETTER FROM THE EDITOR-IN-CHIEF

The Hidden Costs of Medical Liability

The malpractice debate continues, like a rerun of Quincy, M.E.—you know the ending, but pretend to be surprised, if only to justify why you would watch the same show twice (or, in this case, many more).

The issue of medical liability, while it has received little attention in the health reform debate, is perhaps the best example of why changing healthcare, no matter where you want it to go, is so difficult to achieve. Indeed, the medical liability debate will forever be just that: a debate, with no end, with watered down reforms at best. We will never address the core issues because they are simply “too hot to handle.” Much like healthcare reform, everybody wants it, and a majority agree on what is broken, but no one can agree on how to fix it.

Here’s why: On complex issues, humans have a natural tendency to polarize into camps based on perceived moral superiority. This protects them from the insecurity of uncertainty and compromise. Compromise sounds weak, and why should I have to compromise on something when I know I am “right?”

We seek moral comfort on issues; we want black and white, right or wrong. We run away from anything with a touch of grey. The consequence of our morality dilemma is polarization and indecision. We huddle into camps with the like-minded. We demonize the disbelievers as immoral or insensitive.

On one side of the medical liability issue, you have those who highlight how the fear of liability destroys the very fabric of the doctor-patient relationship. Every decision is made with the fear of liability looming. This, we argue, results in unnecessary tests, increased costs, and more time spent away from patients while we document everything we can to protect ourselves.

On the other side are those who fear that without penalties for “negligence,” patient safety will suffer, and evil, no-good doctors will leave patients on operating tables to get in a round of golf.

Now, at the risk of oversimplifying, it is true that patients have suffered significant, life-altering harm from preventable medical errors, many that seem grossly negligent in hindsight. I am often befuddled by how my colleagues’ care can at times seem so substandard, so sloppy, and often filled with arrogance. But I am also very cognizant of the immense pressure we work under: the productivity demands, and a payment system that is not only unjust, but breeds poor care. Not to mention the most complex, unpredictable, and rapidly evolving discipline in the world. No one can be the infallible master of all this.

Now, mistakes are made in every profession, every day. What makes medicine different are the consequences. Life hangs in the balance with every decision. We accepted that when we entered this profession. But punitive liability does nothing to mitigate the very mistakes it is intending to prevent.

Here’s why: When liability hangs over every decision, you simply create an environment of fear. And this has its greatest impact on the most caring, thoughtful, and astute in our profession. The fear of liability significantly changes the practices of the “good,” distracting them from their most critical mission, diluting their potential impact on patient care. All the while, the truly negligent, the arrogant, and the uncaring, are, by nature, oblivious to it all.

Someone forgot to examine the basic human psychology of a punitive system of liability in a profession littered with uncontrollable land mines. Gross, malicious negligence, of the kind malpractice law should punish, is committed by a tiny minority who often consider themselves untouchable anyway. Fear of liability doesn’t faze them.

Many of those opposed to tort reform will argue that a small minority of cases go to trial, and an even smaller minority of jury verdicts favor the plaintiff. They will argue that the system is, therefore, pretty accurate at identifying gross negligence. What they fail to address are the thousands of cases that don’t go to trial, but inflict an incurable harm on those who have had to endure and the fear they have instilled in all of us. You cannot assign a dollar cost to the impact of fear, but I can assure you it has eroded the quality of care our patients receive and the quality of life our physicians deserve.

Lee A. Resnick, MD
Editor-in-Chief
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Treating Common Upper Respiratory Tract Infections in an Era of Increasing Antibiotic Resistance

Common complaints can often be treated with common solutions. When it comes to choosing the appropriate therapy for upper respiratory tract infections, however, a healthy view of the big picture is advisable.

By Joseph Toscano, MD

A Child with Constipation and Swollen Abdomen

Often, malignancies in children are discovered only during investigation of a less ominous complaint, highlighting the importance of being vigilant for relevant signs and symptoms.

By Muhammad Waseem, MD

From dermatophytes to Pityrosporum, infectious fungi are likely to walk through your door on a fairly regularly basis, both literally and figuratively. A review of how to identify and treat the more common types.

WEB EXCLUSIVE

Treatment of Pediatric Community-acquired Pneumonia in an Urgent Care Center

Most cases of community-acquired pneumonia in younger patients can be diagnosed in the urgent care center by non-pediatricians. Subsequent treatment can help reduce the rate of hospitalization. Available only at www.jucm.com.

By Deena R. Zimmerman, MD, MPH, IBCLC, Scott Fields, MD, and Nahum Kovalski, BSc, MDCM

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Frank Leone, MBA, MPH cautions against viewing national contracts as an insurmountable obstacle when pitching occupational medicine services to larger employers.

John Shufeldt, MD, JD, MBA, FACEP casts urgent care’s fight for recognition in a historical context.

David Stern, MD, CPC responds to questions on billing S9083 to multiple carriers, coding for repair of multiple lacerations, maximizing IV reimbursement, proper use of modifier -59, and how to code for measuring oxygen saturation levels.

Do you have an idea for an article or new feature? Maybe an interesting x-ray case to present? Let us know in an e-mail to Lee A. Resnick, MD, JUCM’s editor-in-chief, at editor@jucm.com.

To Submit an Article to JUCM, The Journal of Urgent Care Medicine encourages you to submit articles in support of our goal to provide practical, up-to-date clinical and practice management information to our readers—the nation’s urgent care clinicians. Articles submitted for publication in JUCM should provide practical advice, dealing with clinical and practice management problems commonly encountered in day-to-day practice.

Manuscripts on clinical or practice management topics should be 2,600–3,200 words in length, plus tables, figures, pictures, and references. Articles that are longer than this will, in most cases, need to be cut during editing.

We prefer submissions by e-mail, sent as Word file attachments (with tables created in Word, in multicolumn format) to editor@jucm.com. The first page should include the title of the article, author names in the order they are to appear, and the name, address, and contact information (mailing address, phone, fax, e-mail) for each author.

Before submitting, we recommend reading “Instructions for Authors,” available at www.jucm.com.
A Note of Thanks to Our Peer Reviewers

As we begin our fourth year of publication, *JUCM* remains committed to publishing articles relevant to the practice of urgent care medicine, told in an urgent care “voice.”

A key factor in achieving that objective is to work with the best and the brightest professionals working in urgent care centers and related companies, as well as academic institutions across the country.

Obviously, this is true of the authors whose work has appeared in the journal. You know their names. But there’s another group of individuals without whom we could not in good conscience call *JUCM* a peer-reviewed journal.

The accomplished practitioners who have volunteered to review and comment on articles prior to publication help keep us “honest” and ensure that our content is relevant to the way medicine is practiced in the urgent care setting.

The nature of the peer-review process requires anonymity on a month-to-month basis, but each year we like to collectively thank them for their invaluable contributions.

We are grateful to the following individuals for sharing their time and expertise—some on more than one occasion—in reviewing articles that appeared in Volume 3 (October 2008 through September 2009) of *JUCM*:

Jeffrey P. Collins, MD, MA
Rajesh Davit, MD
Tanise I. Edwards, MD, FAAEM
Ronald J. Ellis, DO, FACOEP
William Gluckman, DO, MBA, FACEP
Joy Green-Hadden, DNP
Akila Iyer, MD
Melvin Lee, MD
Kevin McKee, DO, MS
Genevieve M. Messick, MD
Michael Miller, DO
Matthew P. Mullen, MD
Brian Roberts, MD
Shailendra Saxena, MD, PhD
Joseph Toscano, MD
Adam Wineinger, MD
Mark D. Wright, MD
Donald Yeatts, DDS, MD

If you would like to join our panel of peer reviewers, please e-mail Harris Fleming, editor of *JUCM*, at hfleming@jucm.com.
An old friend of mine, Caroline Herring, is a musician who is launching her fourth album two days from now as I write this. Even though her reviews have been great, she's still nervous—terrified, in fact.

And who wouldn't be? She's spent countless hours of her life coming up with ideas for songs, exploring different ways to present them, finding her voice (not to mention a record contract), performing those songs in a recording studio…and then she waits to see if anyone is listening.

If you've opened an urgent care center, you know the feeling. Think about it: you've spent countless hours of your life coming up with ideas, researching them, gathering expert feedback, putting all the pieces together, getting finances in order, developing a marketing campaign, making all the final touches…and then you wait to see if anyone comes.

I think we all know the feeling. I know that I do.

Every year, about 15 minutes into the start of the annual Urgent Care Convention, we start planning the next one. We make topic lists, we note the great speakers, we start thinking about themes—and for many months after that we narrow down our ideas and choose faculty and put program together. The brochure mails…and we wait to see if anyone registers. It's torture until that first call.

It's the same whether you're opening your first urgent care center or your 31st. The seeds may have been planted when you saw the UCAOA booth at the ACEP or AAFP Scientific Assembly in Boston last month, or when a center opened up in your town, or when you heard about an old friend who is opening one.

“Hmmm…,” you think to yourself. Then you talk to some trusted advisors, pick a site, get some financing, deal with mountains of paperwork, staffing, build-out, marketing. Opening day comes…and you wait for that first patient to pull into the parking lot to confirm it wasn't all a crazy fantasy. It's torture!

It may have been five minutes since that first patient walked into your center, or it could be five months or even five years; you probably still remember that feeling of panic that no one would show up—and the tremendous relief when that first person did! It's a common experience we all share, whether we are “selling” albums, conferences, or healthcare.

We want to gather those experiences from you, whether you were the owner, the office manager, the biller, the provider, or the registrar, and share them with others.

What do you remember about your Opening Day and the weeks leading up to it? What do you recall and what would you pass along to those who are thinking of following in your footsteps?

I think this is a great project, as well as a chance for you to pass along your wisdoms, great and small, and give back to the industry that you began. It's a chance for you help others skip the mistakes you made, and build on your successes.

Whether you have one sentence or many pages to share (or simply have a question about the project), I invite you to send those remembrances to me at lhorwitz@ucaoa.org.

Let's see what we can gather to share with each other and with those who will follow in your footsteps.

By the way, soon we will be able to share some of the details of the 2010 Urgent Care Convention in Orlando, and we hope you will be as excited as we are! If you are not getting the newsletters we e-mail each month, you won't be among the first to hear about early, discounted sign-ups for the convention. This would be a good time to make sure we are on your “safe senders” list and that we have a current e-mail address for you.

Lou Ellen Horwitz is executive director of the Urgent Care Association of America. She may be contacted at lhorwitz@ucaoa.org.
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Introduction

Upper respiratory tract infections (URTIs) are among the most common reasons patients seek assistance in urgent care practice. The common cold, otitis media, acute sinusitis, and acute pharyngitis are well known to patient and provider alike. Acute bronchitis is a lower respiratory tract infection, with features similar to URTIs. These infections are most often self-limited and uncomplicated, but the approach to evaluating each patient should include examining for complications as well as rarer, more severe diseases that can mimic these simpler, common conditions. Treatment should include patient education, symptom management, and the use of antibiotics only if likely to improve the clinical outcome.

Data show a positive correlation between increasing levels of antibiotic use and increasing antibiotic resistance among bacteria. Though in the United States there seems to be a gradually decreasing rate of antibiotic prescription for URTIs in general, overprescription is still common and there is an increasing trend toward the use of broad-spectrum antibiotics for these relatively simple infections.

It is intuitive that a strategy of prescribing antibiotics only when necessary and only of the appropriate antimicrobial spectra will minimize the development of antibiotic resistance. Guidelines describing such use have been published by several organizations (see Table 1 for links) and will be the primary subject of this review.

Clinical Diagnosis, Testing, and Important Complications and Disease Mimics

URTIs are often grouped together because they share a closely related anatomy and pathophysiology. The mu-

Urgent message: Thorough evaluation and thoughtful prescribing can help ensure responsible, effective care and patient satisfaction.

Joseph Toscano, MD
TREATING COMMON RESPIRATORY TRACT INFECTIONS IN AN ERA OF INCREASING ANTIBIOTIC RESISTANCE

Table 1. Clinical Guidelines for the Treatment of Upper Respiratory Tract Infections

<table>
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<td></td>
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</tbody>
</table>

ACP, American College of Physicians; ICSI, Institute for Clinical Systems Improvement; AAP, American Academy of Pediatrics; AAO–HNS, American Academy of Otolaryngology–Head and Neck Surgery; IDSA, Infectious Diseases Society of America

Compendia of all relevant clinical practice guidelines for URTIs in adults and children, updated annually, is available for download at: www.aware.md/HealthCareProfessionals/ClinicalResources.aspx

*Certain ACP and AAP Guidelines are over 5 years old and therefore not considered “current” by those organizations; however, pending updates, these are the most recent recommendations.

The acute onset of cough and higher fever—typically with associated headache and myalgias—generally distinguishes human, swine, and avian influenza. These particularly viral URTIs have higher rates of associated morbidity and mortality and require a different approach than will be discussed in this article. (An excellent review of the testing, evaluation, and care of patients with swine-origin H1N1 virus appeared in the October 2009 issue of JUCM.)

When cough predominates, bronchitis is usually the diagnosis. Wheezing, even in patients without a history of bronchospastic disease, may be noted on exam. When cough is associated with an inspiratory whoop (usually seen only in children) or post-tussive vomiting or is severe and paroxysmal, clinicians should suspect pertussis, especially when symptoms last longer than 14 days. A higher index of suspicion (e.g., any cough illness lasting more than 14 days or severe cough illness of a shorter duration) will apply during an identified pertussis outbreak.4

Prominent ear pain and abnormal otoscopic findings indicate otitis media.

Symptoms of the common cold can include nasal congestion and drainage, sneezing, mild sore throat and cough, and fever. Nasal symptoms usually predominate; otherwise, the widespread nature (sinuses, nose, throat, chest) of generally mild, though often aggravating, symptoms establishes this diagnosis.

Sinus pain can suggest sinusitis.

A chief complaint of sore throat typically indicates pharyngitis.

Sorting through the differential diagnosis of URTIs is largely a clinical exercise. Analyzing a patient’s symptomatology and performing a systematic exam are crucial to the process of diagnosis and the exclusion of significant complications or other diseases that can present in ways similar to milder forms of infection (see Table 2).

On the other hand, there are no clinical criteria (e.g., the presence of fever, level of discomfort, exam findings, the color or characteristics of any produced sputum or mucous) that can be used to reliably distinguish between bacterial and viral etiologies.

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• Serious cases of bronchospasm, including fatalities, have been reported during treatment with RELENZA in patients with and without underlying airways disease. Many of these cases were reported during postmarketing and causality was difficult to assess
• If use of RELENZA is considered for a patient with underlying airways disease, the potential risks and benefits should be carefully weighed. Use in these patients should be done only under conditions of careful monitoring of respiratory function, close observation, and appropriate supportive care including availability of fast-acting bronchodilators
• Discontinue RELENZA and initiate appropriate treatment if an allergic reaction occurs or is suspected
• Patients with influenza, particularly pediatric patients, may be at an increased risk of seizures, confusion, or abnormal behavior early in their illness. Monitor for signs of abnormal behavior
• Safety and efficacy have not been demonstrated in patients with high-risk underlying medical conditions
• RELENZA has not been proven effective for prophylaxis of influenza in the nursing home setting
• RELENZA is not a substitute for early influenza vaccination on an annual basis as recommended by the Centers for Disease Control’s Immunization Practices Advisory Committee
• Influenza viruses change over time. Emergence of resistance mutations could decrease drug effectiveness. Other factors (for example, changes in viral virulence) might also diminish clinical benefit of antiviral drugs. Prescribers should consider available information on influenza drug susceptibility patterns and treatment effects when deciding whether to use RELENZA

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*Clinical significance can not be inferred from surveillance data.
5.5 Bacterial Infections: Serious bacterial infections may begin with influenza-like symptoms and may progress with or occur as complications during the course of influenza. RELENZA has not been shown to prevent such complications.

5.6 Importance of Proper Use of DISKHALER: Effective and safe use of RELENZA requires proper use of the DISKHALER to inhale the drug. Prescriptions should carefully evaluate the ability of young children to use the delivery system if use of RELENZA is considered (see User Instructions in Product Information).

6 ADVERSE REACTIONS

6.1 Clinical Trials Experience: The placebo used in clinical studies consisted of inhaled lactose powder, which is also the vehicle for the active drug; therefore, some adverse events occurring at similar frequencies in different treatment groups could be related to lactose vehicle administration. Treatment of influenza: Clinical Trials in Adults and Adolescents: Adverse events that occurred with an incidence ≥1.5% in treatment studies (patients ≥12 years of age) are listed in Table 1.

<table>
<thead>
<tr>
<th>Table 1. Summary of Adverse Events ≥1.5% Incidence During Treatment in Adults and Adolescents</th>
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<td>Vomiting</td>
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<tr>
<td>Respiratory</td>
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<tr>
<td>Nasal signs and symptoms</td>
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<tr>
<td>Bronchitis</td>
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<tr>
<td>Cough</td>
</tr>
<tr>
<td>Nasal inflammation</td>
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<tr>
<td>Ear, nose, and throat infections</td>
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<tr>
<td>Nervous system</td>
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</table>

Includes studies where RELENZA had administered intranasally (4-8 mg 2 to 4 times per day in addition to inhaled preparation) and/or inhaled more frequently (i.e., >2 times) than the currently recommended dose. Includes additional adverse reactions occurring in less than 1.5% of patients receiving RELENZA, including malaise, fatigue, fever, abdominal pain, myalgia, arthralgia, and urticaria. The most frequent adverse reactions in Phase III placebo-controlled studies included elevations of liver enzymes and CPK, lymphopenia, and neutropenia. These were reported in similar proportions of paroxysmal and lactose vehicle placebo recipients with influenza-like illness. Clinical Trials in Pediatric Patients: Adverse events that occurred with an incidence ≥1.5% in patients 5 to 12 years of age during treatment studies ofRELENZA in 2 Phase III studies are listed in Table 2.

<table>
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<tr>
<th>Table 2. Summary of Adverse Events ≥1.5% Incidence During Treatment in Pediatric Patients</th>
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</tbody>
</table>

Includes a subset of patients receiving RELENZA for treatment of influenza in a prophylaxis study. In 1 of the 2 studies described in Table 3, some additional information is available from children (5 to 12 years old) without influenza-like illness who received an investigational prophylaxis regimen ofRELENZA: 132 children received RELENZA and 145 children received placebo. Among these children, nasal signs and symptoms (20%), ear pain (9%), cough (10%), and rhinorrhea were more frequent with RELENZA than placebos. In a subset with chronic pulmonary disease, lower respiratory adverse events (described as asthma, cough, or viral respiratory infections which could include influenza-like symptoms) were reported in 4 to 7 children and 4 to 8 placebo recipients. Prophylaxis of Infection: Family/Household Prophylaxis: Adverse events that occurred with an incidence ≥1% in the prophylaxis studies (patients ≥5 years of age) are listed in Table 3.

<table>
<thead>
<tr>
<th>Table 3. Summary of Adverse Events ≥1.5% Incidence During 28-Day Prophylaxis Studies in Adults, Adolescents, and Children*</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Adverse Event</strong></td>
</tr>
<tr>
<td>Lower respiratory infections</td>
</tr>
<tr>
<td>Upper respiratory infections</td>
</tr>
<tr>
<td>Headache</td>
</tr>
<tr>
<td>Nasal signs and symptoms</td>
</tr>
<tr>
<td>Nasal inflammation</td>
</tr>
<tr>
<td>Gastrointestinal</td>
</tr>
<tr>
<td>Nasal and throat</td>
</tr>
<tr>
<td>Gastrointestinal</td>
</tr>
<tr>
<td>Malaise and fatigue</td>
</tr>
<tr>
<td>Temperature regulation disturbances</td>
</tr>
</tbody>
</table>

* In prophylaxis studies, symptoms associated with influenza-like illness were captured as adverse events; subjects were enrolled during a winter respiratory season during which time any symptoms that occurred were captured as adverse events.

6.2 Postmarketing Experience: Allergic Reactions: Allergic or allergic-like reactions, including hives, urticaria, angioedema, anaphylaxis, and anaphylactoid reactions, have been reported after accidental or intentional administration of RELENZA. The concurrent use of RELENZA with live attenuated influenza vaccine (LAV) influenza has not been evaluated. However, because of potential interference between these products, LAV should not be administered within 2 weeks before or 48 hours after administration of RELENZA unless medically indicated. The concern about possible interference arises from the potential for antiviral drugs to inhibit replication of live vaccine virus. Treatment of influenza: Clinical Trials in Adults and Adolescents: Adverse events that occurred with an incidence ≥1.5% in 2 placebo studies (patients ≥12 years of age) are listed in Table 2.

<table>
<thead>
<tr>
<th>Table 4. Summary of Adverse Events ≥1.5% Incidence During 28-Day Prophylaxis Studies in Adults, Adolescents, and Children*</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Adverse Event</strong></td>
</tr>
<tr>
<td>Headache</td>
</tr>
<tr>
<td>Nausea and vomiting</td>
</tr>
<tr>
<td>Non-specific</td>
</tr>
<tr>
<td>Temperature regulation disturbances</td>
</tr>
</tbody>
</table>

* In prophylaxis studies, symptoms associated with influenza-like illness were captured as adverse events; subjects were enrolled during a winter respiratory season during which time any symptoms that occurred were captured as adverse events.

6.3 Pancreatitis: Serious, life-threatening, fatalities have been reported in patients receiving RELENZA. The concurrent use of RELENZA with live attenuated influenza vaccine (LAV) influenza has not been evaluated. However, because of potential interference between these products, LAV should not be administered within 2 weeks before or 48 hours after administration of RELENZA unless medically indicated. The concern about possible interference arises from the potential for antiviral drugs to inhibit replication of live vaccine virus. Treatment of influenza: Clinical Trials in Adults and Adolescents: Adverse events that occurred with an incidence ≥1% in the prophylaxis studies (patients ≥5 years of age) are listed in Table 3.

7 DRUG INTERACTIONS: No drug interactions have been evaluated. Since the drug is inhaled and not absorbed into the systemic circulation, no drug interactions need be considered.

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy: Pregnancy Category C. There are no adequate and well-controlled studies of RELENZA in pregnant women. It should be used during pregnancy only if the potential benefit justifies the potential risk to the fetus. Zanamivir has been shown to cross the placenta in rats and rabbits. In these animals, fetal blood concentrations of zanamivir were significantly lower than zanamivir concentrations in the maternal blood.
8.3 Nursing Mothers: Studies in rats have demonstrated that zanamivir is excreted in milk. However, nursing mothers should be instructed that it is not known whether zanamivir is excreted in human milk. Because many drugs are excreted in human milk, caution should be exercised when RELENZA is administered to a nursing mother.

8.4 Pediatric Use: Treatment of Influenza: Safety and effectiveness of RELENZA for treatment of influenza have not been assessed in pediatric patients less than 1 year of age, but were studied in a Phase II treatment study in pediatric patients, where 471 children 6 to 15 years of age received RELENZA or placebo [see Clinical Studies (14.3) of full prescribing information]. Children in the pediatric principal Phase III adult treatment studies. In these studies, 67 patients were 12 to 16 years of age. No definite differences in safety and efficacy were observed between these adolescent patients and young adults.

In a Phase II study of 16 children ages 6 to 12 years with signs and symptoms of respiratory disease, 4 did not produce a measurable peak inspiratory flow rate (PIFR) through the DISKHALER with no adequate inhalation on request, 1 with missing data, 9 had measurable PIFR on each of 2 inhalations, and 3 achieved measurable PIFR on only 1 of 2 inhalations. Neither of two 6-year-olds and one of two 7-year-olds produced measurable PIFR. Overall, 6 of the 16 children (including all those under 6 years old) either did not produce measurable inspiratory flow through the DISKHALER or produced peak inspiratory flow rates below the 60 L/min considered optimal for the device under standardized in vitro testing, lack of measurable flow was related to low or undetectable serum concentrations (see Clinical Pharmacology (12.3), Clinical Studies (14.4) of full prescribing information). Prescribers should carefully evaluate the ability of young children to use the delivery system if prescription of RELENZA is considered.

8.6 Geriatric Use: Of the total number of patients in 6 clinical studies of RELENZA for treatment of influenza, 59 patients were 65 years of age and older, while 24 patients were 75 years of age and older. Of the total number of patients in 4 clinical studies of RELENZA for prophylaxis of influenza in households and community settings, 164 patients were 65 years of age and older, while 34 patients were 75 years of age and older. No overall differences in safety or effectiveness were observed between these patients and younger patients, and other reported clinical experience has not identified differences in responses between the elderly and younger patients, but greater sensitivity of some older individuals cannot be ruled out. Elderly patients may need assistance with use of the device.

In 2 additional studies of RELENZA for prophylaxis of influenza in the nursing home setting, efficacy was not demonstrated (see Indications and Usage (1.3) of full prescribing information).

10 OVERDOSAGE

There have been no reports of overdose from administration of RELENZA.

17 PATIENT COUNSELING INFORMATION

See FDA-Approved Patient Labeling (17.6).

17.1 Bronchospasm: Patients should be advised of the risk of bronchospasm, especially in the setting of underlying airways disease, and stop RELENZA and contact their physician if they experience increased respiratory symptoms during treatment such as worsening wheezing, shortness of breath, or other signs or symptoms of bronchospasm (see Warnings and Precautions (5.3)). If a decision is made to prescribe RELENZA for a patient with asthma or chronic obstructive pulmonary disease, the patient should be made aware of the risks and should have a fast-acting bronchodilator available.

17.2 Concurrent Bronchodilator Use: Patients scheduled to take inhaled bronchodilators at the same time as RELENZA should be advised to use their bronchodilators before taking RELENZA.

17.3 Neuropsychiatric Events: Patients with influenza (the flu), particularly children and adolescents, may be at an increased risk of seizures, confusion, or abnormal behavior early in their illness. These events may occur after beginning RELENZA or may occur when flu is not treated. These events are uncommon but may result in accidental injury to the patient. Therefore, patients should be observed for signs of unusual behavior and a healthcare professional should be contacted immediately if the patient shows any signs of unusual behavior (see Warnings and Precautions (5.3)).

17.4 Instructions for Use: Patients should be instructed in use of the delivery system. Instructions should include a demonstration whenever possible. For the proper use of RELENZA, the patient should read and follow carefully the accompanying Patient Instructions for Use. If RELENZA is prescribed for children, it should be used only under adult supervision and instruction, and the supervising adult should first be instructed by a healthcare professional [see Dosage and Administration (2.4)].

17.5 Risk of Influenza Transmission to Others: Patients should be advised that the use of RELENZA for treatment of influenza has not been shown to reduce the risk of transmission of influenza to others.

17.6 FDA-Approved Patient Labeling and Instructions for Use: See separate label.

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CHF, congestive heart failure; RAD, reactive airways disease; COPD, chronic obstructive pulmonary disease; HIV, human immunodeficiency virus

Table 2. Upper Respiratory Tract Infection Complications and Differential Diagnosis

<table>
<thead>
<tr>
<th>URTI</th>
<th>Complications</th>
<th>Differential diagnoses and “can’t miss” mimics</th>
</tr>
</thead>
<tbody>
<tr>
<td>Common cold</td>
<td>Other URTIs</td>
<td>Other URTIs</td>
</tr>
<tr>
<td>Acute bronchitis</td>
<td>CHF, RAD, COPD</td>
<td>Pneumonia</td>
</tr>
<tr>
<td></td>
<td>Exacerbation of RAD, COPD, or CHF Pertussi</td>
<td></td>
</tr>
<tr>
<td>Acute otitis media</td>
<td>Mastoiditis, tympanic membrane perforation</td>
<td>Eustachian tube dysfunction</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Barotrauma</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Otitis externa</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Mastoiditis</td>
</tr>
<tr>
<td>Acute sinusitis</td>
<td>Intracranial infection, periorbital cellulitis</td>
<td>Common cold</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Chronic sinusitis</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Meningitis</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Wegener’s granulomatosis</td>
</tr>
<tr>
<td>Acute pharyngitis</td>
<td>Peritonsillar or parapharyngeal space infections (though may separate disease process); rheumatic fever and acute glomerulonephritis (for Strep)</td>
<td>Peritonsillar or parapharyngeal space infections; Epiglottis; HIV primary infection; Infectious mononucleosis; Gonococcal pharyngitis; Kawasaki disease</td>
</tr>
</tbody>
</table>

**Differential diagnoses and “can’t miss” mimics**

**CHF**, congestive heart failure; **RAD**, reactive airways disease; **COPD**, chronic obstructive pulmonary disease; **HIV**, human immunodeficiency virus

Atting patients with URTIs. Each disease mimic, however, may have its own diagnostic test(s), a discussion of which exceeds the scope of this review.

Of course, patients who are toxic-appearing or who are immunosuppressed or have other significant comorbidities should be evaluated veryaggressively; the recommendations that follow do not apply to these subsets of patients.

**Appropriate testing**

No diagnostic testing is required for patients with the common cold. However, confirming respiratory syncytial virus (RSV) or influenza infection in febrile pediatric patients with rapid “point-of-care” testing has been shown to reassure clinicians and safely decrease antibiotic prescription and unnecessary further work-ups for other infections.

Complications and mimics of the common cold include any of the other URTIs and allergic and vasomotor rhinitis. Typically, the presence of fever suggests URTI, while ongoing nasal congestion and rhinorrhea in a patient without fever suggest noninfectious rhinitis.

No diagnostic testing is needed to confirm acute bronchitis, though a chest radiograph should be performed if pneumonia is suspected. Pneumonia may be more likely in an older or ill-appearing patient; if there is fever, hypoxemia, tachycardia, or tachypnea; or if abnormalities are present on lung exam.
Those with acute bronchitis symptoms and a history of asthma, chronic obstructive pulmonary disease, or congestive heart failure may be having an exacerbation of chronic disease, either as their primary problem, or triggered by a concomitant chest infection. An assessment of past history and risk factors, as well as physical exam and, when needed, chest radiograph findings can usually establish the diagnosis.

In most situations, for suspected cases of pertussis, the recommended strategy involves both polymerase-chain reaction (PCR) testing and culture, health department reporting, empiric treatment, and close follow-up. A lower threshold for empiric treatment will apply during an identified pertussis outbreak. Local infectious disease specialists, health departments, and the CDC are important resources to consult to balance the need to identify and treat this disease while avoiding treating everyone who has a prolonged cough with antibiotics.

No testing is necessary to make the diagnosis of acute otitis media or sinusitis. There is no proven beneficial role for sinus radiographs in the diagnosis or treatment of sinusitis, and CT scanning of the sinuses should be reserved for refractory or severe cases being treated in conjunction with specialty care.

The diagnosis of otitis media should include acute onset of ear pain and physical exam evidence of tympanic membrane (TM) inflammation and middle-ear effusion (i.e., bulging TM, air-fluid level, otorrhea, or decreased TM mobility on pneumatic otoscopy). Mimics of otitis media are usually distinguishable on exam, and specific palpation of the mastoid process is important in any patient with ear pain.

Features traditionally associated with the clinical diagnosis of sinusitis—nasal obstruction, purulent nasal discharge, pain on bending forward, maxillary toothache, presence of a two-stage illness with sinus symptoms following a URTI—have variable sensitivity and specificity.

One of the more common mimics of acute sinusitis is the exacerbation of chronic sinusitis. There is no specific number, but any patient presenting with a history “usual sinus infection” three or more times per year probably requires a different approach than just an antibiotic prescription and a pat on the back. For these patients, strongly consider a work-up for chronic sinusitis and its causes (anatomic osteomeatal disease, chronic rhinitis, etc.), typically in conjunction with a specialist. And, though it occurs rarely, patients with sinusitis combined with signs of pulmonary and/or renal disease should be promptly referred for work-up for Wegener’s granulomatosis.

Diagnostic testing does play a role in the management of acute pharyngitis. Older guidelines presented options for purely clinical diagnosis, emphasizing the cost effectiveness of such an approach, but the most recent recommendations emphasize obtaining positive rapid antigen testing or culture for group A beta-hemolytic Streptococcus (GABHS) before beginning antibiotic treatment. Some guidelines recommend that, in the face of an initial negative rapid antigen test, patients with a high chance of GABHS (based on age and other risk factors) should have a throat culture obtained before discharge from the clinic. Patients with a prior history of rheumatic fever are at high risk of recurrence and should be followed very closely when they develop pharyngitis or any possible Streptococcal infection.

Any patient with sore throat should be thoroughly examined for swelling or other abnormalities of the uvula and peritonsillar and other parapharyngeal spaces. Drooling, stridor, and trismus are nonspecific but typically indicate severe disease and the need for urgent specialty consultation or ED transfer.

In the absence of such severe symptoms, finding any of these disease mimics at the earliest possible stage requires consideration of the full range of possible diagnoses—maintaining a high index of suspicion—for every patient with a sore throat.

Of the many viral etiologies for pharyngitis, some can result in higher rates of morbidity, including Epstein Barr virus and cytomegalovirus, both of which can cause an acute-mononucleosis-type syndrome of fever, malaise, lymphadenopathy, and frequently splenomegaly and usually mild hepatitis.

Primary infection with human immunodeficiency virus (HIV) can result in a similar clinical picture and should be considered in patients with appropriate risk factors.

Suspicion of gonococcal pharyngitis is also engendered by risk factor assessment. Kawasaki disease presents more often as stomatitis than as pharyngitis, but it is important to keep this diagnosis in mind due to the potential complication of affected children developing coronary artery aneurysms. Suspect the diagnosis and obtain urgent consultation for children under 10-years-old (particularly under 3 years of age) with fever for five days or more, and a syndrome including conjunctivitis, polymorphous rash or desquamation, cervical lymphadenopathy, and any combination of fissured lips, stomatitis, pharyngitis, and/or strawberry tongue.
RESPIRATORY TRACT INFECTIONS

Treatment and Disposition
Toxic-appearing and otherwise unstable patients—those with airway, breathing, and circulatory compromise—require an aggressive approach, with initial rapid evaluation and stabilization (supplemental oxygen and intravenous fluid boluses and airway interventions, if within the scope of the clinic and clinician) and prompt ambulance transport to the emergency department.

Patients who have compromised immunity, significant comorbidities (e.g., chronic obstructive pulmonary disease, pulmonary fibrosis, cystic fibrosis, congestive heart failure, hepatic and renal disease, etc.), or refractory, persistent, or frequently recurrent URTIs require more complex decision-making than that described here.

In general, however, stable, otherwise healthy patients with uncomplicated URTIs who are maintaining their hydration—who will be the overwhelming majority of patients in most practices—can be treated very simply at home.

In every situation, explain to patients what they should expect and discuss precautions for immediate re-evaluation, as well as specific timing for return if not improving. Schedule next-day follow-up for patients for whom the level of illness is unclear. Because every disease has a time course and even uncommon things will occur the longer one practices, use good communication and close clinical follow-up as your safety net for every patient.

Common cold, bronchitis, and viral pharyngitis
Existing clinical practice guidelines emphasize the importance of not prescribing antibiotics for patients with a common cold, acute bronchitis, and viral pharyngitis. Symptomatic care can include acetaminophen or a non-steroidal anti-inflammatory medication (if there are no contraindications) for fever, aches, and pain. Stronger analgesics may be reasonable in patients who fail to get relief with these, e.g., to facilitate oral fluid intake in those with pharyngitis.

Often, patients desire relief from cough; unfortunately, no preparation has consistently shown clinical benefit. A potential limitation in this research, however, is that comparison is often made with a placebo, yet no placebo exists for clinicians to prescribe or recommend! It is probable that, as long as the possible side effects are considered by the patient and provider, prescription of some sort of cough suppressant is reasonable.

In patient with bronchitis, some studies have shown variable benefit for the use of beta-agonist inhalers, like albuterol, to help with cough and chest congestion; the presence of wheezing on exam may indicate a greater chance of benefit in a particular patient.

Otitis media and acute sinusitis
For both acute otitis media and sinusitis, the decision to treat with antibiotics may be based on available guidelines, plus the knowledge that placebo-controlled studies have shown rates of up to 80% resolution without antibiotics for these conditions.7

A study published in 2007 in the British Medical Journal8 estimated that over 4,000 patients with otitis media would need to be treated with an-
 Antibiotics to prevent a serious complication (e.g., mastoiditis) in one patient; a similar “number needed to treat” of over 4,000 applied to preventing serious complications of URTI and sore throat.

Specifically, antibiotic treatment is recommended for all of those under 6 months of age who have a diagnosis of otitis media. There is an option to observe and withhold antibiotics in children between 6 months and 2-years-old if the diagnosis is uncertain or the condition is not severe, and for those 2 years and older unless the diagnosis is certain and the disease is severe.

When antibiotics are used, focused-spectrum therapy is recommended (see Table 3). Amoxicillin is still first line, although because of the prevalence of drug resistance among pneumococcus, a high-dose regimen (80 mg/kg/day to 90 mg/kg/day, divided BID) is recommended.

For all patients with otitis media, attention to analgesia (oral and topical) is strongly emphasized. Decongestants and antihistamines have not been shown to be helpful.9 There are no specific guidelines for adults with otitis media.

For acute sinusitis, existing guidelines recommend using antibiotics in patients with severe symptoms or moderate symptoms that are worsening after five to 10 days or not improved after 10 days. Again, focused-spectrum antibiotics (Table 3) are first line for uncomplicated infections if antibiotics are felt to be necessary.

Studies have yielded a range of results regarding the use of nasal steroid sprays, and a Cochrane Review10 of the literature found them to be possibly effective. Antihistamines may cause drying of nasal secretions and impede drainage, and are generally avoided in patients with sinusitis.

Interestingly, some new evidence11 suggests that, though a specific patient may indeed have sinusitis, there may be no reliable clinical indicators to tell a clinician whether antibiotics might be helpful or harmful to that patient.

GABHS pharyngitis
Penicillin-resistance among GABHS has been reported to be nonexistent or extremely rare. (In contrast, macrolide resistance is rising.) Concerns have been raised regarding penicillinase activity among other organisms inhabiting the throat at the same time as a GABHS infection, but these seem to impact mostly disease-related outcomes such as culture-proven eradication of the pathogen, rather than patient-oriented outcomes such as duration of illness or the development of complications.

For all of these reasons, guidelines continue to recommend penicillin as first-line antimicrobial therapy for GABHS pharyngitis in patients who are not allergic to it. For penicillin-allergic patients, a narrow-spectrum alternative antibiotic should be used (Table 3).

Adjunctive systemic corticosteroids for one to three days (at most) may help decrease pain associated with GABHS pharyngitis.

Continued on page 28

Table 3. Recommended Antibiotic Regimens for Uncomplicated Respiratory Tract Infections

<table>
<thead>
<tr>
<th>Respiratory tract infection</th>
<th>First-line antimicrobial therapy</th>
<th>Alternate therapy</th>
</tr>
</thead>
<tbody>
<tr>
<td>Common cold</td>
<td>None</td>
<td>None</td>
</tr>
<tr>
<td>Acute sinusitis</td>
<td>None or amoxicillin</td>
<td>Doxycycline</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Trimethoprim/sulphamethoxazole</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Cefdinir, cefprozil, cefuroxime, cefpodoxime</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Amoxicillin/clavulanate</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Respiratory fluoroquinolones</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Clarithromycin</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Azithromycin</td>
</tr>
<tr>
<td>Acute pharyngitis</td>
<td>Penicillin, if Strep testing is positive</td>
<td>For penicillin-allergic* patients:</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Second-generation cephalosporins</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Erythromycin</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Clindamycin</td>
</tr>
<tr>
<td>Acute bronchitis</td>
<td>None</td>
<td>None</td>
</tr>
<tr>
<td>Acute otitis media</td>
<td>Amoxicillin (high-dose) 80-90 mg/kg daily divided BID</td>
<td>Amoxicillin/clavulanate</td>
</tr>
<tr>
<td></td>
<td></td>
<td>For penicillin-allergic* patients:</td>
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<tr>
<td></td>
<td></td>
<td>Cefuroxime</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Cefdinir</td>
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<tr>
<td></td>
<td></td>
<td>Cefpodoxime</td>
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<tr>
<td></td>
<td></td>
<td>Azithromycin</td>
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<tr>
<td></td>
<td></td>
<td>Clarithromycin</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Ceftriaxone</td>
</tr>
</tbody>
</table>

* Some patients who report prior allergy to penicillin also have allergic reactions to cephalosporins; if a person has had anaphylaxis or other severe allergy to penicillin, it is safest to avoid cephalosporins.
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Case Report

A Child with Constipation and Swollen Abdomen

Urgent message: Malignancies in children are often discovered only inadvertently, in conjunction with seemingly less dire presentations. Awareness of relevant signs and symptoms by the urgent care clinician can be invaluable in identifying tumors that might otherwise escape notice until they are at an advanced stage.

Muhammad Waseem, MD

Introduction

The identification of a palpable abdominal mass in an urgent care center or emergency department is quite concerning, as it represent a serious underlying disorder. Any abdominal mass in a child is usually considered malignant until proven otherwise. Abdominal tumors are uncommon in children and can present with pain, vomiting, or abdominal mass.

Here, we report on a child who presented with history of constipation and an abdominal mass that was identified on physical examination and subsequently diagnosed as neuroblastoma.

Case Study

A 15-month-old female presented with a two-day history of constipation. There was no vomiting, fever, or abdominal pain. She previously had been in good health and had seen her pediatrician regularly. Her parent also noted her to have a swollen abdomen, which they attributed to “constipation.” She had been evaluated for this by her pediatrician and was taking stool softeners.

Findings

Physical examination revealed a well-developed, comfortable child with:
- temperature 99°F (36.9°C)
- pulse 108 beats per min
- BP 99/62 mmHg
- respiration 20 per minute.

The patient’s abdomen was mildly distended, with fullness on the left side and flank. A somewhat firm, nontender, nonmobile mass with rounded margins was palpable in the left lower quadrant and left lumbar regions. No guarding, rigidity, or tenderness was noted. Bowel sounds were audible. Rectal examination
Other findings included:
- white blood cell count $9.3 \times 10^3 /\text{mCL}$ (11.1x10^9/L)
- 29.1% neutrophils

- 56.4% lymphocytes
- 12.7% monocytes
- 1.2% eosinophils
- 0.6% basophils
- hemoglobin 11.3 g/dL
- platelet count was $733 \times 10^3 /\text{mCL}$ ($733 \times 10^9$/L)
- serum electrolytes were normal
- aspartate aminotransferase 43 U/L
- alanine aminotransferase 16 U/L
- albumin 3.8 g/dL
- bilirubin 0.3 mg/dL
- alkaline phosphatase 188 U/L
- lactic dehydrogenase 317 U/L

Urinalysis yielded normal findings.

The initial renal ultrasound showed a retroperitoneal mass in the left flank without hydronephrosis or intrinsic renal mass (Figure 1). Subsequently, a contrast computed tomography (CT) scan of the abdomen revealed a left retroperitoneal mass between the left kidney and upper lumbar spine, with some displacement of the left kidney (Figure 2). The urinary catecholamine metabolites, homovanillic acid (HVA), and vanillylmandelic acid (VMA) were also elevated.

The patient was referred to an affiliated tertiary care center for sub-specialty care.

**Discussion**

The child with an abdominal mass presents a unique challenge. A palpable mass in the abdomen of a child is a serious finding, with a fairly extensive differential diagnosis.

Abdominal masses may occur at any age, and may have a wide variety of clinical presentations. It is the urgent care clinician’s role to differentiate between benign conditions such as constipation and more serious causes of abdominal mass (Table 1).

When an abdominal mass is discovered on physical examination, the immediate goal is to determine its nature and extent. A tentative diagnosis can often be made from the presenting symptoms, age of the child, and the site of mass. Physical examination should focus on the location, size, and mobility of the mass, as well as other
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†In vitro data are not always indicative of clinical success or microbiological eradication in a clinical setting.

*Remember to use the full course of therapy—7 days.

IMPORTANT SAFETY INFORMATION
VIGAMOX® solution is indicated for the treatment of bacterial conjunctivitis caused by susceptible strains of the following organisms: Corynebacterium species‡, Micrococcus luteus‡, Staphylococcus aureus, S. epidermidis, S. haemolyticus, S. hominis, S. warneri‡, Streptococcus pneumoniae, Streptococcus viridans group, Acinetobacter Iwoffii‡, Haemophilus influenzae, Haemophilus parainfluenzae‡, Chlamydia trachomatis (efficacy for this organism was studied in fewer than 10 infections). VIGAMOX® solution is contraindicated in patients with a history of hypersensitivity to moxifloxacin, to other fluoroquinolones, or to any of the components in this medication. NOT FOR INJECTION. VIGAMOX® solution should not be injected subconjunctivally, nor should it be introduced directly into the anterior chamber of the eye. In patients receiving systemically administered quinolones, including moxifloxacin, serious and occasionally fatal hypersensitivity (anaphylactic) reactions have been reported, some following the first dose. As with other anti-infectives, prolonged use of VIGAMOX® solution may result in overgrowth of non-susceptible organisms, including fungi. The safety and effectiveness of VIGAMOX® solution in infants below 1 year of age have not been established. The most frequently reported ocular adverse events were conjunctivitis, decreased visual acuity, dry eye, keratitis, ocular discomfort, ocular hyperemia, ocular pain, ocular pruritus, subconjunctival hemorrhage, and tearing. These events occurred in approximately 1%–6% of patients.
**Table 1. Common Causes of Abdominal Mass in Children**

<table>
<thead>
<tr>
<th>Cause</th>
<th>Percentage of Cases</th>
</tr>
</thead>
<tbody>
<tr>
<td>Wilms’ tumor</td>
<td>20%</td>
</tr>
<tr>
<td>Neuroblastoma</td>
<td>10%</td>
</tr>
<tr>
<td>Lymphoma</td>
<td>5%</td>
</tr>
<tr>
<td>Polycystic or dyplastic kidney</td>
<td>2%</td>
</tr>
<tr>
<td>Hydrocephrosis</td>
<td>2%</td>
</tr>
<tr>
<td>Hepatoblastoma</td>
<td>2%</td>
</tr>
<tr>
<td>Teratoma</td>
<td>2%</td>
</tr>
<tr>
<td>Ovarian cyst</td>
<td>2%</td>
</tr>
<tr>
<td>Constipation/fecal mass</td>
<td>2%</td>
</tr>
</tbody>
</table>

**Clinical Presentation**

NB may have diverse clinical features because of its variable sites of origin, propensity to metastasize, and secretion of hormones. Clinical features at presentation depend on the location and primary cause of the tumor, and whether the tumor has abnormalities on examination. Examination should also include measurement of serial blood pressures and a complete neurological assessment.

The presence of an abdominal mass may indicate the existence of colonic stools or tumors. In constipation, a mass can be palpated, which is usually associated with fecal impaction on rectal examination. Wilms’ tumor and neuroblastoma (NB) are the two common intra-abdominal masses in children. These are more prevalent in children younger than 4 years.

NB is the most common extracranial tumor in children, accounting for 10% of pediatric malignancies. After tumors of the central nervous system, it is the most common solid tumor in children. It is also the most common malignant tumor of infancy, with 50% of cases occurring in children younger than 2 years and 75% diagnosed by the fourth year of life.

The median age at diagnosis is 2 years, but the tumor may present in the neonate or even in adolescents and adults. NB occurs most commonly in the abdomen (65%), either in the sympathetic chain or more commonly in the adrenal gland. After the abdomen, the thorax is the second most common location of NB (15%) followed by the neck (1%) and the pelvis (2% to 3%).
metastasized. Between 50% and 70% of patients with NB may have metastasis at the time of their presenta-
tion.7 Large masses may cause respiratory distress. The common sites for distant metastasis include the bone marrow, liver, and skin.8

The signs and symptoms of this tumor are dependent on the location of the primary tumor, which may occur anywhere along the peripheral sympathetic nervous system, and the sites of metastatic disease. Typically the initial symptoms are nonspecific (general malaise, weight loss, unexplained fever), as in most patients the tumor is either retroperitoneal or in posterior mediastinum.9

Common symptoms
The common symptoms are:

- abdominal pain or discomfort
- sensation of fullness
- fever
- weight loss.

Although most children present with abdominal pain or a palpable mass, many present with manifestations of their metastatic disease, including bone or joint pain and periorbital ecchymosis.

Diagnostic Evaluation
An abdominal mass in a child must be considered malignant until proven otherwise. As such, any studies described here but not feasible for the urgent care setting should be facilitated via referral. They are included here for the sake of presenting as complete a picture of the diagnostic process as possible.

When an abdominal mass is detected in a child, imaging studies should be performed. Ultrasound is helpful in the initial evaluation of a child with an abdominal mass, as it is useful in determining the origin of the mass. It will also help determine whether the mass is cystic or solid.

Abdominal CT scan is a commonly used imaging modality for the assessment of a child with abdominal mass. It offers several advantages in the differential diagnosis of a possible Wilms’ tumor, such as confirmation of the intrarenal origin of the tumor, detection of multiple masses, determination of the extent of tumor, and evaluation of the opposite kidney.

Initial laboratory studies should include a complete blood count to identify anemia or thrombocytopenia suggestive of bone marrow invasion. NB produces catecholamine metabolites, which are used as tumor markers; therefore, a search for specific tumor markers secreted by the suspected tumor should also be made.

Elevation of urinary catecholamine metabolites, homovanillic acid (HVA), and vanillylmandelic acid (VMA) is used as a diagnostic screen,10,11 as this is present in 75% to 90% of patients with NB.12

Differential Diagnosis
Most abdominal masses in infants are due to problems of the urinary tract. Hydronephrosis and multicystic kidney are common causes of flank masses at this age.13 If a mass appears to be in the flank in an older infant or child, Wilms’ tumor and NB should be considered.

Additional features suggestive of Wilm’s tumor include fever, abdominal pain, or hematuria.

Hepatoblastoma is the most common primary hepatic tumor in young children. A high serum alpha-fetoprotein is often noted.

Lymphoproliferative conditions can also present with an abdominal mass. A suprapubic mass may be due to a distended bladder, which in turn may be secondary to urinary tract obstruction.

Conclusion
The presence of an abdominal mass in children includes a wide spectrum of diseases. This case emphasizes the value of performing a careful and thorough physical examination upon presentation. The presence of an abdominal mass should generate the suspicion for the possibility of tumor.

References
ABSTRACTS IN URGENT CARE

On Retail Clinics, Bell Palsy, and Abscesses: to Pack or Not to Pack?

NAHUM KOVALSKI, BSc, MDCM

Each month, Dr. Nahum Kovalski reviews a handful of abstracts from, or relevant to, urgent care practices and practitioners. For the full reports, go to the source cited under each title.

Cost and Quality of Care at Retail Clinics

Key point: Quality scores at retail clinics rivaled those at urgent care centers, physician offices, and EDs.


Professional organizations have raised concerns about the quality of care that is delivered at store-based retail clinics. Using claims data from a Minnesota health insurer, investigators searched for episodes of initial care for pharyngitis, otitis media, and urinary tract infections; they identified 2,100 episodes that occurred in retail clinics and matched them with 13,070 episodes that occurred in urgent care centers, physician offices, or emergency departments (EDs). In addition to location and ailment, cases were matched by age, sex, comorbidities, and income.

Aggregate quality scores (proportion that met indicators specific for each condition) were similar for retail clinics, physician offices, and urgent care centers (64%, 61%, and 63%, respectively) and lower at EDs (55%). Urine cultures for high-risk patients were ordered less often at retail clinics than in other settings (30% vs. 55%-58%). Costs per episode (health plan reimbursements plus copayments) were lower for retail clinics ($110) than for physician offices ($166), urgent care centers ($156), and EDs ($570).

Many physicians smirk when they hear about care that is delivered at a retail clinic or at a freestanding “doc-in-the-box.” This study won’t settle questions about quality of care in such settings, in part because all settings scored low and measures were obtained exclusively from claims data. But, clearly, these services have arisen in response to perceived needs.

[Published in J Watch Gen Med, September 17, 2009—Richard Saitz, MD, MPH, FACP, FASAM.]

Do Corticosteroids and Antiviral Agents Have Benefits for Patients with Bell Palsy?

Key point: New evidence has emerged regarding the use of corticosteroids and antiviral agents in Bell palsy.


Eligible studies were randomized controlled trials comparing treatment with either corticosteroids or antiviral agents with a control and measuring at least one of the following outcomes: unsatisfactory facial recovery (four months), unsatisfactory short-term recovery (six week to less than four months), synkinesis and autonomic dysfunction, or adverse effects. Eighteen trials involving 2,786 patients were eligible. Regression analysis identified a synergistic effect when

Nahum Kovalski

Nahum Kovalski is an urgent care practitioner and assistant medical director/CIO at Terem Emergency Medical Centers in Jerusalem, Israel.
corticosteroids and antiviral agents were administered in combination compared with alone.

Meta-analysis using a random-effects model showed corticosteroids alone were associated with:

- a reduced risk of unsatisfactory recovery
- a reduced risk of synkinesis and autonomic dysfunction
- no increase in adverse effects.

In addition:

- Antiviral agents alone were not associated with a reduced risk of unsatisfactory recovery.
- When combined with antiviral agents, corticosteroids were associated with greater benefit than antiviral agents alone.
- When combined with corticosteroids, antiviral agents were associated with greater risk reduction of borderline significance compared with corticosteroids alone.

Simple Abscesses—Can We Poke without the Pack?

Key point: Packing is not necessary.


Visits to the emergency department for cutaneous abscesses more than doubled between 1996 and 2005. Incision and drainage are the mainstay of treatment. The authors of these articles challenge the common wisdom that packing is critical to the care and healing of cutaneous abscesses.

Researchers conducted a prospective, randomized, single-blinded study of 48 adult subjects presenting to a single emergency department with simple cutaneous abscesses. All adult patients with cutaneous abscesses on the trunk or extremities that required I&D were eligible. Exclusion criteria included abscesses greater than 5 cm in diameter, pregnancy, comorbid medical conditions with possible immunosuppression including diabetes, HIV, malignancy, and chronic steroid use, as well as abscesses on other areas of the body and head, and a sulfa allergy.

Standard incision, drainage, and irrigation were performed on each abscess. Subjects were then randomized to packing with ¼ inch plain gauze or no packing. Pain scores were measured in the ED using a standard 100-point visual analog scale (VAS); subjects were asked to record their VAS pain score twice daily until they were seen in a return visit in 48 hours. All subjects were prescribed trimethoprim-sulfamethoxazole (TMP-SMX), ibuprofen, and oxycodone/acetaminophen.

A physician blinded to the study examined each wound at
ABSTRACTS IN URGENT CARE

the 48-hour visit and determined the need for further intervention and measured wound erythema, induration, and fluctuance. Measurements were repeated by a second, similarly blinded physician. All patients were contacted by phone 10-15 days after the initial visit to determine if their abscesses had required additional interventions.

Patients were randomized to the packing group (n=23) or the non-packed group (n=25). Only 34 subjects (66%) returned for the 48-hour follow-up visit. Thirteen were from the non-packed group and 21 from the packed group. Four of the patients in the packed group and five of the patients in the non-packed group required intervention at follow-up.

Ten of the 11 patients in the non-packed group who did not return for follow-up and were contacted by phone reported that they did not think the abscess required re-evaluation and that they were pain free. Only one of the three patients in the packed group who did not follow up was reached and reported moderate pain but did not return to the ED.

There was no difference between the groups in pre-procedural pain scores. Subjects in the packed group reported higher pain scores in both the immediate post-procedural period and at the 48-hour follow up visit. There was no significant difference in the amount of ibuprofen taken, but patients in the packing group took a mean of 3.1 narcotic pain pills, compared with a mean of 0.91 pills in the non-packed group.

Given the prevalence of community acquired methicillin-resistant Staphylococcus aureus, it is unlikely that we will see a reduction in the prevalence of cutaneous abscesses. However, if the evidence bears out, elimination of packing of simple abscesses will save time and money and reduce patient discomfort.

URTI, continued from page 18

A note on dehydration

A complication of any of the URTIs in children, and sometimes adults, is dehydration. Fever and other mechanisms can increase insensible fluid loss, and malaise and sore throat can decrease fluid intake. Discuss fever control, analgesia, and appropriate oral hydration with each patient; occasionally providing intravenous fluid rehydration may be necessary.

Patient Satisfaction

There is no evidence that patient satisfaction is related to getting an antibiotic prescription for a URTI. In addition, data show that clinicians are not able to determine whether any particular patient expects such a prescription or not.

Studies do link patients’ satisfaction to their receiving discussions of their diagnoses, as well as attention to alleviation of their symptoms.

Several years ago, the concept of a delayed or “safety net” prescription was introduced. This strategy involved giving a patient an antibiotic prescription, along with instructions to wait for several days of no improvement before filling and beginning to take it. This approach was shown in several studies to be safe, to reduce antibiotic use, and to be satisfactory to patients. However, a recent review combining many studies showed that prescribing no antibiotic, rather than giving a safety net prescription, resulted in similar clinical and patient satisfaction outcomes, assuming clinicians felt that it was safe not to prescribe antibiotics for a URTI.

Conclusion

Antibiotic prescribing has a direct impact on the development of antimicrobial resistance. URTIs are a common chief complaint in urgent care practice, and the tendency to overprescribe antibiotics exists. A variety of guidelines and data from the medical literature can assure the clinician that antibiotics are not necessary for the majority of uncomplicated URTIs in most patients.

References

In each issue, JUCM will challenge your diagnostic acumen with a glimpse of x-rays, electrocardiograms, and photographs of dermatologic conditions that real urgent care patients have presented with.
If you would like to submit a case for consideration, please e-mail the relevant materials and presenting information to editor@jucm.com.

The patient is a 19-year-old male who complains of pain after receiving a blow to the shoulder.
Range of motion is limited due to pain. The patient is otherwise healthy.
View the x-ray taken (Figure 1) and consider what your diagnosis and next steps would be.
Resolution of the case is described on the next page.
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The correct reading of the image is pseudosubluxation of the shoulder, which refers to inferior displacement of the humeral head relative to the glenoid by fluid in the joint cavity, either hemarthrosis or lipohemarthrosis. This is not a true dislocation.

The radiograph demonstrates a crescent-shaped, low density region representing fat layering above blood within the joint space. The humeral head is inferiorly displaced—but not dislocated—due to distension of the joint.

No fracture line was seen on this film.

Early follow-up with an orthopedist is important. Immediate management is a sling and ice.

Acknowledgment: Case presented by Nahum Kovalski, BSc, MDCM, Terem Emergency Medical Centers, Jerusalem, Israel.
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## 2009/10 UCAOA PRACTICE MANAGEMENT WEBINAR SERIES SCHEDULE

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                        | Patty Riskind, Patient Impact                                      |
| November 5th at 1 p.m. (cst) | Considerations for Expanding or Relocating a Center  
                        | Michael Zelnick, Equity Partners                                   |
| December 3rd at 12 p.m. (cst) | Evaluating Media Purchases – TV, Newspaper, Radio  
                        | Ira Bloomfield, Market Welby Urgent Care                          |
| January 21st at 1 p.m. (cst) | How to Motivate Urgent Care Staff to Deliver Exceptional Service  
                        | Marty Martin, PhD, DePaul University                               |
| February 18th at 1 p.m. (cst) | Top Ten Urgent Care Coding Mistakes: Impact on Compliance and Revenue  
                        | Dr. David Stern, Practice Velocity                                |
| March 4th at 1 p.m. (cst) | Integrating Urgent and Primary Care: Billing/Service issues; Differentiating Models  
                        | Jennifer Stephenson, PrimaCare Medical Centers                     |

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I just returned from Boston, where the American College of Emergency Physicians held its national conference. While there, along with learning (and relearning) some emergency medicine, I had the chance to walk along the Freedom Trail and enhance my understanding of our battle for independence. What continually amazes me is how fortunate we were to actually succeed. Many times the only thing which turned the tide and saved the day was the persistence of our founders.

Many historians consider the Siege of Boston (starting after the battles of Lexington and Concord) to be the beginning of the Revolutionary War. During the siege, militiamen surrounded Boston, attempting to prevent the British Army, which was garrisoned within Boston, from receiving supplies. To fortify his army’s position and prevent the British Navy from supplying the British Army, General Washington sent a 25-year-old bookseller named Henry Knox to bring heavy cannons that had been captured at Fort Ticonderoga in New York all the way to Dorchester Heights, MA, which overlooked Boston’s harbor. Over a wet and freezing winter, Knox and his small group moved 60 tons of artillery by boat, horse-drawn sledges (which they built), and sheer persistence 300 miles along snow-packed trails, across two semi-frozen rivers, and through forests and swamps to the Boston area in 56 days.

Historian Victor Brooks called Knox’s feat “one of the most stupendous feats of logistics” of the entire war. Ultimately, the effectiveness of these cannons marked the turning point which eventually forced the British out of Boston.

One of my favorite quotes comes from Calvin Coolidge: “Nothing in this world can take the place of persistence. Talent will not; nothing is more common than unsuccessful people with talent. Genius will not; unrewarded genius is almost a proverb. Education will not; the world is full of educated derelicts. Persistence and determination alone are omnipotent. The slogan ‘press on’ has solved and always will solve the problems of the human race.”

You may be asking yourself, how is this relevant to urgent care medicine? More and more I am realizing the “secret sauce” which separates the failing (or, at best, marginally successful) business from the highly profitable endeavor is simply the persistence of the leadership team.

Urgent care medicine is engaged in our own Siege of Boston. There are a number of unknowns which will influence the eventual outcome of our new specialty. Retail clinics will eventually figure out how to turn a profit for their investors. When this happens, how much more of our current patient volume will be usurped by these groups? How will healthcare reform impact our business?

It seems logical that under any plan to reduce healthcare costs, urgent care should fair well. However, will we survive until that happens? How will our rates be affected? We are already at the low end of the reimbursement continuum and even trivial rate cuts could erase our already thin margins.

When will the malpractice insurers realize that urgent care medicine is a risky proposition and raise our premiums? When will emergency departments improve their throughput and lower their cost to compete with urgent care centers and retail clinics?

So Mr. Knox, what’s the game plan? How are we going to move the cannons and protect our already besieged turf? Here are some strategies to consider:

- **Don’t settle.** Health plans are in business to make a profit for their shareholders. To accomplish this, they have three options:
  1. Charge higher rates.
  2. Reduce utilization.
  3. Pay providers less.

Out of those three, guess which one is the easiest? You guessed it, pit the providers against each other, and contract with the urgent care who will accept the least amount of reimbursement. Ever hear of divide and conquer? It is happening to us. How do we combat it without being accused of price fixing? Educate the plans on the value of contracting with urgent care centers and hold the line on reimbursement.

**John Shufeldt** is the founder of the Shufeldt Law Firm, as well as the chief executive officer of NextCare, Inc., and sits on the Editorial Board of *JUCM*. He may be contacted at jsf@shufeldtlaw.com.
HEALTH LAW

- **Practice quality medicine.** What influences our malpractice rates? The collective industry malpractice experience. Therefore, if our industry has a disproportionate amount of adverse events, it will affect all of our rates, no matter our own center’s malpractice history.

  I have the “benefit” of reviewing a large number of malpractice cases brought against urgent care centers all across the U.S. The vast majority are very preventable by following a few simple ideas: Urgent care centers (like emergency departments) typically have “one-off” encounters where the tolerance for error is low. Therefore, protect yourself—and most importantly, the patient—by ruling out the things that will kill them.

  I advocate using a liberal number of diagnostic tests. No one ever died from getting an unnecessary EKG, CXR, D-dimer, troponin, etc. We typically have one shot to get it right, so do what is necessary to assure yourself that the patient does not have an unusual presentation of a deadly condition.

- **Use standing orders.** Important tasks and tests must not be overlooked during high volume times. Every malpractice case I have reviewed could have been easily prevented by using rational standing orders.

- **Use informed consent and document the conversation.** Engage patients in their own healthcare. Appropriately documented informed consent is medicine’s equivalent to Monopoly’s “Get out of Jail Free” card.

- **Invest in a state-of-the-art electronic health records system.** This will allow the provider to accurately record the patient’s treatment, discharge instructions, and informed consent. Let the patient take home an electronic or paper copy of the record so that they can share it with their personal physicians.

- **Stick together.** Don’t denigrate your competitors despite what they may be saying. There will be enough people criticizing urgent care centers without us trash-talking our own. As Ben Franklin said, “We must hang together, gentlemen...else, we shall most assuredly hang separately.”

As a discipline, if we simply hold the line on reimbursement, practice quality medicine, and stick together we will enjoy the longevity that our specialty and our patient’s deserve.

In other words, we have to persist and thrive in the face of a myriad of challenges.

So what happened to Henry Knox? He was repeatedly promoted and was eventually named the first Secretary of War. He went on to champion rights for Native Americans and later retired to Maine. Most importantly, his extraordinary accomplishments paved the way for our eventual independence.
**Q.** We bill S9083 to several carriers. Occasionally, a patient will have secondary insurance. If the primary insurance is contracted to pay the S9083 code but transfers the balance to the deductible, how do we bill the secondary carrier if they do not accept the code?

- **Question submitted by Paula Seify, Back Office MD**

**A.** Many secondary payors do not accept S9083, but these payors still will often cover the actual services that were rendered under this code if you bill them using typical fee-for-service codes.

To my knowledge, there is no official CMS or AMA guideline for appropriate coding to a secondary payor under this situation. Most coders would suggest that you recode the services using standard coding methods, i.e., E/M, CPT, and HCPCS codes as appropriate.

In order to avoid compliance issues, the total dollars billed for the aggregate of these codes, however, should not exceed the initial amount billed to the primary insurance under the S9083 code.

---

**Q.** A 25-year-old new patient presented to our urgent care center with two lacerations—a 2 cm laceration on the face and a 2.2 cm on the leg. He was otherwise healthy. Can I code an E/M code since I spent 45 minutes for both suture repairs, then add 12011 for the facial laceration repair and 12001 for the leg with modifier -51, as there are more than one laceration? I know that we can add the two cuts into one and use 12004, but I wonder if billing this way you would get less reimbursement since there are two different sites.

- **Physician, Name withheld, California**

**A.** Let me answer in several parts. First, the time spent repairing the laceration should not count toward an E/M code, as this time is already accounted for in the code(s) for the laceration repair.

Second, is it appropriate to code an E/M code? You say that he was “otherwise healthy,” so I assume that you took an appropriate history and performed an appropriate physical exam. Is this information documented and separately identifiable in the chart? If so, then you may code for the E/M that is appropriate for this documentation and medical complexity. You need to add modifier -25 to the E/M code to indicate that the E/M was performed in addition to the procedure(s).

Third, the coding method that gets more reimbursement should not determine code selection. Instead, use the compliant method per CPT, CMS, or other applicable payor.

Fourth, if the laceration repairs are of the same complexity and if the laceration repairs are located in anatomic regions that are grouped together under the same general heading in the code descriptors for that complexity of laceration repair, then the coder should add the lengths of the two lacerations together to determine the appropriate CPT code. In your example, however, the anatomic locations (face and left leg) are not grouped under the same general heading for simple laceration repairs. Thus, you should not add the lengths of the lacerations together. Instead, you should code each laceration separately, with 12001 (Simple repair of superficial wounds of scalp, neck, axillae, external genitalia, trunk, and/or extremities (including hands and feet), 2.5 cm or less) and 12011 (Simple repair of small wound of facial area, 2.5 cm or less).

Remember to measure the lacerations after the repair is completed.

---

**Q.** We had a patient who needed intravenous fluids and monitoring for five hours. We found the CPT codes 96360-96361 to use for the intravenous therapy. However, the doctor cannot believe the paltry reimbursement for these codes is correct. We did bill an office visit...
Dealing with prospective client companies such as Walmart that are theoretically wed to a national contract is a relatively common obstacle that may seem virtually impossible to overcome.

Do not, however, view such contracts as impenetrable walls; they can be dealt with in many ways.

**All Contracts are Not Created Equal**

When an employer cites a contractual obligation, much may be left unsaid. Indeed, contracts come in various sizes and shapes. A contract can be:

- local, regional, or national
- prospect-specific (relevant only to that company in that locale) or part of a contract that affects an entire national chain (e.g., Home Depot)
- limited in scope or all-inclusive
- binding to the prospect company or discretionary
- simply a price discount that can be matched or sold over by another provider.

Given these variants, there are many ways that an urgent care clinic can approach a contractually bound prospect. You must first, however, ferret out the nature of the contract or perceived obligation. You should pose questions such as:

- Is the contract for a specific time period?
- Do you or your local office have unilateral authority in arriving at a contractual relationship? If not, who is responsible for a final decision?
- Exactly what services does the contract cover?
- What other types of occupational health-related services does your company use that are not specifically bound to the contract?
- To what degree is the contract binding to your company?
- In what way, if any, might another provider bid for your company’s business?
- (If the contract is merely a guaranteed price discount): Besides price, what other things does your company consider when valuating the return on your investment for health and safety support?

In sum, probe to get the facts, look for openings when they occur, and learn to separate opportunities from closed doors.

**Remember: You Have Other Things to Sell**

Your urgent care center’s occupational health strategy should include product expansion, rather than simply maintaining a static, narrow program. A portfolio-expanding strategy also provides your clinic with a series of fallbacks if and when you are unable to overcome a contractually bound objection.

**Existing Contracts Can Mean Opportunity**

Assume for a moment that a national player makes even greater inroads with national contracts and that many of your best prospects become wed to such contracts. Such a theoretical crisis should be viewed as an opportunity.

As provision of core occupational health services becomes less cost effective and as national obligations potentially erode relationships with your clients or prime prospects, it becomes necessary to develop and offer supplemental services (specialized screenings, wellness services, executive health, travel medicine, etc.).

**You Don’t Have to Bat 1.000**

At .365, the Minnesota Twins’ Joe Mauer had the highest batting average in the major leagues in 2009. Accolades aside, Mr. Mauer’s .365 batting average also indicates that he...
"Outright resignation negates opportunity that may be a question or two away."

failed to get a hit 63.5% of the time. So it is with a sales professional in almost any industry. You have to anticipate a mix of home runs, base hits, walks, and strikeouts.

A company’s national contract obligation may in fact close the door on your clinic—at least for the time being. It is inevitable and part of your sales management function. However, remember to ask questions and probe to seek opportunities when dealing with such companies. Nowadays, outright resignation seems to be the norm, thus negating opportunities that may be only a question or two away.

National contracts suggest that occupational health services are a commodity, when in fact they should be seen in terms of a relationship. Armed with such an understanding, the national contract roadblock should be viewed as less onerous than it is in many cases.

The accompanying table presents a series of appropriate responses to common contract scenarios.

<table>
<thead>
<tr>
<th>Condition</th>
<th>Strategy</th>
</tr>
</thead>
<tbody>
<tr>
<td>Long-term contract</td>
<td>Probe for other service needs</td>
</tr>
<tr>
<td>Short-term contract</td>
<td>Position to bid for business at conclusion of contract</td>
</tr>
<tr>
<td>Contract with actual company</td>
<td>Stay in touch with decision maker</td>
</tr>
<tr>
<td>Contract with national office</td>
<td>Identify contact and correspond with national office</td>
</tr>
<tr>
<td>Meeting with decision maker</td>
<td>Learn hot buttons; stay in touch</td>
</tr>
<tr>
<td>Decision maker in home office</td>
<td>Identify contact and correspond with national office</td>
</tr>
<tr>
<td>Contract covers limited scope of services</td>
<td>Tout value of your broad, integrated services</td>
</tr>
<tr>
<td>All, or nearly all, exclusive contract</td>
<td>Look for missing pieces</td>
</tr>
<tr>
<td>Contract compliance is mandatory</td>
<td>Look for missing pieces</td>
</tr>
<tr>
<td>Contract compliance is discretionary</td>
<td>Treat as traditional prospect; emphasize ROI</td>
</tr>
<tr>
<td>Contract involves price discount</td>
<td>Emphasize ROI issues</td>
</tr>
<tr>
<td>Contract involves vague obligation</td>
<td>Treat as traditional prospect</td>
</tr>
</tbody>
</table>

in addition to the intravenous hydration. Is this all we can bill? Does this seem right to you?

- Question submitted by Nicole, First Health Medical, Fresno, CA

A. You are using the correct codes. Don’t forget, however, to list the 96361 multiple times (once for each additional hour after the first hour) when appropriate. If the visit in question is properly documented, for example, you would code an E/M code (e.g., 99203). 96360, 96361 x 4.

Q. Is it appropriate to add modifier -59 to after-hour codes?

- Question submitted by Sharon Dear

A. Using modifiers on these codes is not helpful for compliance or reimbursement. Modifier -59 is for pointing out to a payor that you are referring to a service that might otherwise be bundled into another code, but because of special circumstances, they are really distinct. For the NCCI edits, the primary purpose of modifier “-59” is to indicate that two or more procedures are performed at different anatomic sites or during different patient encounters.

Q. Can 94760 (Non-invasive ear or pulse oximetry for oxygen saturation; single determination) be reimbursed in addition to an E/M code?

- Question submitted by Linda, Keith & Co., El Cajon, CA

A. The code 94760 should be used only when the physician orders a single measurement of oxygen saturation (O2sat) level. Do not use this code when the clinic is documenting pulse oximetry as a routine part of patient intake.

Some basic guidelines for coding 94760 (per Medicare) include:

1. It is only covered if the patient exhibits any signs or symptoms that may be suggestive of oxygen desaturation.
2. A physician order for the pulse oximetry must be documented in the medical record.
3. When pulse oximetry for oxygen saturation is utilized to monitor a patient’s respiratory status, oxygen saturation (during a surgical procedure or conscious sedation) oximetry is considered included in the primary service and not separately reimbursable.

Many, but not all, payors follow similar guidelines. If you are following these guidelines but a payor is denying payment for this code, you should consider an appeal. As always, however, any individual payor may have a policy to deny payment for any particular service.

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Disclaimer: JUCM and the author provide this information for educational purposes only. The reader should not make any application of this information without consulting with the particular payors in question and/or obtaining appropriate legal advice.
URGENT CARE OPPORTUNITY - Seeking BC/BE primary care physician for urgent care in Bryan/College Station Texas. 8 hour shifts, flexible schedule, paid malpractice. For more information contact Lauren with Emergency Service Partners at 888-800-8237, or Lauren@eddocs.com.

URGENT CARE - Seeking primary care physician for hospital-based urgent care near Tyler, Texas. 12 hour shifts, flexible schedule, paid malpractice. For more info contact Julianne with Emergency Service Partners at 888-800-8237, or Julianne@eddocs.com.

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Carolina's HealthCare System is the largest health care system in the Carolinas and operates one of the most successful urgent care networks in the southeast. Our facilities are located in the Charlotte, NC metro area which was ranked in 2008 as the No. 1 city in which to live by Relocate-America.com. Charlotte is conveniently located between the Blue Ridge Mountains and the beautiful Carolina coast!

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Sarah.foster@carolinashealthcare.org or call: 800-847-5084
Fax: 704-355-5033

NEVADA - Laughlin Urgent Care EPMG is seeking a part-time urgent care physician for our clinic in Laughlin, NV or FP with related experience considered. Emergency Physicians’ Medical Group (EPMG) has been providing outstanding partnership opportunity since 1973. EPMG offers democratic governance, open books, and excellent compensation. Contact Bernhard Beltran directly at: 909-509-3073, or 800-828-0898. Email: bbeltran@epmg.com, fax: 330-491-4077, or send CV to: EPMG, 4535 Dressler Road NW, Canton, OH 44718.

Dunkirk and Solomons, Maryland

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Excellent benefits, competitive salary, J-1 Visa, NHSC and Loan Repayment opportunities. Located on the western shore of Lake Michigan, Milwaukee provides Old World charm with world-class arts, cultural and sporting activities, easy access to natural resources and a low cost-of-living environment. 80 miles north of Chicago.

Email interest to gail.paschall@sschc.org, or via U.S. Mail to: Gail Paschall, Sixteenth Street Community Health Center, 1032 S. Cesar E. Chavez Dr., Milwaukee, WI 53204

URGENT CARE OPPORTUNITY – STOCKTON, CALIFORNIA

Gould Medical Group, Inc., California’s premier multispecialty group, is currently seeking two BC/BE emergency, family medicine, or internist physicians to staff their new urgent care department, which will be housed in a brand new 130,000 square foot office building scheduled to open in November of 2009. Candidates should have a full range of urgent care skills, be ACLS certified, and have an interest in working with an innovative group.

Excellent work environment includes:
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Medical Officer
As a Medical Officer, you will be required to provide care to patients presenting at the clinic with a range of urgent care and minor injury presentations. We are looking for doctors with significant GP experience who also have experience in Emergency Medicine and urgent care and are confident of working in an independent setting.

Senior Medical Officer
As the Senior Medical Officer, you will be required to assume a clinical leadership responsibility and to ensure the clinic operates to the highest clinical quality standards. You will be required to have the equivalent experience as the Medical Officer.

Group Medical Officer
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For advertising information contact Trish O’ Brien at: (800) 237-9851, ext. 237 or email jucm@rja-ads.com.
In early 2008, UCAOA revamped its annual survey in conjunction with researchers at Massachusetts General Hospital and Harvard University with the goal of assuring that the UCAOA Benchmarking Committee’s efforts produced a scientifically valid report.

Here, we present some of the data from this landmark survey, to which 436 urgent care centers responded.

In this issue: Which payors foot the biggest portion of the bill among responding urgent care centers?

The results are not dramatically different from the last time the question was asked on a UCAOA benchmarking survey (2006); slightly higher percentages are coming from Medicaid and Medicare, a slightly lower percentage of patients are paying out-of-pocket, and private insurance is covering exactly the same proportion, according to the current report.

The current UCAOA survey report also offered comparison data with primary care and emergency medicine. The portion of payments coming from the occupational medicine segment was dramatically higher in urgent care than in either primary or emergency care, while Medicaid paid a much higher percentage in emergency care than in either urgent care or primary care. Other categories saw less variation among the three settings.

Acknowledgment: Data submitted by Robin M. Weinick, PhD, at the time of the survey assistant professor, Harvard Medical School and senior scientist, Institute for Health Policy, Massachusetts General Hospital. Dr. Weinick is also a member of the JUCM Advisory Board. Financial support for this study was provided by UCAOA.

If you are aware of new data that you’ve found useful in your practice, let us know via e-mail to editor@jucm.com. We’ll share your discovery with your colleagues in an upcoming issue of JUCM.
The Urgent Care Association of America® congratulates the following centers who were recently presented their Certified Urgent Care designation.

- Acadiana Urgent Care Center
  - LaFayette, LA
- AcuteCare of Elizabethtown, PLLC
  - Elizabethtown, KY
- Advocate Condell Immediate Care
  - Buffalo Grove, Gurnee, Round Lake, Vernon Hills, IL
- Armistice Urgent Care and Occupational Health
  - Pawtucket, RI
- Bee Caves Urgent Care
  - Austin, TX
- Covenant Clinic
  - Las Cruces, NM
- ER QuickCare
  - Naples, FL
- Henry Ford Macomb Urgent Care
  - Romeo, Chesterfield, Clinton, MI
- Immediate Care of the South
  - Mobile, AL
- Lake Health
  - Chardon, Madison, Mentor, Willowick, OH
- MD Urgent Care
  - Albuquerque, NM
- MedCare Express
  - North Charleston, SC
- Newport Urgent Care
  - Newport Beach, CA
- OnCall Medical Services
  - Troy, NY
- OnCall Urgent Care
  - Santa Fe, NM
- Physicians Immediate Care
  - East Port St. Lucie, Fort Pierce, Port St. Lucie, FL
- Physicians Immediate Care and Medical Center
  - Richland, WA
- Premier Urgent Care Center, LLC
  - Pompano Beach, FL
- Provena Immediate Care
  - Plainfield, IL
- Ross Urgent Care Plus
  - Hamilton, OH
- San Juan Regional Medical Center
  - Farmington, NM
- Staten Island Physicians Practice
  - Staten Island, NY
- Urgent Care of Green Country
  - Bixby, Claremore, Owasso, Pryor, OK
- Urgent Care Santa Fe
  - Santa Fe, NM
- Valley FirstCare
  - Espanola, NM
- Valley Immediate Care, LLC
  - Grants Pass, Medford, OR
- Wichita Clinic Immediate Care – East
  - Wichita, KS
- Willow Urgent Care and Diagnostic Center
  - Fresno, CA

For more information on how to become a Certified Urgent Care, visit www.ucaoa.org.
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- web demo of our EMR

David Stern, MD, CPC • Practice Velocity

* Map: Numbers per state are accurate, but specific locations not designated.