The Zohydro ER Affair
The conspiracy theory behind the drug approval

Just Say No to Zohydro ER
The abuse potential of this new hydrocodone product makes it a possible contributor to the opioid abuse epidemic

The conspiracy theory behind the drug approval

Just Say No to Zohydro ER
The abuse potential of this new hydrocodone product makes it a possible contributor to the opioid abuse epidemic

During his four years of residency at the University of Southern California Los Angeles County General Hospital, Ryan McGarry, MD, recorded more than 500 hours of emergency department footage. Now, one year out of residency and an assistant professor of emergency medicine at New York-Presbyterian/Weill Cornell Medical College, Dr. McGarry is about to release the 82-minute documentary those 500 hours produced: Code Black. The documentary is named after the highest-level code

CONTINUED on page 18

Even from behind the scenes, these four professionals have made significant contributions to EM
You may know by now that EMP places a high value on having fun and living life wholeheartedly. But our mission: To care for patients, is where we began 22 years ago, and it’s where we begin each day. At EMP, we’re all in. Putting our hearts into everything we do means every patient receives the best care imaginable, and every EMP physician has the opportunity to thrive in a group where fun and discovery never end. Are you all in?
### ACEP Council Speaks Out

ACEP Councillors were asked whether or not they feel certain topics should be addressed at the Council meeting in Chicago this October. Here are the results:

<table>
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<tr>
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<td>51.9%</td>
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<td>5. Safe harbors for tort reform</td>
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<tr>
<td>6. Legalization or decriminalization of marijuana</td>
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<td>9. ED opioid prescribing</td>
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<td>15.2%</td>
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<tr>
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<td><strong>NO</strong></td>
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<tr>
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<tr>
<td>18. CMS 3-day stay rule</td>
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**STRONG OPINION**

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**Strongly Opposed**

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<td>4. Psychiatric patient boarding</td>
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<td>5. Safe harbors for tort reform</td>
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**Opinion**

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**No**

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Dr. Howard Mell Named ACEP’s 2014 Spokesperson of the Year

For his work with national media outlets, including extensive interviews conducted after the 2014 Report Card on Emergency Medicine release, Howard Mell, MD, MPH, FACEP, has been named ACEP’s Spokesperson of the Year. He received his award at last month’s ACEP Leadership and Advocacy Conference in Washington, DC.

Dr. Mell is an emergency physician in the Cleveland, Ohio area and an outstanding member of ACEP’s Spokespersons’ Network. In the past year, he was quoted multiple times in news organizations such as The Chicago Tribune, New York’s Daily News, San Francisco Chronicle, Minneapolis’ Star Tribune, ABC News, and several more on a variety of emergency medicine topics.

Following the publication of a series of articles on energy drinks, attorneys with the beverage industry threatened Dr. Mell with legal action if he didn’t retract his quotes. Dr. Mell stood by his medical experience and did not respond, and the threats never materialized.

As a Report Card spokesperson, Dr. Mell was quoted in multiple media sources, including The Columbus Dispatch, WTWN in Ohio, and CBS Radio in Las Vegas. He also is very active with social media, helping to promote ACEP through Twitter and the Public Relations Committee’s Tweet Team. He is also a member of the sub-group of the Public Relations Committee that evaluates. "Annals of Emergency Medicine studies for promotion to media.

Dr. Michael Miller is 2014 Emergency Department Director of the Year

For his successful collaborative work at the University of Iowa Hospital and Clinics and 18-plus years of commitment to improving patient care, Michael Miller, MD, FACEP, was named 2014 Emergency Department Director of the Year at a May 21, 2014, ceremony in Dallas. Presented by Blue Jay Consulting and the Emergency Medicine Foundation, the award is given to an ED director who demonstrates collaborative relationships with nursing and ancillary departments to implement and improve operational and clinical standards based on evidence-based practice.

Dr. Miller captured the fifth annual award because of several initiatives he has championed, including his strategies to improve door-to-balloon times for ST segment elevation myocardial infarction (STEMI), elevated sepsis recognition with an electronic nursing and ancillary departments to implement, and the Emergency Medicine Conference in Dallas. Presented by Blue Jay Consulting and the Emergency Medicine Foundation.

In the past six months, the emergency department’s “patients left without being seen” percentage has dropped from 3 percent to 0.2 percent, and patient satisfaction survey results have increased from the 30th percentile among similar hospitals to the 84th. The length of stay for all emergency department visitors has decreased from 3.5 hours to 3.0 in the same time frame.

Dr. Miller is the chief safety officer for the health system and one of the hospital’s associate chief medical officers. He also directs the administrative curriculum for EM residents, teaching them how to function as physician administrators in their own emergency departments.

“Lasting change in a complex health care system requires building relationships and teamwork across historic silos to achieve success,” Dr. Miller said. “It is important to maintain optimism in the power of creating a learning environment.”

Dr. Ryan Stanton Recognized by 911 Network

The 911 Legislative Network Member of the Year Award for 2014 goes to Kentucky emergency physician Ryan Stanton, MD, FACEP. This honor is bestowed upon 911 Network members who have gone the extra mile in advocating for the specialty of emergency medicine to federal legislators and

NOW APPROVED FOR MULTIPLE INDICATIONS

Now can be used for:
- Reduction of the risk of stroke and systemic embolism in patients with non-valvular atrial fibrillation
- For the treatment of deep venous thrombosis and pulmonary embolism in patients who have been treated with a potent anticoagulant for 5-10 days
- To reduce the risk of recurrence of deep venous thrombosis and pulmonary embolism in patients who have been previously treated

**Important Safety Information About Pradaxa**

**Indications and Usage**
- Pradaxa® (dabigatran etexilate mesylate) capsules are indicated:
  - To reduce the risk of stroke and systemic embolism in patients with non-valvular atrial fibrillation
  - For the treatment of deep vein thrombosis and pulmonary embolism in patients who have been treated with a potent anticoagulant for 5-10 days
  - To reduce the risk of recurrence of deep vein thrombosis and pulmonary embolism in patients who have been previously treated

**WARNING:** (A) Premature Discontinuation of Pradaxa Increases the Risk of Thrombotic Events. (B) Spinal/Epidural Hematoma

(A) Premature Discontinuation of Pradaxa Increases the Risk of Thrombotic Events

Premature discontinuation of any oral anticoagulant including Pradaxa increases the risk of thrombotic events. If anticoagulation with Pradaxa is discontinued for a reason other than pathological bleeding or completion of a course of therapy consider coverage with another anticoagulant.

(B) Spinal/Epidural Hematoma

Spinal or epidural hematoma may occur in patients treated with Pradaxa who are receiving neuraxial anesthesia or undergoing spinal puncture. These hematoma may result in long-term or permanent paralysis. Consider these risks when scheduling patients for spinal procedures. Factors that can increase the risk of developing spinal or epidural hematoma in these patients include:

- Use of indwelling epidural catheters
- Concomitant use of other drugs that affect hemostasis, such as non-steroidal anti-inflammatory drugs (NSAIDs), platelet inhibitors, other anticoagulants
- A history of traumatic or repeated epidural or spinal punctures
- A history of spinal deformity or spinal surgery
- Optimal timing between the administration of Pradaxa and neuraxial procedures is not known

Monitor patients frequently for signs and symptoms of neurological impairment. If neurological compromise is noted, urgent treatment is necessary. Consider the benefits and risks before neuraxial intervention in patients who are or will be anticoagulated.

Please see boxed WARNING and accompanying brief summary of full Prescribing Information.

NVAF = non-valvular atrial fibrillation; DVT = deep venous thrombosis; PE = pulmonary embolism.
Mr. Tierney

Mr. Gawande

Mr. Gawande received this year’s Award of Excellence in the magazine category for “Why Boston Hospitals Were Ready.” His piece shed light on the value of emergency medicine as it described what happened following the bombings at the 2013 Boston Marathon.

Have an Idea or Suggestion?
Submit a Council Resolution

The deadline to submit a resolution to the Council is July 28, 2014. Over the course of two days, the Council will consider dozens of resolutions that will shape the direction of ACEP for the coming year and beyond. Get your idea or policy considered at the Council meeting Oct. 25–26, 2014, in Chicago by following these guidelines.

1. Resolutions must be submitted by at least two ACEP members or by any component body represented in the Council.
2. Resolutions may be submitted by mail, fax, or email (preferred), Resolutions are due at least 90 days before the Council meeting.
3. Resolutions consist of a descriptive Title, a Whereas section, and a Resolved section. The Council only considers the Resolved when it votes, and the Resolved is what the Board of Directors reviews to direct College resources.

4. There are two types of resolutions: general resolutions and bylaws resolutions. General resolutions require a majority vote for adoption, and bylaws resolutions require a two-thirds vote.
5. Councilors receive the resolutions prior to the annual meeting along with background information and cost information developed by ACEP staff. Resolutions are assigned to refer-

continuing on page 6
8. When considering a resolution, the Council’s options are to adopt the resolution as written; adopt, as amended by the Council; refer to the Committee on Rules, the Council Standing Commit- 
tee, or the Bylaws Interpretation Committee; or not adopt (defeat or reject) the resolution.
9. ACP has more resources on the resolution process at www.acp.org/council. Review the “Guidelines for Writing Resolutions” for tips.
10. Writing and submitting councils resolutions keeps our College healthy and vital. A Coun-
selor resolution is a great way for members to provide information to their colleagues and ACP 
Leadership. Please take advantage of this opportunity and exercise your rights as part of our emer-
gency medicine community.
11. Stop reading, and go write your resolution.
Praise for Electronic Patient Communications

Thank you for the "Text Rx" article that appeared in the March 2014 issue of Academic Emergency Medicine.

I share your enthusiasm for reaching patients electronically to check on their well-being. My ED uses a system recommended by the Robert Wood Johnson Foundation to contact discharged patients by text and email. We have more than one site of bleeding.

Confidence interval

In the RE-LY study, the rate of any gastrointestinal bleeds in patients receiving PRADAXA 150 mg twice daily was 0.7% (3/408 patients) on dabigatran at a dose of 110 mg twice daily, vs. 2.2% (9/408 patients) on warfarin (n = 416 patients receiving PRADAXA 150 mg twice daily). The RE-LY study was a placebo-controlled trial in which 1430 patients received PRADAXA 150 mg twice daily for a median duration of treatment of 165 days. Table 5 shows the number of patients experiencing bleeding events in the study.

Table 3: Bleeding Events in RE-COVER and RE-COVER II Treated Patients

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<th>Treatment</th>
<th>N (%)</th>
<th>Hazard Ratio (95% CI)</th>
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<td>PRADAXA 150 mg twice daily</td>
<td>37/201 (1.8)</td>
<td>1.43 (0.95, 2.13)</td>
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<tr>
<td>Placebo</td>
<td>51/201 (2.5)</td>
<td>1.60 (1.04, 2.45)</td>
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Table 4: Bleeding Events in RE-MEDY and RE-SONATE Treated Patients

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<tr>
<th>Treatment</th>
<th>N (%)</th>
<th>Hazard Ratio (95% CI)</th>
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<tr>
<td>PRADAXA 150 mg twice daily</td>
<td>101 (4.0)</td>
<td>1.06 (0.74, 1.51)</td>
</tr>
<tr>
<td>Placebo</td>
<td>170 (6.7)</td>
<td>1.12 (0.77, 1.62)</td>
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I would like to point out the error in this argument. The etiology of idiopathic intracranial HTN is different than getting a CT for other etiologies. In this condition, LP is safe because the pressures are equal since it is communicating hydrocephalus, thus pressure in ventricles and the subarachnoid space are equal to that of the lumbar cistern, thus LP is safe.

For other causes, I would argue LP before CT is not safe. In the study you cite, 52 of 56 patients had uneventful LP in spite of abnormal CT—well, what if one of those four was a family member? I don’t care for stats always when that small percent is me, I would be demanding a CT before the LP. Also, what happened to those four patients? I did not find the study so was not able to see.

Thank you, and great article!

—Mark Rollins, MD
Atlanta, Georgia


dr. Klauser Responds

I bring up a very reasonable point worth careful consideration. If a provider feels more comfortable obtaining a CT prior to LP, they should probably order one.

However, it is important to note that increased ICP is not truly associated with herniation following LP; brain shift is the phenomenon we should be concerned about.

In the Hasbun study you noted, the 56 patients had abnormal findings on CT and the four did not have bad outcomes associated with LP, but the clinician decided not to perform the LP due to noted mass effect (three severe and one mild).

In addition, the authors note that all four had no more than two critical predictors predicting this finding.

In closing, despite this age-teaching, there is little evidence supporting the supposition that CT must be performed routinely prior to LP.

Thank you for your letter.

—Kevin Klauser, DO, EJD, FACEP
Canton, Ohio

Table 3: Bleeding Events in RE-COVER and RE-COVER II Treated Patients

<table>
<thead>
<tr>
<th>Treatment</th>
<th>N (%)</th>
<th>Hazard Ratio (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>PRADAXA 150 mg twice daily</td>
<td>37/201 (1.8)</td>
<td>1.43 (0.95, 2.13)</td>
</tr>
<tr>
<td>Placebo</td>
<td>51/201 (2.5)</td>
<td>1.60 (1.04, 2.45)</td>
</tr>
</tbody>
</table>

Dr. Klauser,

I was reading today (3/25/14) “Myths in Emergency Medicine: Part 2” in ACEP Now (March 2014, p. 17).

I wanted to ask you about myth number four: CT before LP. You state: “Consider the number-one treatment for idiopathic intracranial hypertension (pseudotumor cerebri). Not only is it safe to LP these patients without risk of herniation, it’s recommended.”

I would like to point out the error in this argument. The etiology of idiopathic intracranial HTN is different than getting a CT for other etiologies. In this condition, LP is safe because the pressures are equal since it is communicating hydrocephalus, thus pressure in ventricles and the subarachnoid space are equal to that of the lumbar cistern, thus LP is safe.

For other causes, I would argue LP before CT is not safe. In the study you cite, 52 of 56 patients had uneventful LP in spite of abnormal CT—which, what if one of those four was a family member? I don’t care for stats always when that small percent is me, I would be demanding a CT before the LP. Also, what happened to those four patients? I did not find the study so was not able to see.

Thank you, and great article!

—Mark Rollins, MD
Atlanta, Georgia

Revisiting CT Before LP

I was reading today (3/25/14) “Myths in Emergency Medicine: Part 2” in ACEP Now (March 2014, p. 17).

I wanted to ask you about myth number four: CT before LP. You state: “Consider the number-one treatment for idiopathic intracranial hypertension (pseudotumor cerebri). Not only is it safe to LP these patients without risk of herniation, it’s recommended.”

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Thank you, and great article!

—Mark Rollins, MD
Atlanta, Georgia
have Medicaid expansion, there’s simply not a payer class. With the states that do not have Medicaid plus 20 to 30 percent of the charge, which leaves a significant “balance bill” due from the patient. The accessibility of the provider charges less the unreasonable low reimbursement by the health plans is being transferred to the patient, and the patient’s having to pay that in states where there isn’t a restriction on balance billing, and that’s a real problem for emergency medicine specialty exist on a Medicaid-based model. So if we look out over the next five to 10 years, can the emergency medicine specialty exist on a Medicare- or Medicaid-style reimbursement methodology? I don’t think it can exist in the way it does today with independent ED group practices. The recently published Health Affairs study showed reimbursements set to Medicare and Medicaid would produce double-digit losses for most EDs and that historically privately insured patients have subsidized all other ED payer classes. With the states that do not have Medicaid expansion, there’s simply not enough reimbursement in the system to pay for the moral imperative, the Emergency Medical Treatment & Labor Act (EMTALA) imperative, and the imperative of emergency medicine to care for all comers to the ED. I think that’s the single biggest challenge we have in EM.

**GG:** I’d say adapting to all of the changes brought about by the ACA would be the generic answer. My number-one concern would be the prevalence of the high-deductible health plans (HDHP) being offered through the exchanges. What you really see is simply a new form of self-pay when you have $2,000 and $3,000 deductibles and higher, shifting patients from uninsured to the high-deductible plan. A second concern would be the morphing of the Physician Quality Reporting System into a penalty phase in the next couple of years coupled with the arrival of value-based purchasing modifier systems, which arise out of the ACA. Those two things, in combination, bring exceptional complexity, and when you see the flow chart of those two things together, it introduces almost a sense of despair and skepticism to the house of medicine. In terms of early arrivals, the GOT in 2013, this past calendar year, has resulted in major national commercial payers systematically lowering their non-par reimbursements and justifying it from the GOT regulation. Of course, two of those three rates in the GOT criteria are set by the payers and are thus in a black box unknown by providers. So payers have realigned, and their legal counsel has sanctioned the systematic ratcheting downward of these payments, and that poses serious challenges this year and beyond.

**CE:** One of the things that I think we are all still reeling from is this ICD-10 schizophrenia. We had physician groups gearing up for it and hospitals paying millions of dollars to get ready for all of the big change that didn’t happen. So do we continue to prepare for it? Do we wait until the last minute and see if we want to spend some more money keeping people’s skills sharp and then take a chance it’s going to get put on the back burner again?

Additionally, somehow the patients have become customers, and with them as customers, the hospitals are driven to assure top levels of patient satisfaction. You have your doctor-to-doctor time; you have your patient comments about the care they’re receiving. Many times, none of that has to do with the quality of medical care that’s being provided, and I think we’ve seen physicians who take a little bit more time with their patients and are a little bit slower than everyone else get called to the carpet for that. I see that happening more and more as we talk to more groups across the country. Patients have been given more power, but I’m not sure they’ve been educated on how to use it, and that concerns me a lot.

I see the changes with so many of the payers. Many of them have now set up their own internal audit departments to audit our claims and track whether our charges go up the slightest little bit or if one physician is charging higher than another. Sometimes they don’t really care about the reasons; they just want to bust your chops about it, and we have to defend ourselves. I see that more and more groups are spending more and more money on nonclinical care just to stay ahead of the audits and stay ahead of the impact of electronic medical records (EMRs) and their documentation issues. So, what I see is money tightening up and nonclinical expenses very necessarily going through the roof.

**KK:** Those are great follow-up comments, and I have to apologize: when you’re not going first, it leaves you a little less to say, but you’ve found some great things to add to the conversation. So, John, what do you think?

**JJ:** There are a couple of things I always like to include in the top health care challenges in general: one being the development of health information exchanges and the second being health care going mobile or health care going retail. Shifting to the top emergency medicine issues, I really believe there are three of them, one of which is defining emergency medicine’s role and function in the care continuum. The second one is emergency medicine self-defining the value metrics by which it will be measured, and the third challenge is the specialty has to really assess and address some macro issues that I truly believe are directly impacting upon two core issues, namely EMTALA and the prudent layperson definition of an emergency. These issues are the infusion of the newly insured; the urgent care explosion; the high-deductible plans, as my three colleagues have mentioned; hospitals going into the insurance business; and what I would call the “retailization” of health care. When patients can self-direct their own lab tests today, the landscape has truly changed. I believe these landscape changes are going right at the heart of core issues of the specialty.
**KK:** I think those are huge, and I appreciate the segue into the specific challenges for emergency medicine. Let’s reverse the order here on the challenges specific to EM. Caral, what do you think? Any particular challenges you see that are unique to emergency medicine that haven’t been mentioned already or that are more global to health care in general but something specific to emergency medicine?  

**CE:** I think we’re still the front door to much of the health care that is provided, but what happens on the other side of that door has been significantly redefined and will continue to be redefined. We have Walgreens and Walmart now providing urgent care, and patients don’t quite know where they fit and where they’re supposed to go. I think that’s going to be a huge challenge for emergency departments. The hospitals are pushing the emergency physicians to get into some involvement with urgent care, some involvement with hospital medicine, some involvement with Walgreens and Walmart type urgent care to assure quality in their areas, to try to compete with them, and our physicians are doing a lot more than seeing patients.

**GH:** I think simply getting the message out about the unique payer mix and payment stream for emergency medicine is a major and vital challenge. I’ve developed a suite of tools or talking points for advocates within our organization and even utilized it to some degree within our trade association. I liken the emergency department group to swimmers in a very turbulent ocean, and that’s because fully 50 percent or more, often 60 to 70 percent, of the patients who arrive are either low pay, that is Medicaid, or no pay, that is uninsured. The average would be 20 percent uninsured and 30 percent Medicaid. Now, what other business has 50 to 60 percent of its customers in a status of no pay or low pay and whose revenues fund about one-third of the cost to deploy the service? None.

So what are the lifelines for this swimmer who’s in very turbulent waters indeed? Well, there are two lifelines: the one is fair commercial payments—that is, from the Blues and from the other national payers, such as Aetna, Cigna, United—and the second would be a hospital subsidy. A lifeline is necessary whenever there is inadequate and unfair payment from the commercial payers to offset the losses from the uninsured and Medicaid, and what you see now with the “Greatest of Three” taking shape is a massive shift back to the very patient who is unable or unwilling to pay. The question is: will the hospitals now be able to step up and increase their subsidies—or, in some cases, start subsidies for the first time—in order to strengthen the one remaining lifeline keeping the emergency group intact? Of course, I don’t know for sure, but I am worried about the fiscal health of many of our hospital partners in an age in which disproportionate share support has been scaled back massively and value-based measures will result in steep penalties to hospitals.

“I can see the ED of the future having more to do than provide ED care...I can see our role changing, and if we’re indeed going to be responsible for that and accountable for these outcomes, we’re going to need a lot more influence...” –Caral Edelberg, CPC, CPMA, CAC, CCS-P, CHC

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**KK:** Ed, do you want to comment about 2007?

**EG:** We frequently hear from Medicaid agencies that the way they’re going to reform Medicaid, or a principle way they’re going to reform Medicaid, is to keep people out of the emergency department, and we’ve seen those attempts despite the EMTALA mandate. We’ve seen that in Washington state initially with their very restrictive diagnosis and ED visit limitations. We’ve seen expanded cost sharing (eg, coinsurance) for non-emergency use of the ED in what I call the “Arkansas Model” of Medicaid expansion. This premium support model has been picked up by Pennsylvania, Iowa, and several other states seeking to use federal matching funds to purchase Medicaid health plan policies for the working poor. In North Carolina, Medicaid is looking at Accountable Care Organizations (ACOs) to drive significant percentages of patients out of the ED. Yet the Oregon Medicaid expansion study showed that when Medicaid expanded in Portland, ED utilization went up approximately 50 percent. It highlights the challenges we have of the moral and legal imperative of the EMTALA mandate and the issues surrounding the chronic conditions, which you know much better than all of us, in terms of that Medicaid patient base and how we deal with it. How do we arm the state chapters to be able to go and make the case that reforming Medicaid on the backs of emergency physicians is not the answer? Maybe 20 years ago, we heard, “Stand behind EMTALA and prudent layperson, and don’t engage in the conversation.” I don’t hear that much anymore—emergency physicians want to be at the table. As Dr. Lynn Massingale of TeamHealth has said for years, “We’re either at the table or on the table,” and that is very hopeful for the future that the level of engagement of ED physicians has really changed for the better.

CONTINUED on page 10
**UNSGN HEROES OF EMERGENCY MEDICINE**

**CONTINUED FROM PAGE 9**

**KK:** Something you mentioned the readers might want to hear more about: you noted cost-sharing models that are out there. Is there anything you would like to expand beyond just high-deductible plans shifting burden to the patient?

**EG:** You know the ACO, the shared savings models, and the bundled payment experiments that were done in what was called the ACE demonstration, which is now the Bundled Payments for Care Improvement Initiative on the Medicare side. I think both of those types of programs appear from a 40,000-foot perspective as an attempt to move away from fee-for-service to something else. The question is how do you move chronic care and patients with multiple comorbidities into a bundled payment? Maybe a bundled payment works great on hips and knees or coronary artery bypass grafting, but is it really going to affect how Dr. Klauser is going to work up the nursing home patient who is weak and dizzly, has several chronic conditions like congestive heart failure, and does not know why he doesn’t feel good? How much is his ordering or diagnostic treatment protocol going to impact the total bundle payment at the end of the day? I think the good news for us in nobody’s totally figured out emergency medicine yet from an ACO or bundled payment perspective—we’re in the very early innings. We’re the X factor. When you listen to officials at leading hospitals who have experimented with bundled payment and ACOs talk about bundled payment arrangements or ACOs, they scratch their heads and say, “Well, but we’ve had to carve out emergency medicine.”

**KK:** When they can’t figure us out, we have two ways to look at it. It does provide great opportunity for us to shape their understanding and create, perhaps, a larger scope for emergency medicine. But if they don’t understand EM, they might also interpret this in another direction, as many have already done: that the emergency department is an expensive place to receive care and people shouldn’t go there, clearly a preposterous assumption about our specialty.

**EG:** But one of the biggest challenges we have, Kevin, is the old pit-doctor mentality from 20 years ago that I do not want to understand how these changes may impact my practice and my livelihood, that I’m going to work my shift, I’m going to go home, I’m going to take care of my family, I’m going to educate my kids and whatever, and I’m not going to get involved. The biggest threat is that a doctor who thinks all of this stuff is a lot of white noise and somebody else is going to do it for him or her. We all can make a difference, and we should consider how we all stand on the shoulders of the giants of emergency medicine who carved this specialty out of solid rock—they all stepped up time and time again, and we all need to follow their leadership and example.

**KK:** We’re going to do it collaboratively, and we all have to be involved; at the least, we have to be informed. Do you remember that book from several years ago, Who Moved My Cheese? Well, they’re moving our cheese. We have to evolve and move with it. If we don’t change the way we think and the way we practice, this specialty will be at real risk. I have a question for each of you, based on a comment that Ed made about utilization. How many tests does it take to diagnose a hip fracture? Most emergency physicians when they ini-

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**INDICATIONS AND USAGE**

ADASUVE® (loxapine) inhalation powder, for oral inhalation use, is a typical antipsychotic indicated for the acute treatment of agitation associated with schizophrenia or bipolar I disorder in adults. Efficacy was demonstrated in 2 trials in acute agitation: one in schizophrenia and one in bipolar I disorder.

**HOW LONG CAN YOU WAIT?**

**IMPORTANT SAFETY INFORMATION**

**WARNING: BRONCHOSPASM and INCREASED MORTALITY IN ELDERLY PATIENTS WITH DEMENTIA-RELATED PSYCHOSIS**

Bronchospasm

ADASUVE can cause bronchospasm that has the potential to lead to respiratory distress and respiratory arrest. Administer ADASUVE only in an enrolled healthcare facility that has immediate access on-site to equipment and personnel trained to manage acute bronchospasm, including advanced airway management (intubation and mechanical ventilation). Prior to administering ADASUVE, screen patients regarding a current diagnosis, history, or symptoms of asthma, COPD and other lung diseases, and examine (including chest auscultation) patients for respiratory signs. Monitor for signs and symptoms of bronchospasm following treatment with ADASUVE. Because of the risk of bronchospasm, ADASUVE is available only through a restricted program under a Risk Evaluation and Mitigation Strategy (REMS) called the ADASUVE REMS. Increased Mortality in Elderly Patients With Dementia-Related Psychosis

Elderly patients with dementia-related psychosis treated with antipsychotic drugs are at an increased risk of death. ADASUVE is not approved for the treatment of patients with dementia-related psychosis.

- ADASUVE is contraindicated in patients with the following:
  - Current diagnosis or history of asthma, chronic obstructive pulmonary disease (COPD), or other lung disease associated with bronchospasm
  - Acute respiratory signs/symptoms (eg, wheezing)
  - Current use of medications to treat airways disease, such as asthma or COPD
  - History of bronchospasm following ADASUVE treatment
  - Known hypersensitivity to loxapine or amoxapine. Serious skin reactions have occurred with oral loxapine and amoxapine
- ADASUVE must be administered only by a healthcare professional
- Prior to administration, all patients must be screened for a history of pulmonary disease and examined (including chest auscultation) for respiratory abnormalities (eg, wheezing)
- Administer only a single 10 mg dose of ADASUVE within a 24-hour period by oral inhalation using the single-use inhaler
ADASUVE® (loxapine) inhalation powder

HELP DEFUSE THE SITUATION BEFORE AGITATION ESCALATES FURTHER

ORAL INHALATION

Breath-actuated, single-use, ready-to-use inhaler

FAST ONSET

Statistically significant reduction in agitation at 2 hours, with improvement rapidly achieved at 10 minutes post-dose

Reduction from baseline in agitation symptoms

ENDPOINT          SCHIZOPHRENIA    BIPOLAR I DISORDER

AT 2 HOURS (SECONDARY) ADASUVE           49%           53%       PLACEBO          33%           27%

AT 10 MINUTES (PRIMARY) ADASUVE           19%           23%       PLACEBO          10%           10%

The mean baseline PEC scores in all treatment groups were 17.3 to 17.7.

IMPORTANT SAFETY INFORMATION (continued)

- After ADASUVE administration, patients must be monitored for signs and symptoms of bronchospasm at least every 15 minutes for at least 1 hour
- ADASUVE can cause sedation, which can mask the symptoms of bronchospasm
- Antipsychotic drugs can cause a potentially fatal symptom complex called Neuroleptic Malignant Syndrome (NMS), manifested by hyperpyrexia, muscle rigidity, altered mental state, irregular pulse or blood pressure, diaphoresis, and cardiac dysrhythmia. Associated features can include elevated serum creatine phosphokinase (CPK) concentration, rhabdomyolysis, elevated serum and urine myoglobin concentration, and renal failure. If NMS occurs, immediately discontinue antipsychotic drugs and other drugs that may contribute to the underlying disorder, monitor and treat symptoms, and treat any concomitant serious medical problems
- ADASUVE can cause hypotension, orthostatic hypotension, and syncope. Use with caution in patients with known cardiovascular disease, cerebrovascular disease, or conditions that would predispose patients to hypotension. In the presence of severe hypotension requiring vasopressor therapy, epinephrine should not be used
- Do not use ADASUVE with caution in patients with a history of seizures or with conditions that lower the seizure threshold. ADASUVE lowers the seizure threshold. Seizures have occurred in patients treated with oral loxapine and can also occur in epileptic patients
- Use ADASUVE when driving or operating machinery. ADASUVE can impair judgment, thinking, and motor skills
- The potential for cognitive and motor impairment is increased when ADASUVE is administered concurrently with other CNS depressants
- Treatment with antipsychotic drugs caused an increased incidence of stroke and transient ischemic attack in elderly patients with dementia-related psychosis; ADASUVE is not approved for the treatment of patients with dementia-related psychosis
- Use of ADASUVE may exacerbate glaucoma or cause urinary retention
- The most common adverse reactions (incidence ≥2% and greater than placebo) in clinical studies in patients with agitation treated with ADASUVE were dysgeusia, sedation, and throat irritation
- ADASUVE Category C. Neonates exposed to antipsychotic drugs during the third trimester of pregnancy are at risk of extrapyramidal and/or withdrawal symptoms after delivery; ADASUVE should be used during pregnancy only if the potential benefit justifies the potential risk to the fetus
- Discontinue drug or nursing, taking into account the importance of the drug to the mother
- Discontinue drug or nursing, taking into account the importance of the drug to the mother
- The safety and effectiveness of ADASUVE in pediatric patients have not been established

References:
1. ADASUVE [package insert]. Horsham, PA: Teva Select Brands, a division of Teva Pharmaceuticals USA, Inc; December 2013.

Please see Brief Summary of Prescribing Information, including Boxed Warnings, on following pages.
sentially responsible for driving front-end cost for health care, which is everything up to, and possibly including, the admission or everything to the consultant coming in. I can see our role changing, and if we’re indeed going to be responsible for that and accountable for these outcomes, we’re going to need a lot more influence and a lot more power over what happens to the patients we’re responsible for.

KK: That’s a great perspective, I hadn’t thought about it that way. I really like the way you think on that, Greg. Caral, any additional thoughts on that?

CE: I see our specialty changing. I don’t want to use the word “gatekeeper” because it has such a negative connotation for so many health care professionals, but it’s a role I think we’re going to end up playing. We’re going to be directing more than just the emergency care. I can see the emergency department of the future having more to do than provide emergency department care. I can see us essentially responsible for driving front-end cost for health care, which is everything up to, and possibly including, the admission or everything to the consultant coming in. I can see our role changing, and if we’re indeed going to be responsible for that and accountable for these outcomes, we’re going to need a lot more influence and a lot more power over what happens to the patients we’re responsible for.

KK: That’s a great perspective, and that actually leads me to my next question. I’m going to ask you all just for one line about whether you think the scope of practice in emergency medicine will be shrinking or expanding in the next couple of years.

EG: Expanding.
we were doing to support the specialty, and obviously we couldn’t be members of ACEP because we weren’t physicians. We needed a trade umbrella to represent the professional inter-
ested stakeholders to go to Washington and go to the state capitals, and become the vehicle for us to be able to channel all of our energies.

It was started in a crisis, the Medicare reimag-
nification, best practices committee, and the inter-
sted person and Medicaid restrictive diagnosis and triage fee crises hit us right afterwards in the late ’90s. We’re helped emergency medicine because, as one of our colleagues said, when we come together in EDPA, “we take off our respective company uniforms and focus on the greater good and on achieving results.” It’s re-
warding on many levels because we have been fortunate to be part of something larger than each of our companies or ourselves. The as-
sociation allowed us to represent our clients with ACEP and other EM stakeholders. It’s been a fantastic working relationship, and I believe we have made a difference because we cared and we worked together.

JH: I consider it my duty, my responsibility, to be a practice manager and resource for success for the cli-
icians’ shoulders and to create and implement high-performing solutions in the areas of infor-
mation technology, government compliance, three-party audits, third-party enrollment, coding, billing, cash collection, management negotiations, and fee schedule compliance.

CE: I feel like I have to keep reinventing myself. I own a very large technology coding and com-
pliance company. In order to do that and repre-
sent what we do to support our clients, which are hospitals, physicians, and payers and a little bit more of everything, I’ve never been able to step away from the sense that I have to know the details. I feel like in order to represent the priori-
ties and potential solutions to our physicians and our clients, I really have to understand the issues, which keep changing all the time. I al-
ways feel like I have to be down in the weeds in

Table 1. Adverse Reactions in 3 Pooled Short-Term, Placebo-Controlled Trials (Studies 1, 2, and 3) in Patients with Schizophrenia or Bipolar Disorder

<table>
<thead>
<tr>
<th>Adverse Reaction</th>
<th>Placebo (n = 263)</th>
<th>ADASUVE (n = 259)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Dysgeusia</td>
<td>0%</td>
<td>5%</td>
</tr>
<tr>
<td>Sedation/Somnolence</td>
<td>12%</td>
<td>17%</td>
</tr>
<tr>
<td>NMS</td>
<td>0%</td>
<td>0%</td>
</tr>
<tr>
<td>Gastrointestinal</td>
<td>0%</td>
<td>0%</td>
</tr>
<tr>
<td>Hematologic</td>
<td>0%</td>
<td>0%</td>
</tr>
<tr>
<td>Neurologic</td>
<td>0%</td>
<td>0%</td>
</tr>
<tr>
<td>Cardiac</td>
<td>0%</td>
<td>0%</td>
</tr>
</tbody>
</table>

**Reference**

1. Wilson M, Cutler D. Emergency department profits are likely to continue as the affordable care act expands coverage. Health Aff 2014;33(5):792-795.
7.2 Anticholinergic Drugs

Elderly patients with dementia-related psychosis treated with antipsychotic drugs are at an increased risk of death compared to placebo patients (generally one to five deaths per 1,000 patients treated). While the overall risk of death may be increased for all antipsychotic drug therapies, the risk is most prominent in elderly and suicidal patients. There were suicides in placebo and drug groups; however, they were more frequent in the drug groups in one of the short-term studies of elderly patients with dementia-related psychosis. In general, cardiac monitoring should be used in patients weighing 55 kg or more who are taking 4 mg or more daily, and in any patient who is taking 10 mg or more daily. In patients weighing less than 55 kg who are taking 4 mg or more daily, cardiac monitoring should be done periodically during the first month of treatment. Patients with a ≥ 20% decrease in FEV1 from baseline in the Healthy Volunteer, Asthma, and COPD categories are cumulative; i.e., a subject with a maximum decrease of 21% is included in all 3 categories. Patients with a ≥ 20% decrease in FEV1 did not receive a second dose of study drug.

The data in Figure 7: LS Mean Change from Baseline in FEV1 in Patients with Asthma show that patients receiving ADASUVE 10 mg had a greater decrease in FEV1 compared to placebo. The mean decrease in FEV1 in patients receiving ADASUVE 10 mg was 7.1% compared to 3.6% in patients receiving placebo. A similar trend was observed in the Healthy Volunteer and COPD categories. The mean decrease in FEV1 in patients receiving ADASUVE 10 mg was 8.9% and 8.6%, respectively, compared to 3.8% and 3.9% in patients receiving placebo.

In the 3 short-term (24-hour), placebo-controlled trials of ADASUVE in 259 patients with agitation associated with schizophrenia or bipolar disorder, 52% (35/67) of patients receiving ADASUVE had akathisia compared to 30% (20/67) of patients receiving placebo. Akathisia (motor restlessness) has also occurred.

ADASUVE is contraindicated in patients with a known hypersensitivity to loxapine or any of the other components of ADASUVE. ADASUVE is not recommended for use in patients with a history of seizures or who are at risk of seizures due to other causes. The use of antipsychotic drugs in the treatment of patients with a known history of seizures or a history of seizures during treatment with other antipsychotic drugs is not recommended.

ADASUVE is generally well tolerated, and patients treated with ADASUVE in clinical trials had a lower incidence of adverse events than patients treated with placebo. The most common adverse events associated with ADASUVE treatment were dizziness, headache, and nausea.

In long-term trials, the most common adverse events associated with ADASUVE treatment were dizziness, headache, and nausea. These events were generally mild to moderate in severity and were observed in a similar frequency in patients treated with ADASUVE and placebo.

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In long-term trials, the most common adverse events associated with ADASUVE treatment were dizziness, headache, and nausea. These events were generally mild to moderate in severity and were observed in a similar frequency in patients treated with ADASUVE and placebo.
A quarter of doctors captured more than a third of the $77.4 billion in payments, and one in three of the top earners were ophthalmologists.

Granovsky. “If your hospital has a lot of specialty services, a large volume of Medicare patients will be seen, and they will be more complex.”

Even factors such as proximity of urgent care centers will filter out lower-acuity patients and result in a higher ratio of more complex patients treated in the ED. Nearby nursing homes, too, can bring up the cost of providing care by simply contributing large volumes of patients.

The data release comes amid ongoing debate over how to better control costs in the Medicare program and how to rein in unnecessarily expensive care while improving patient outcomes. Medicare spending is near $600 billion annually, including payments to hospitals and physicians and costs for drugs. Cutting wasteful and fraudulent payments is one way to slow cost growth.

While the transparency intended by releasing the data might help some consumers choose which doctors they would like to see, when it comes to care in the ED, Dr. Granovsky is skeptical the information is useful for patients. “I am not sure it is a valuable reference tool to help select an emergency department for care,” he said.  

KELLY APRIL TYRELL is a freelance journalist based in Wilmington, Delaware.
THE ZOHYDRO ER AFFAIR

The conspiracy theory behind the drug approval

BY PAUL KIVELA, MD, MBA, FACEP

Whether you buy into a Food and Drug Administration conspiracy theory or not, there are many problems with the approval of Zohydro ER. Although there is certainly disagreement over the benefits and risks of the medication, almost everyone can agree that Zohydro is one the most controversial recent drug approvals by the FDA.

The opposition to the FDA’s approval of Zohydro includes consumer groups, doctors, state and federal leaders, and multiple governmental organizations. FDA Commissioner Margaret Hamburg has received letters protesting the decision to approve Zohydro ER from at least 28 state attorneys general and multiple US senators. Massachusetts Gov. Deval Patrick and Vermont Gov. Peter Shumlin, issued directives to either ban or make it more difficult to prescribe and dispense the medication. Health insurers claim that opioid abuse costs more than $70 billion a year in direct health care costs.

The potential conflicts involving Zohydro ER illustrate the involved process and complicated business operations and influence that the pharmaceutical industry might have on the approval process.

The Zohydro ER story centrally involves a company called Alkermes. This company owns a product called Vivitrol, an extended-release version of naltrexone that is given in a monthly injection to treat opioid dependence. The medication was supported by several very controversial studies and, in October 2010, was approved by the FDA to treat and prevent relapse in patients with opioid dependence who have undergone detoxification treatment. The product has had a very slow start and has failed to meet sales expectations.

In 2011, Alkermes purchased a part of another pharmaceutical company, Elan, that originally made Zohydro ER, and, with it, an exclusive marketing deal with a pharmaceutical company called Zogenix, which has the right to market the drug. Interestingly, Alkermes chose not to list Zohydro ER as one of its products and instead left the product under the Zogenix product line. Not surprisingly, many think it is a conflict of interest for Alkermes to have a product that treats opiate abuse as well as a product that has the potential to greatly increase opiate abuse.

In May 2012, a bipartisan Senate committee launched an investigation into the conflicts of interest and issued a series of subpoenas to various “consumer” groups, including the American Pain Foundation, University of Wisconsin Pain & Policy Studies Group, the American Pain Society, the Federation of State Medical Boards, and The Joint Commission. Within days, the American Pain Foundation shut down, and it was disclosed that the Federation of State Medical Boards commissioned the production of its Responsible Opioid Prescribing—A Physician’s Guide and other publications by the American Pain Foundation, which received 90 percent of its $5 million in funding from pain medication manufacturers. Interestingly, the American Pain Foundation was very active in lobbying for greater pain medication use. In 2011, under investigation, Scott M. Fishman, MD, the main author of the pain guide and the past president of the American Pain Foundation, acknowledged that he had to add additional disclosures including honoraria from the pharmaceutical industry not previously disclosed.

In October 2013, the Milwaukee Journal Sentinel disclosed that the University of Rochester created two organizations: IMMPACT (Initiative on Methods, Measurement, and Pain Assessment in Clinical Trials) and ACTTION (Analgiesic Clinical Trial Translations, Innovations, Opportunities, and Networks). Some alleged potential pay-to-play arrangements involving meetings between pharmaceutical industry representatives and FDA officials occurred with these organizations. Pharmaceutical industries allegedly paid between $20,000 and $35,000 to send one representative to a two-day meeting. Zohydro ER’s original manufacturer may have participated in those meetings. Allegedly as a result of the meetings, the FDA approved a method, known as enriched enrollment, which allowed pharmaceutical companies doing pain studies to remove certain patients who did not respond well to a medication or could not tolerate it before the study began. This made it much easier for pharmaceutical companies to prove their medications were safe and effective.

On March 10, 2014, several senators called for a special investigation into these pay-to-play allegations against several pharmaceutical companies, physicians, and FDA officials. Sen. Joe Machin (D-WV) and Rep. Stephen Lynch (D-MA) introduced legislation to prohibit the FDA from approving similar drugs unless they are formulated to prevent abuse. Several Republican senators also demanded the FDA release safeguards to prevent abuse.

In March, Purdue Pharma announced it had developed a tamper-resistant version of its hydrocodone product for extended release. Interestingly, the company that markets Zohydro ER announced it will likely have a tamper-proof version of Zohydro ER ready by 2016.

On April 15, 2014, a US District Court issued a preliminary injunction citing that the ban ordered by Massachusetts Gov. Patrick was unconstitutional. The investigations are just beginning, and the controversy will undoubtedly continue. It is an unintended, and unfortunate, coincidence that Zohydro ER happens to have the word “ER” in it.

The Official Voice of Emergency Medicine
JUST SAY NO TO ZOHYDRO ER

The abuse potential of this new hydrocodone product makes it a possible contributor to the opioid misuse epidemic

BY FRANK LOVECCHIO, DO, MPH, FACEP

Despite the United States encompassing just 5 percent of the world’s population, it accounts for 84 percent of oxycodone (Oxycontin) and a whopping 99 percent of hydrocodone (Vicodin, Lortab) global consumption. Unfortunately, the opiate epidemic is worsening.

New, Controversial Opioid Option

Since March 2016, patients and physicians have had a new option with a hydrocodone tablet, Zohydro ER (hydrocodone bitartrate). Hydrocodone (Zohydro ER’s sole ingredient) is one of the most frequently prescribed and, unfortunately, abused opioids. One of the advantages of Zohydro ER is that it is safer, or high, can last up to 12 hours per dose. Zohydro ER is specifically indicated for the management of pain severe enough to require daily, around-the-clock, long-term opioid treatment and for which alternative treatment options are inadequate. Following Zohydro ER ingestion in pre-marketing studies, hydrocodone levels peak in five to six hours.

According to the package insert, the starting dose for patients who are not opioid tolerant is Zohydro ER 10 mg orally every 12 hours; this is the lowest dose. The package insert defines patients who are opioid tolerant (the target population) as those receiving, for one week or longer, at least 60 mg oral morphine per day, 25 mcg transdermal fentanyl per hour, 30 mg oral oxycodone per day, 8 mg oral hydrocodone per day, 25 mg oral oxymorphone per day, or an equianalgesic dose of another opioid.3

The presumed benefit of Zohydro ER—other than its strength—is that it doesn’t contain acetaminophen, as do Vicodin and Percocet. It’s on this basis that its maker, Zogenix, has argued that it’s safer than the alternatives. Currently, there are other opioid painkillers on the market that don’t contain acetaminophen. Current recommendations to minimize any risk of hepatotoxicity following chronic acetaminophen use resulted in the Food and Drug Administration recommending reducing the daily dose of acetaminophen to one capsule. Following overdose or accidental ingestion, it is prudent for patients to undergo an extended observation period. In my opinion, such patients should be observed at least 12 hours, assuming no naloxone was used, until future data are collected. In summary, the FDA approved Zohydro ER for the management of pain severe enough to require daily, around-the-clock long-term opioid treatment and for which alternative treatment options are inadequate. At least initially, if you prescribe this drug, be aware of this indication. Be aware that the peak may not occur for six hours in “normal” patients in “ideal” circumstances. Be aware that Zohydro ER is metabolized via P450 interactions P450 3A4, and drugs such as macrolides orazole antifungal inhibitors that inhibit this cytochrome may increase hydrocodone levels. Be aware that ethanol and other CNS depressants potentiate the effect. Be aware that clearance is altered, namely decreased in patients with hepatic and renal disease (not further identified). An FDA-approved patient medication guide, which is available with the product information and can be accessed at www.fda.gov/downloads/Drugs/DrugSafety/UCM370009.pdf, must be dispensed with this medication. Currently, it is approved as a Schedule II drug and can only be dispensed through a physician’s written prescription, and no refills are allowed. There are also stringent recordkeeping, reporting, and physical security requirements for Schedule II controlled substances. Considering the risk-benefit ratio, is this really much to be aware of for one drug?

Patients and physicians want better pain relief and improved patient satisfaction. Mandates warrant “significant decreases in pain scores,” further increasing the demand for a drug such as Zohydro ER. Logically, drug companies will increase supply based on demand. Regulations, lobbying, restrictions, and patient and physician education may decrease the demand for opioids. As individuals, and hopefully as a group, the simplest way for us to decrease demand for Zohydro ER and similar products is not to prescribe them. In conclusion, further data are available—especially addressing post-marketing safety—it is best to “say no to Zohydro” in the ED.

NEWS

Congress Questioning Ethics Behind FDA Zohydro Approval

Recent hydrocodone opioid approval raises concerns

BY JESSICA KINSELLA

The FDA’s approval of the hydrocodone drug Zohydro ER has been met with criticism from health care professionals and lawmakers alike, and members of Congress, including Rosa DeLauro (D-CT), Harold Rogers (R-KY) and Stephen Lynch (D-MA), have requested an investigation into the matter from the Office of the Inspector General.

Zohydro ER, approved in October 2013, contains five to 10 times more hydrocodone than any other drug on the market and lacks safeguards to prevent immediate release through the snorting or injecting of the drug’s easy-to-crush form. The FDA’s Anesthetic and Analgesic Drug Products Advisor Committee displayed overwhelming opposition to Zohydro ER, with an 11-2 vote against its approval. This recommendation, combined with efforts to reclassify hydrocodone from a Schedule III drug to a Schedule II drug, has led many members of Congress to question why Zohydro ER was approved at all.

Media sources, including Milwaukee’s Journal Sentinel and The Washington Post, claim that pharmaceutical companies influenced FDA officials during their participation in pay-to-play initiatives set up by professors at the University of Rochester and the University of Washington. Opioid misuse is becoming more frequent across the United States, with opioid overdoses accounting for an average of 15,000 deaths per year. In an effort to reduce opioid misuse, Massachusetts attempted to ban the sale of Zohydro ER, although a federal judge denied the request.

Congressional leaders are not the only ones taking notice of Zohydro ER’s controversial approval; many emergency physicians are aware of the questions surrounding the new prescription painkiller. While the investigation into Zohydro ER’s approval continues, it is up to individual emergency physicians to choose whether they prescribe the controversial drug or not.

—Jessica Kinsella is a writer based in Hoboken, New Jersey.

References

in LA County’s institutional disaster plan, one which stands for a 30-hour wait in the emergency department and no beds upstairs. Code Black was the norm in the country’s busiest ED at the time of the filming, so Dr. McGarry had his hands full learning the medicine, producing the documentary, and trying to maintain a personal life.

*Code Black* has astonished film festival attendees, winning awards at the Los Angeles Film Festival, the Aspen Filmfest, the Staxx Denver Film Festival, and the Hamptons International Film Festival, along with a special recognition at the Vancouver International Film Festival. Emergency physicians and laypeople alike may experience the difficulties and triumphs of being an emergency medicine resident in one of the nation’s busiest EDs, illustrating how difficult and challenging this environment really is. The film gives the real-world view from the inside, hopefully capturing the hearts and minds of those who know little about the safety net of our health care system yet are always reliant on its existence.

*Code Black* will be premiering June 20th in New York City at the IFC Center. In July and August, it will be released in 40 cities throughout the United States.

Dr. McGarry, who wrote and directed *Code Black*, recently sat down with ACEP Now’s medical editor in chief, Kevin M. Klauer, DO, EJD, FACEP, to talk about the process of making the movie and what he hopes to accomplish with the film.

**FRANCESCA BARATTA** is a freelance writer based in New Jersey.

**KK:** Give us an overview of *Code Black*.

**RM:** *Code Black* is a feature documentary film that I directed and was released to the film festivals in 2013. The film asks the questions: If you signed up to be a doctor today, and you came in with expectations for how you wanted to practice medicine, how long would it take before the greater system and the ideas of billing, profit, regulation, documentation, and medical legal practice start to chip away at those ideals? How are you going to protect them? And, once they are threatened, how do you fight back?

**KK:** When did you have the epiphany that “I’m qualified to do this, and this story has to be told”?

**RM:** I can speak to the filmmaking qualifications first: I had none. I considered an MSA in film school, but I declined that on the notion that if I did want to pursue filmmaking someday, I’d have a lot more access to the human condition as a physician than as a film student. The documentary really formed itself when I was a rotating medical student at USC. I was with the old LA General hospital and in the middle of this special area of the ER called C-Booth. I was struck by the intensity of that space. I wanted to know, “Why does intensity matter?” And my answer is that intensity shows us priorities. We found that physicians and health care staff find nostalgia in that notion of “remember when you were just a training doc and your biggest priority was patient care.” I miss that most basic of concerns. Now I’m worried about the right RVUs, statisticians, documents, and Press Ganey.

**KK:** Does it come out in the documentary that you were focused on care, but you lost focus because of all the regulatory red tape?

**RM:** That’s a major theme. The three acts of the film mirror how an emergency medicine physician might mature along the way, especially in a big, busy trauma center. When physicians watch it, they grasp onto that. When laypeople watch it, I think they start to say, “I didn’t realize that this is going on. Why is the system so focused on not us and not my doctor?”

**KK:** Who’s working with you on this project?

**RM:** In the first days of the project, we had the support of some of the USC faculty: Drs. Ed Newton, Billy Mallon, Jan Stoenberger, and Diku Mandavia, along with the hospital administration and the LA County Board of Supervisors. Beyond that, there was the filmmaking world: Mark Jonathan Harris, who’s a three-time Academy Award winner in documentary film, and Marti Noxon, who scripted for *ER*. 

**PHOTOS FROM CODE BLACK**

Top to bottom: Danny Cheng, MD, and Jaime Eng, MD (at left) with patient; Dave Pomeranz, MD, Ryan McGarry, MD, and Billy Mallon, MD, at bedside; Jamie Eng, MD, with patient; C-Booth at LA County Hospital.
**KK:** Do you think you’ll make another film?

**RM:** [The market research] says that a part two with just Billy Mallon might sell really well! As far as Code Black part two, we’ve probably said enough on this topic in the feature.

**KK:** How did it affect your personal life?

**RM:** It was significant. I didn’t expect to make a film as a resident. It was three years of no vacation, and any day off I had went toward this effort. The goal of residency is to learn how to be a proficient physician, and I had to be careful in order to assure that the film could never be more important than that. I have to thank every one of my classmates and my program director for a lot of support along the way, whether for last-minute trades or just the ability to take a 20-minute nap on a shift. Even still…residency by default should consume you. You put them together at the same time, and there’s going to be some fallout.

**KK:** Do you regret the decisions you’ve made?

**RM:** I regret allowing the pressure to dictate how I acted and how I thought. I certainly don’t regret making Code Black at the time that I did. One thing that I feel very lucky to have experienced is nonmedical people sitting through some pretty tough stuff in cases we show, and at the end of the film people give us a standing ovation. I wish I could share that with every physician, nurse, and X-ray tech who leaves a really tough shift.

**KK:** Is this just a venture of altruism, or is there the potential for you to have some well-deserved financial benefit?

**RM:** It takes in the low seven figures to produce documentaries. Your best-case scenario is a film of this scope. Nobody makes a buck off a documentary like this.

**KK:** Roughly, what does it take to produce a film of this scope?

**RM:** The film is supported by SonoSite Fuji-Film. We made an agreement with them that said in exchange for private support for the production, we would make a true story about filmmaking by default did this without any control with regards to creative oversight or required brand placement. We went through 16 cuts on Code Black. For a documentary, you’re building an essay from a vast vocabulary of footage that exists as a library in volume. So you have an editor, an assistant editor, an archivist, a sound designer, a sound mixer, a color corrector, and, of course, all of the producers it takes to make a documentary.

**KK:** I think it can change perspectives a great deal about what emergency medicine is. I hope that you have a successful career as a director, and I hope we don’t lose you from emergency medicine too soon because of it.

**RM:** I wouldn’t give up my shifts for anything. I’ve met Academy Award winners who would love to have the ability to bridge their time in between projects with intermediate employment but don’t have the opportunity to do what we do. It really gives that crowd an entirely different perspective when they’re worried about a publicist or a new script and you’re saying, “You know, I just helped save a life.” It’d be crazy to ever give up on that investment because it’s really a gift.

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The availability of inpatient psychiatric care has worsened significantly and progressively over the past four years on state and national levels. As inpatient psychiatric beds have become increasingly scarce, the number of patients seeking or requiring psychiatric assistance has also increased. These patients are spending increased time “boarding” in emergency departments, and with beds scarce and increasingly far afield, many require transfer to facilities many miles away. In the meantime, emergency physicians and other emergency department personnel must dedicate significant time and resources to not only searching for placement, but also attending to patients’ needs while ensuring the safety of both patients and departmental staff for the duration of patients’ ED stays. This leads to increased throughput times for other patients, a frightening environment for delivering care, patient safety issues, and decreased satisfaction for patients and providers.

A brief review of the literature and national statistics on mental health care confirms what most of us already know from experience: the number of inpatient psychiatric beds nationally falls woefully short of what is necessary to meet current demand. The Treatment Advocacy Center recommends that each state should have 50 public inpatient psychiatric beds for every 100,000 people in a state’s population.1

According to the ACEP 2014 State-by-State Report Card, only three states (Mississippi, Missouri, and Arkansas) hit this target number, while 31 states had 50 percent or fewer of the target number of beds.2 Unfortunately, there is little hope for improvement in these numbers as state budgets continue to cut billions of dollars from public mental health spending.

The problem of the inadequate supply of inpatient psychiatric beds affects both psychiatric patients and emergency providers. The external stimuli associated with the busy emergency department environment have been shown to increase patient anxiety and agitation, leading to increased risk of symptom exacerbation or elopement of patients seeking treatment for mental health or substance abuse issues, which poses a danger to patients and staff.3 Elopement before screening and treatment is dangerous and leads to increased risk of self-harm and suicide.4 Furthermore, the need for increased security and additional ancillary staff to monitor and protect these patients, emergency department staff, and other patients leads to increased labor costs.5 Additionally, the significant number of resources and personnel required to provide adequate care for these patients for extended periods may lead to delays in care of other ED patients. Poor clinical outcomes and increased morbidity and mortality have been directly linked to ED overcrowding and a lack of available ED beds.

The financial impact of boarding these patients is profound. A recent study at Wake Forest University Health Sciences Center found admitted psychiatric patients are associated with a 40 percent decrease in average physician reimbursement as compared to nonpsychiatric patients. Furthermore, the increased length of stay for each of these patients was determined to prevent the ED from caring for an additional 2.2 patients, leading to an overall financial loss to the system of approximately $2,400 per boarded psychiatric patient.1

The effects of budget cuts to public mental health care can be felt at the ground level in many emergency departments across the nation. Per the ACEP State-by-State Report Card, Iowa ranks 16th in the nation in number of psychiatric inpatient at 28 beds per 100,000 population. Although I work at the state’s only tertiary referral hospital, we lack sufficient psychiatric beds for our needs and are frequently forced to transfer patients to other facilities. We recently conducted a review of patient records for

With fewer places left to turn, patients in need of mental health services are increasingly flocking to emergency departments across the country. But in EDs, mental health patients are diverting attention from medical patients, tying up emergency beds for days while awaiting inpatient or community beds. They also are at increased risk of poor outcomes due to care that is delivered in a less-appropriate setting. Nationally, there are an average of 26.1 psychiatric beds per 100,000 people, according to ACEP’s 2014 Report Card. Meanwhile, the National Alliance on Mental Illness reports 61 million Americans experience mental illness each year, but state funding for mental health continues to drop. From 2009 to 2012, states cut $1.8 billion in funding, a decrease of 10 percent.

Recently, the shortage of psychiatric beds contributed to an overall D+ grade on ACEP’s 2014 Report Card on Emergency Medicine, which ranked access to care more heavily than any other measure. “We can do emergent care, stabilize individuals, but people need to go to the appropriate places to get the appropriate care,” said Jon Mark Hirshon, MD, MPH, PhD, FACEP, associate professor in the Department of Emergency Medicine at the National Study Center for Trauma and Emergency Medical Systems in Baltimore and the Report Card Task Force chair. For instance, late last year, Virginia State Sen. Creigh Deeds was repeatedly stabbed by his 24-year-old son, who then fatally turned a gun on himself. Deeds’ son received an emergency mental health evaluation the day before the incident, but he was released when no psychiatric bed could be found for him. These problems don’t typically start in the ED. Instead, they are symptomatic of larger flaws within the health care system. Dr. Hirshon recently testified at a Congressional hearing meant to address the psychiatric bed shortage. Within two hours of the start of a recent shift, ACEP President-Elect Michael Gerardi, MD, FAAP, FACEP, an emergency physician at Morristown Medical Center and Goryeb Children’s Hospital, both in Morristown, New Jersey, had already evaluated four psychiatric patients.

He expected at least one of them would board in the ED for two or three days. In addition to a shortage of available beds and staff, he said hospitals are often reluctant to admit patients whose insurance may not pay for mental health. This situation places EDs and hospitals at odds. EDs must treat every patient who comes through the doors. Hospitals are not bound by the same code. Dr. Gerardi is waiting to see if the Affordable Care Act changes the game.

Dr. Hirshon said some of his colleagues report that patients have boarded between two weeks and 42 days in the ED. He said it’s going to take creative solutions to solve this problem. Nationwide, a 2008 ACEP survey showed that 99 percent of emergency physicians admit psychiatric patients each week, and nearly 80 percent report psychiatric patients are boarded in the ED. “I think no single source is going to have all the answers,” Dr. Hirshon said. “The ED is a spot that works 24-7. 365 days a week and we’ve there to help people….Our mission is to help people, but that’s problematic when they have needs that are above and beyond what we have in the ED.”

KELLY APRIL TYRELL is a freelance journalist based in Wilmington, Delaware.
all psychiatric patients transferred out of the University of Iowa Hospitals and Clinics (UIHC) for 2010–2013. The average length of stay for psychiatric patients requiring transfer out of our ED was more than doubled in that time (from 5.6 hours in 2010 to 13.8 hours in 2013). It is not unusual for patients to wait more than 24 hours in the ED while providers attempt to locate an available psychiatric bed.

The primary problem is inadequate funding. The recent economic downturn forced states to cut approximately $4.35 billion in public mental health spending in the period between 2009–2012, the largest reduction since deinstitutionalization in 1960s. According to Steve Blanchard, the department administrator of UIHC Psychiatry, most hospitals run their inpatient psychiatric units at a deficit. Due to the chronic, disabling nature of mental illness, many patients seek care with coverage through government payers, which generally pay below cost. Duration of stay in an acute inpatient setting may be lengthened due to the inadequacy of outpatient community resources. In Iowa, the supply of inpatient psychiatric beds has continued to decline as inpatient facilities close due to lack of funding and retirement or exodus of Iowa psychiatrists to states with better reimbursement and more support staff. The problem of the inadequate supply of beds is exacerbated by the poor distribution of beds; many rural areas have no access to services. UIHC recently had a patient remain on one of the acute inpatient psychiatric units for 462 days because it was nearly impossible to locate a community-based option that could accept the patient, despite contacting more than 100 facilities.

While some of these problems may be specific to Iowa, they are symptomatic of the larger national crisis of lack of adequate mental health care. In the absence of promises for increased funding for mental health from state or federal sources, alternative solutions should be pursued. For example, telepsych, or the remote psychiatric evaluation, is one alternative being piloted at UIHC. Additionally, low-cost collaboration between EDs and community outpatient alternatives has been shown to decrease emergency department boarding. This collaboration could include using mental health clinicians to train ED staff in the management and care of patients with serious mental illnesses or having a social worker in the ED who can connect patients with community services at the time of discharge. The involvement of law enforcement may help. Federal grants of up to $250,000 over two years are available for the planning, implementation, or expansion of collaborative programs between criminal justice and mental health partners, including specialized training of law enforcement officers. As emergency care providers, whether in rural Iowa or inner-city New York, we are all impacted by the shortage of inpatient psychiatric beds. Cost, quality of care, ED throughput, and patient safety are all negatively impacted by this crisis. Solutions, such as collaborating with community mental health services, educating ED staff about the management of the boarding mental health patient, and using telemedicine are all viable strategies that will protect a subset of ED patients who often cannot advocate for themselves.

In the absence of the promises for increased funding for mental health from state or federal sources, alternative solutions should be pursued.

References

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Dr. Kitchen is an emergency physician resident R2 at the University of Iowa Hospital and Clinics in Iowa City.
Recently cared for a 35-year-old woman who presented to the emergency department for evaluation of palpitations. The symptoms lasted for 10 minutes and produced a mild sense of lightheadedness, but there was no chest pain, dyspnea, diaphoresis, syncope, or other typical cardiopulmonary symptoms. The patient reported that she had palpitations a few times in the prior month, and she had presented to another ED after the first episode. At that time, she had an electrocardiogram that was normal, and she had no further workup. She had no other medical problems, took no medications, and had no primary care physician. I was unable to identify any precipitants for the palpitations: no recent changes in diet, medications, illicit drug use, or stress and no use of tobacco, stimulants, or alcohol. Her physical exam, ECG, and electrolytes were completely normal.

The patient I described is not unusual to anyone working in the ED. We often see patients like this and debate the management. Given the absence of significant cardiopulmonary complaints, it would be difficult to justify admission, and even a 24-hour ED observation for cardiac monitoring is likely to be low-yield given the infrequency of her symptoms. My normal approach to this patient would be to recommend that she see her primary care physician or a cardiologist within a day for placement of a Holter monitor or event monitor, but given her lack of a primary care physician and the difficulty of obtaining a rapid appointment within our crowded system, I knew that I was not going to be able to help this patient find a quick diagnosis and treatment.

A Possible Solution?

A solution for scenarios like this may be on the way. The ZIO XT Patch is a single-channel continuous-recording ECG monitor, available by prescription, that can be worn up to 14 days by patients being evaluated for possible cardiac dysrhythmias. As stated in the product manual, “it is indicated for use on patients who may be asymptomatic or who may suffer from transient symptoms such as palpitations, shortness of breath, dizziness, lightheadedness, presyncope, syncope, fatigue, or anxiety.” There are no contraindications to its use.

The ZIO XT Patch was applied and worn for up to 14 days or until the patient had symptoms to trigger an event. The overall diagnostic yield for detection of a dysrhythmia was 63 percent. Almost half the patients (48 percent) were noted to have at least one significant dysrhythmia during their triggered events, indicating a nondysrhythmic cause of symptoms. The median time to detection of dysrhythmia was one day (interquartile range 0.2–2.8 days), and the median time to first symptomatic event was 1.5 days (interquartile range 0.4–6.7 days), suggesting that traditional 48-hour Holter monitors would have detected a majority, but not all, of the dysrhythmias.

The ZIO XT Patch offers a promising alternative to Holter or event monitors for the outpatient evaluation of patients with possible dysrhythmias. However, a major question comes to mind as I consider the future use of this device: given the potential for widespread availability of the device, will the ZIO XT Patch become yet another overused test in very low- or no-risk populations?

A major question comes to mind as I consider the future use of this device: given the potential for widespread availability of the device, will the ZIO XT Patch become yet another overused test in very low- or no-risk populations?

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**Reference**


**DR. MATTU** is professor and vice chair of the Department of Emergency Medicine and director of the Emergency Cardiology Fellowship at the University of Maryland School of Medicine in Baltimore.
Above 115 heartbeats per minute, fine motor control is compromised. Above 145, gross motor control is affected.

The Conspiracy of Stress: Part II

by RICHARD M. LEVITAN, MD, FACEP, AND MICHAEL ASKEN, PHD

“Too much time is dedicated to the acquisition of technique and too little to the preparation of the individual for participation.”
—Bruce Lee, martial artist, actor, director

“The probability that the total system will perform correctly is the probability that the hardware/software will perform correctly, times the probability that the operating environment will not degrade the system operation, times the probability that the user will perform correctly. By defining total system this way, human performance is identified as a component of the system.”

In many instances, the “difficult airway” is a relative term—relative to the operator. Early in my career, I recall missing an intubation. I panicked and called an anesthesiologist. Picking up the same instrument, she inserted the tracheal tube without difficulty. It would have made me feel much better if she had struggled with the tube, but alas, she made it look easy. For some time, I wondered what I did wrong and whether I could have done it better. For some time, I wondered what I did wrong and whether I could have done it better. Twenty years later, I was able to successfully intubate a patient in whom anesthesia missed the tube. The patient was shot in the central box; anesthesia had placed a tube, but by direct visualization of the lungs (thoracotomy in progress), it was clear the tube was not in the trachea. I picked up the same laryngoscope, came down the tongue, suctioned the mouth, identified the epiglottis, and intubated the patient as if it were easy.

Looking back on my multidecade obsession with the techniques of airway management, I realize in hindsight how much the individual’s mindset is critical to successful performance in crisis. Proper techniques are essential: patient positioning, the mechanics of mouth opening, epiglottoscopy (finding the epiglottis before making any attempt to expose the larynx), understanding the subtleties of epiglottis elevation, knowing laryngeal anatomy (even when partially viewed), and the nuances of tube insertion.

The operator’s mindset, however, is what allows for the proper application of techniques in the moment of crisis. It is one thing to know how something should be done but quite another to actually then pull it off in the real life-and-death, high-pressure situation. Related to the ultimate stress—fear—the Spartan commander Braxidas observed: “Fear makes men forget, and skill which cannot fight is useless.”

Fear Is the Mind Killer

We each have a genetic disposition to handle stress. Looking back at my initial years in emergency medicine, I now understand that my inherent adrenaline response made it very difficult for me to perform well. I got too stressed, and the adrenaline dump that ensued made it very difficult for me.

Like everyone I know in EM, I was never given psychological performance skills to maximize my response in crisis situations. I just assumed, and was led to believe by my teachers, that through some kind of desensitization, I would just get better. Over decades of practice I did improve, but recent interactions I have had with military personnel have now convinced me that we can train to do better by directly confronting the gorilla in the room and addressing the psychological aspects of procedural performance.

I am so convinced of this that I have added a psychologist, Michael Asken, PhD, my co-author on this article, to the faculty of my airway course in Baltimore. I now view every component of airway management with a different perspective. I think about how stress conspires to make us fail and how it must be handled at each step in the process.

“Fear is like fire. It can cook for you. It can heat your house. Or it can burn you down.”
—Cus D’Amato, boxing manager, trainer

Adrenaline increases our heart rate, dilates our blood vessels, and widens our pupils; it gets us ready for the increased physical demands of a fight-or-flight situation. Excess adrenaline, however, becomes very dangerous, especially when we are required to perform complex tasks (as opposed to just running away from a predator). In the procedural performance situation, the mismatch between the perceived demands of a task and one’s perceived abilities creates “performance stress.” When the mismatch is dramatic, the adrenal dump that occurs becomes detrimental. Above 115 heartbeats per minute, fine motor control is compromised. Above 145, gross motor control is affected. Time perception gets altered. Our ability to appreciate external cues (ie, listening, accurately observing the situation) becomes limited. Frozen by the stress, operators become “stuck on stupid,” repeating the same response over and over (even though it’s not working). The Brits jokingly refer to this as “wearing brown trousers” because in superstressful situations bowel control is compromised. Tactical operators emphasize the importance of the “battle crap” before beginning a mission. At a recent conference in Australia, I heard an EM doc refer to using bike clips with their brown trousers—now that’s stressed! While it was once believed that crisis functioning or mental toughness was the right stuff, something that you either had or did not have, we now know this is not the case. The military, professional and Olympic athletes, and police agencies have all recognized this and created psychological training programs to maximize performance in high-stress and life-threatening situations. Emergency medicine has lagged far behind in this critical area.

In the procedural performance challenge of emergency airway management, what can we do to manage our stress appropriately? We need to have the right mindset. By adjusting our perceptions (perceived demands versus abilities), we can reduce our overall performance stress. We need to consider and design our procedural (and team) processes in ways that recognize operator stress as a risk factor for error. When actually performing the procedures, we should factor in mechanics, ergonomics, lighting, and other environmental variables so we can do our best. We will address these solutions in future columns.
EDs, Clean Up Your Act

by SHARI WELCH, MD, FACEP

One of the most commonly observed features of the practice of emergency medicine is that workflow depends upon access to many and varying supplies. In any emergency department anywhere in the country right at this moment, myriad staff members (including physicians) can be observed scurrying around their departments, searching for and gathering supplies necessary for treating the next patient. Most EDs tolerate a level of clutter and disorganization that most providers would not tolerate in their own homes.

Kaizen is a Japanese workplace philosophy that means “improvement” and focuses on making small continual improvements that keep a business at the top of its field. The philosophy involves everyone in the organization, managers and leaders and workers alike, and urges them to make never-ending efforts for improvement. The foundation for Kaizen was laid after World War II when the country was attempting to rebuild infrastructure and rethink many systems. American leaders like W. Edwards Deming and Joseph Juran came to Japan to lecture and teach, and this led to the Training Within Industry (TWI) programs, which subsequently gave way to Kaizen in the 1950s. The philosophy has been central to the cultures of companies like Toyota and Canon, where suggestions are regularly solicited from each employee, written down, shared, and implemented.

Kaizen has five key principles:
1) There is heavy reliance on teamwork. Everyone’s opinion is valued and considered.
2) Workers have a strong personal discipline, and morale must improve under Kaizen.
3) Workers should be confident about offering suggestions, even when a system is already functioning adequately.
4) Kaizen recognizes that there is always room for improvement.
5) The system uses worker groups (also called quality circles) that meet and work together to both solve problems and come up with innovations.

This improvement model would seem particularly appropriate in the emergency department, which has many parallels to factory-floor processes. In emergency medicine, our product is urgent medical care, we work in teams to provide it, and many steps are involved in the delivery of that urgent care that have no intrinsic value to the patient (customer). Another key feature of Kaizen has been termed the Five S of Kaizen. This is a method for organizing a workplace, especially a shared workplace like an emergency department.

Seiri (Sort): Tidiness, keeping out only essential items out.
Seiton (Set in order): Orderliness, eliminating extra motion.
Seiso (Shine): Cleanliness, keeping the workplace clean.
Seiketsu (Standardize): Standardizing work practices.
Shituke (Self-discipline): Sustaining and maintaining discipline and reviewing standards.

Kaizen in the ED

In a perfect example of Kaizen principles at work (though it was never dubbed as such), Alan Weier, MD, medical director for the emergency department at Baylor Regional Medical Center at Plano in Texas, spearheaded a project that involved the physician partners of 10 facilities within the Baylor Health Care System. The medical directors met for a strategic planning session in 2011, and one of their initiatives involved the standardization of equipment at all of their sites. They had three main areas of focus; two involved high-risk procedural equipment (airway and critical procedures), and a third focused on wound care.

The ED council, which in effect served as a Kaizen quality circle, worked together to standardize the equipment for three important workflows in the ED: airway, critical procedures (chest tubes, invasive vascular access, lumbar puncture) and wound care (lacerations, incision and drainage). They developed master lists of equipment for each cart and purchased the carts and supplies at a reduced rate for all 10 campuses. These carts can be moved anywhere in the department, and this is in keeping with the new model for EDs in which any care can be rendered in any room. This standardization helped reduce clutter within the department and the need to run around the department in search of supplies for both high-acuity and high-frequency conditions within the department, standardized care. It also added discipline to the culture, and reduced costs.

The implementation phase for the change was just as important as the development of the carts. Dr. Weier and his team used staff meetings, education sessions, and checklists with photos as part of the introduction of the carts, which took place at all sites. Continual feedback from providers ensured that the fine-tuning of the carts and their contents would be ongoing. The ED staff and physicians are so convinced that these standardized carts have improved care and increased efficiency and value that they are in the process of developing other carts for use throughout the system.

Dr. Weier and his team presented this work at the ED Innovations 2014 conference held in Las Vegas February 19–21. It will be my pleasure to share with you more of the innovations that were showcased as posters at the conference. And apropos of the work done in Plano, I challenge you to return to your own department and “clean up your act!”

Left: One of Baylor Regional Medical Center at Plano’s ED carts.
Above: Organization system for airway supplies.
How Much Money Do You Need to Retire?

by JAMES M. DAHLE, MD, FACEP

Retirement rules of thumb say you need 20 times your annual income to retire. Twenty times a typical emergency physician income of $275,000 is $5.5 million, far more than the vast majority of physicians need to enjoy a wonderful retirement. The best way to figure out how large your nest egg needs to be in order to retire without having to worry about ever running out of money is to first determine your expenses in retirement and then determine if you have the resources to pay those expenses.

The Good News

The best estimate of your expenses in retirement is what you are spending just before retirement along with some common-sense adjustments. The good news for physicians is that the majority of their expenses may completely disappear upon reaching financial independence and retiring. Consider Table 1, an example of a physician making $300,000 and his pre-retirement and post-retirement expenses.

This particular physician finds that he only needs $68,492 per year, or 23 percent of his pre-retirement income, to maintain his standard of living. However, that 23 percent is by no means a rule of thumb and is highly individualized. You may find you only need 20 percent of your pre-retirement income, or perhaps you may need as much as 50 percent. However, it is unlikely that you will need the 70 to 80 percent that some financial planners estimate once you subtract your savings, insurance costs, payroll taxes, mortgage payments, and expenses related to your children.

The Bad News

There is also some bad news associated with retirement planning. Financial professionals use a concept called the safe withdrawal rate, which is the amount of money you can withdraw from a reasonable portfolio each year, adjusted to inflation, while expecting that portfolio to last throughout your retirement. Although this number varies slightly over time and no one can predict future market returns, most experts agree the number is somewhere around 4 percent. That means a portfolio of $1 million can safely support an income of only about $40,000 per year, adjusted upward each year for inflation. Using this number, you can quickly see that an annual income of $220,000 will require a portfolio of $5.5 million. To make matters worse, if all or most of that portfolio is in tax-deferred accounts like 401(k)s and traditional IRAs, the after-tax income will be even lower.

Other Income Sources

Other sources of income decrease the expenses your portfolio must support. The most common of these is Social Security. The Social Security Administration sends you a statement each year with an estimate of the income you will receive at your full retirement age. There are a few things to keep in mind when evaluating that figure. First, it assumes you will continue working until your full retirement age. If you retire early, such as at age 55, that number may be significantly lower. Social Security averages the highest 35 years of earnings in determining your payment. If you only work for 25 years, Social Security will use 10 years’ worth of $0 earnings to determine your payment. Second, delaying Social Security payments to age 70 is one of the best ways to insure against your own longevity. But if you plan on retiring at 50 or even 60, you will need a plan to bridge the gap to Social Security at age 70 (not to mention Medicare at age 65).

Third, while many fear that Social Security will disappear completely, this seems highly unlikely given the popularity of the program. However, changes to the program are inevitable and may include raising the retirement age, lowering payments, and/or increasing the amount of Social Security tax paid. The bottom line is you should expect Social Security to provide an income of $20,000 to $40,000, at least in the latter half of your retirement years. Other sources of income include pensions, the income of a spouse who continues to work after you retire, inheritances, and rental property. Each of these can be used to reduce the amount of income required from your portfolio.

Other Options

Once you have determined your expenses and matched them against other sources of income, you may find that your portfolio is not large enough to support your remaining needs. There are a couple of options to make up the difference, but both involve giving up control of assets. The first is to use a portion of your portfolio to purchase a single premium immediate annuity (SPIA). This is an insurance contract where you pay the insurance company a lump sum of money and, in exchange, the company pays you a set amount each year for the rest of your life. While many annuities are complicated high-expense products designed to be sold and not bought, SPIAs are a straightforward and competitively priced way to purchase a pension. Unlike life insurance, which becomes more expensive as you get older and sicker, SPIAs become less expensive as you age and develop illnesses. The major benefit of a SPIA is that, unlike portfolio withdrawals, the income is guaranteed (although when you die, your heirs do not receive anything). A SPIA purchased on a healthy 70-year-old male currently pays about 8.3 percent per year, more than twice as much as the safe withdrawal rate of 4 percent.

Another method of increasing income is to use a reverse mortgage. While this industry has been appropriately maligned for high fees and inappropriate sales practices, a reverse mortgage allows you to convert your home equity into income while staying in your home as long as you are able.

A better option for most doctors facing a retirement shortfall is to work a few more years. A few more years of work, even part-time work, can make a huge difference in your spending level in retirement. Working longer allows for more savings, more time for prior savings to compound, and fewer years in which your portfolio must support you. Fewer years of work could increase retirement income by 80 percent or more.

The nonfinancial aspects of retirement should not be ignored. Losing your identity as a practicing physician is difficult for many. Filling your time with worthwhile activities is hard for others. Your relationship with your spouse may also undergo a difficult adjustment when you work less. Emergency physicians and other shift workers are lucky in that they can often ease themselves into retirement by gradually reducing shifts, minimizing these issues compared to many specialists.

Appropriate retirement planning, done on your own or with an appropriate professional, will minimize financial worries in your later years.

Table 1. Expenses Before and After Retirement

<table>
<thead>
<tr>
<th>EXPENSE</th>
<th>PRE-RETIREMENT</th>
<th>POST-RETIREMENT</th>
</tr>
</thead>
<tbody>
<tr>
<td>Retirement savings</td>
<td>$60,000</td>
<td>$0</td>
</tr>
<tr>
<td>College savings</td>
<td>20,000</td>
<td>0</td>
</tr>
<tr>
<td>Child related expenses</td>
<td>30,000</td>
<td>0</td>
</tr>
<tr>
<td>Payroll taxes</td>
<td>23,208</td>
<td>0</td>
</tr>
<tr>
<td>Mortgage</td>
<td>36,000</td>
<td>0</td>
</tr>
<tr>
<td>Life insurance</td>
<td>1,500</td>
<td>0</td>
</tr>
<tr>
<td>Disability insurance</td>
<td>4,800</td>
<td>0</td>
</tr>
<tr>
<td>Commuting/work expenses</td>
<td>9,092</td>
<td>0</td>
</tr>
<tr>
<td>Income taxes</td>
<td>50,000</td>
<td>10,000</td>
</tr>
<tr>
<td>Charity</td>
<td>20,000</td>
<td>5,000</td>
</tr>
<tr>
<td>Health savings account/health care</td>
<td>8,000</td>
<td>2,000</td>
</tr>
<tr>
<td>Other insurance</td>
<td>12,000</td>
<td>12,000</td>
</tr>
<tr>
<td>Property taxes</td>
<td>5,000</td>
<td>5,000</td>
</tr>
<tr>
<td>Food</td>
<td>12,000</td>
<td>12,000</td>
</tr>
<tr>
<td>Vacations</td>
<td>5,000</td>
<td>15,000</td>
</tr>
<tr>
<td>Other items</td>
<td>7,492</td>
<td>7,492</td>
</tr>
<tr>
<td>TOTAL</td>
<td>300,000</td>
<td>68,492</td>
</tr>
</tbody>
</table>
By JEREMY SAMUEL FAUST, MD, MS, MA

The Free Open Access Medical Education movement (#FOAMed) often focuses its attention around the hottest topics in emergency medicine. Would you use a bougie when performing a cricothyrotomy or not? What is your preferred ratio of blood products for massive blood transfusion in trauma? How do you assess response to resuscitation in septic patients? These debates rage on in the Twitterverse, and if you want in on those conversations, I heartily recommend following the Twitter feed of PHARM (Prehospital & Retrieval Medicine) podcast creator and host Minh Le Cong, MBBS (@gfldoc). RFDS, as all Aussies but few Americans know, stands for Royal Flying Doctor Service. Dr. Le Cong seems to run a small ICU from his plane as he covers vast swaths of the Australian bush. While passing the time on long flights, he enjoys serving as a lightening rod in the #FOAMed conversation on Twitter, bringing his extensive knowledge and experience to these debates, along with his tenacity and good humor. From high-yield pearls to frequent links to new papers, his feed is certainly a busy one. So beware: following Dr. Le Cong is akin to drinking the Twitter Kool-Aid. You will learn a lot, but once you’ve followed him, there’s no turning back.

Increasingly, emergency medicine providers are using Twitter as a tool to disseminate more traditional bread-and-butter medical knowledge, the information found in those bounded collections of pages held together with glue and thread. Ah, yes...books.

This month, there were a number of tweets that referenced “traditional” medical education that caught my eye. The first came from the feed @Master_USMLE. This account is devoted to board-review pearls found in various review books and has amassed more than 53,000 followers—many being medical students and residents. The feed mainly consists of mnemonics that you may not remember and probably don’t need to. However, you might like the occasional EM-relevant entry. One recent standout: “Vertigo differential: VOMITs: Vestibulitis, Otoxic drugs, Ménière’s disease, Injury, Tumor, Spin (benign positional vertigo).” Not bad, but this tweet was missing something—the one cause of vertigo you simply can’t afford to miss: cerebellar stroke! I uploaded using VOMITs as a mnemonic for the differential diagnosis of vertigo but made this glaring omission known in my tweeted response. Mnemonics seem to work best when the acronym of the mnemonic is in some way associated with the medical problem it is used for. Vertigo tends to causes emesis, so you’re more likely to remember it and use it. But the real reason to follow @Master_USMLE is that your medical students probably read it, and no one wants to be pimped by their students!

More Twitter-based PR for old-fashioned book learning came from Michael Stone, MD, emergency ultrasound fellowship director at Brigham & Women’s Hospital in Boston (@bedsideonsen). Last month, Dr. Stone was busy tweeting a slew of high-yield pearls from his own boss, Ron Walls, MD, author of the Manual of Emergency Airway Management and chair of EM at “the Bej.” Dr. Walls’ checklist for the assessment of airway difficulty is second nature to many EM providers, but it’s always worth repeating: “Walls - LEMON, L

Rule of 50s to correct sugar: % dextrose x cc/kg=50. Adult gets D50 at 1cc/kg. Kid gets D25 at 2cc/kg. Infant gets D10 at 5cc/kg. Look externally (gestalt), E - evaluate 332 (that’s shorthand to say that in patients with “easier” airways, you should be able to fit three fingers between their incisors, the mandible length should be at least three-fingers wide, and the distance between the hyoid bone and the thyroid bone should be at least two-fingers wide), M - mallampati, O - obstruction/obesity, N - neck mobility.”

Pik Mukherji, MD, EM/IM attending at Long Island Jewish Medical Center in New Hyde Park, NY (@er-cowboy), seems to bask in his role as self-appointed FOAM skeptic and is known for his rhyming Twitter profile, “Devil’s Advocate (by choice and intent).” Offense (if given) Never Meant.” Dr. Mukherji also enjoys a reputation as a master educator. His points on Twitter are always succinct and relevant, like this excellent reminder for resuscitating hypoglycemic patients of all ages: “Rule of 50s to correct sugar: % dextrose x cc/kg=50. Adult gets D50 at 1cc/kg. Kid gets D25 at 2cc/kg. Infant gets D10 at 5cc/kg. #EMConf.”

The final entry for this month’s installment of “The Feed” doesn’t exactly fit the “traditional medical education” category, but it’s so good that I have to include it. From University of Maryland ED pharmacist and toxicologist and frequent Academic Life in Emergency Medicine (www.academicleifem.com) contributor—and arguably its MVP—Bryan Hayes, PharmD (@PharmERToGuy), comes, “The 2014 list of Oral Dosage Forms That Should Not Be Crushed. From @ismpl http://www.ismp.org/tools/donotcrush.pdf #FOAMed.” This online PDF from the Institute of Safe Medication Practices, a non-profit patient safety organization, contains a list of all medications that should not be crushed. For each entry, the list includes the active ingredient, the relevant formulation (tablet versus capsule, etc.), and a brief and precise reason the medication shouldn’t be crushed. Some of these are obvious and trivial (such as the advice to avoid crushing any extended-release formulation), while others are obscure yet important and downright fascinating. For example, did you know that you should never crush Celebrex (meclofenamate mefenyl, an immunosuppressive agent for transplant patients) because direct exposure to the active ingredient can enhance tumor production? I sure didn’t. In-sights like these are what cause so many of us to keep drinking from the endless fountain of FOAM.
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Ohio – Northeastern Ohio

Physicians Emergency Services, Inc. is a progressive, single hospital, independent democratic group seeking another BC/BE physician to join its team.

The hospital is located in Ravenna and has a 22 Bed ED with electronic medical record system. Annual census is 37,000. Competitive salary. Excellent benefit package. Equal shareholder at 2 years. Eight-hour shifts rotate amongst all physicians except two existing physicians work exclusively nights. ED Physician coverage is 40 hours per day and PA/NP coverage 20 hours per day.

A description of some our practice advantages along with a more detailed summary of our salary and benefit package is available.

For more information please contact Brian Adams, MD, FACEP 440-864-4242 or by email at phys_app@pesmed.com.
Award-winning care  
we do that here

Northwestern Medical Center is looking for a full-time physician to join our thriving Emergency Department team!

Our ideal candidate is BE/BC in Emergency Medicine, and will maintain ACLS certification. Our Emergency Department has 36 hours of physician and 18 hours of APP coverage daily, 7 days per week. Full-time physicians work 10 shifts per month, with some weekends and nights required. We register patients at the bedside, chart via NextGen EMR, have bedside ultrasound, and rarely board patients in the department.

We're blessed with our unique location in the northwest corner of Vermont. It gives us great access to hiking, the lake, skiing—any number of ways you can interact with nature. Working at NMC gives you the opportunity to work at an award-winning institution while having a great quality of life. Competitive compensation, including excellent benefits ($6,500 per year in CME plus paid time off for CME or vacation!). Relocation and education reimbursement negotiable.

For details, contact: Kimberly Rubinsak at 727-507-3631 or Kimberly.Rubinsak@EmCare.com
The University of Florida Department of Emergency Medicine is recruiting motivated & energetic emergency physicians to join our new UF Health – Northside Emergency Department in Jacksonville, Florida.

Live and play at the beach. Work and learn with academic colleagues on the cutting edge of simulation, ultrasound, advanced airway management, critical care and wellness. Be part of a growing and supportive academic faculty that will work to help you establish your professional goals.

UF Health – Northside will begin as a 28 bed full-service, free-standing emergency department with six observation beds. There will be comprehensive radiology and laboratory services, and consultation will be available from all UF Health specialty and sub-specialty services. Phase 2 of this project will include the addition of 99 inpatient beds to this facility. This is a rare opportunity to get in on the ground floor of an exciting project, and take care of patients in a beautiful, state-of-the-art emergency department.

Join the University of Florida Faculty and earn an extremely competitive community-based salary as a UF assistant or associate professor in a private practice setting. Enjoy the full range of University of Florida State benefits including sovereign immunity occurrence-type medical malpractice, health, life and disability insurance, sick leave, and a generous retirement package.

All physicians are ABEM / ABOEM Board Certified / Board Eligible.

E-mail your letter of interest and CV to Dr. Kelly Gray-Eurom
Kelly.grayeurom@jax.ufl.edu

EOE/AA Employer
Why Take the National Emergency Medicine Board Review Course?

The fact is that taking certifying or recertifying board examinations is a stressful and time-consuming experience. The sheer mass of information that needs to be reviewed, combined with the press of occupational and personal responsibilities makes finding the time to study very difficult. Even with adequate time to study, the volume of material to be studied is staggering: Rosen's 2006 edition is 3179 pages long and the latest edition of Tintinalli has 1917 pages. Bottom line – preparation for these exams can be a daunting process. The National Emergency Medicine Board Review was created 18 years ago to specifically address the needs of busy emergency physicians required to take their certification or recertification examinations and who wanted a highly focused, no-fluff course that delivers the information they need in a concentrated, high-yield manner.

If you Don't Pass, You Don't Pay! Period!* Plus Over $1000 in Free CME!

*That's right – 100% of your tuition refunded, plus the opportunity to attend selected future CME programs at no charge. There is no fine print and no administrative fees are withheld.

"Excellent – best I've attended in 30+ years"

"Learning should always be this easy and so much fun."

"Excellent educational opportunity. I highly recommend this course."

"Some of the best lecturers out there. It's hard to keep everyone's interest while reviewing the entire EM core curriculum, but you all pulled it off!"

"This course was a focused educational experience, and I could not think of a better way to prepare for the exam."

"Great News! I got a 93% on my ConCert Exam. The NEMBR course was instrumental. Ten years ago, after I took the course, I got a 94%."