Recently I had the pleasure of speaking with anesthesia residents and faculty at a well-known progressive academic anesthesiology department. Opportunities like this are among the high points of my professional life because I invariably know more when I leave these presentations than when I arrive. This time was no different.

My recent professional focus has been on working with hospitals and health systems to identify workflow enhancements and quality improvement initiatives to streamline care delivery and deliver greater total value. On a more theoretical level, I have been identifying and developing novel ways to produce comparable or better perioperative medical care in terms of price, quality, and service by using nontraditional processes or clinicians in nontraditional ways. With few exceptions, however, these latter efforts fall mainly into what one would call product development — showing promise but not yet ready for prime time.

The topic, then, for this visit was the role of disruptive innovation in the

Continued on page 4
What Defines Success in Today’s Healthcare Environment?

It is a privilege to bring you another article from Michael R. Hicks, MD, MBA, this time on Successfully Competing in Anesthesia Services Today. Throughout his career as an anesthesiologist and executive, Dr. Hicks has developed unique insights into the qualities that make for success. As a physician, he writes for his peers honestly and without trepidation. Dr. Hicks’s wisdom is among the most valuable information we have published in the Communiqué. In the current issue, he addresses anew the concept of disruptive innovation in anesthesia practice—but as he notes, “successful companies within the anesthesia space are still focused on implementing and executing sustaining innovations” such as quality and process improvement and “better management practices built upon fiscal and behavioral discipline.” The needed innovations will come from five different strategies identified by Dr. Hicks:

1. Actively manage the performance of the practice and its members, recognizing that neither the group nor its individual clinicians should be seen as commodities;
2. Seek ways to reduce your cost to your hospital, and in the process learn about operations management;
3. Embrace accountability and use your data;
4. Learn how to communicate. “Every interaction that we have in the perioperative process is at is essence a negotiation,” and
5. Continue developing your special professional skills.

Keep in mind, Dr. Hicks also counsels, that both direction and proper execution of a strategy are requisites for success. For the time being, given the state of the competition, execution can even trump strategy.

One way to lose to the competition is to fail to recognize it, according to our frequent contributor Mark F. Weiss, Esq. Mr. Weiss invites you to consider Are You Making This Mistake Concerning Competition? “This” mistake would be neglecting the threat from within the group. Other group members sometimes offer not only direct competition, e.g., by breaking off to form their own group, but they may also enable an outside group to take the place of the incumbent. It is important to build protective measures around the partnership/owner/employee relationship.

As anesthesia practice leaders, we have to stay on top of another “big C” compliance. Vicki Mykowiac, Esq. discusses the importance of preventive strategies in her article Anesthesia and Chronic Pain Compliance Risk Areas: Compliance Advice from Benjamin Franklin and Francis Bacon. If we listen, the government tells us clearly how to get into—and how to stay out of—trouble.

With the publication of the federal government’s final regulations on the HIPAA Privacy, Security and Breach Notification in January, 2013, we entered into a new era of HIPAA rights and responsibilities, which Neda M. Ryan, Esq. reviews in HIPAA Omnibus Rule: What Anesthesiologists Must Do Now. Christopher Ryan, Esq. focuses on one very important aspect of security—preventing loss of confidential information from cell phones and tablets—in Taking Security on the Road: Steps You Can Take to Secure Your Mobile Devices.

Finally, some less obvious compliance risks come from the new technologies themselves. Joette Derricks, CPC reviews the limitations of electronic health records and the erroneous documentation that they can easily engender in Health Information Management Challenges in the World of EHR.

One recurring topic that we have not touched upon in this issue is payment for anesthesia and pain medicine services. As we go to press, the two percent across-the-board cut to Medicare physician payments mandated by the federal budget sequester is set to begin on April 1, 2013, CMS confirmed in a recent announcement. We know by now that budget deals happen down to the wire, and we hope that our worries about sequestration will be moot by the time you read the Communiqué. Whatever happens, we can assure you that we will continue to keep you up to date.

With best wishes,

Tony Mira
President and CEO
**Are You Making This Mistake Concerning Competition?**

Mark F. Weiss, Esq.
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**“The Competition”**

These days I hear that term from more and more anesthesia group leaders, and I’m sure that you’re thinking about it more than you’d like.

From the Latin root competitio-nem, its meaning originated in the sense of rivalry, of a contest for something. Since at least the 1790s, it’s been used to describe rivalry in the marketplace.

Ask yourself what “the competition” means to you. What comes to mind?

In working with anesthesia group leaders across the country, my regular experience is that they envision the competition as another anesthesia group, whether from across the county or across the country. These days, the image that often first comes to mind is that of the predatory staffing-service model.

I certainly can’t fault these group leaders because, especially these days, there is tremendous competitive pressure from outside entities coveting your facility contracts.

So, for most group leaders protecting their practice from competitors involves looking outward to block outside groups from breaching the walls of your facility relationships.

Accordingly, the best of those group leaders would be doing all they could to create an Experience Monopoly, the unique experience that the group provides to its “customers” — hospitals, referring physicians and patients, one that, even if the competitors could observe what was going on, they couldn’t replicate. And, those group leaders would be taking other action to create barriers to block entry into their market by those outside competitors.

Of course, that focus on outside competitors and the action taken to keep them at bay and to render them ineffectual, are both necessary and required.

Unfortunately, if that’s all that’s done, your group is still at risk.

That’s because being so intently focused on outside competitors can blind group leaders from seeing another, often equally lethal, predator seeking to capture your group’s business; a predator so dangerous it might destroy your group’s ability to survive.

Let me use an example from outside of healthcare to illustrate this second category of threat.

A country can take great efforts to protect state military or even physical access to a sensitive installation by, among other things, erecting barriers, both physical and virtual, to outside intrusion. Similarly, industry spends countless millions of dollars each year protecting essential trade secrets from competitors.

But that’s only part of the story, because as much as a country’s military and as industry have to be concerned about protecting their secrets from someone who’s on the outside, they have to be equally concerned about guarding against espionage from someone who’s already on the inside — by someone who is thought to be “on the team.”

*Continued on page 7*
Successfully Competing in Anesthesia Services Today

Continued from page 1

shaping of anesthesia practices in the future and how the nature of competition among those providing anesthesia services might change with new entrants and funding mechanisms. Before venturing into the world of disruption and how things might look radically different, however, I felt strongly that the residents needed some background and explanation of the market as it currently exists so that the disruptive innovations I highlighted would have context as they were introduced to the audience. As a result, I began with an overview of both the business models of traditional private practice groups and the larger regional and national practices as well as some of the evolving global models for health care delivery that I see developing around the country.

As the presentation proceeded, a key discussion point centered on the drivers of capital flow into the specialty, particularly the sources of funds that would deliver the expected return on the invested capital. Very early a question arose as to what unique skills and competencies are possessed by large practices and private equity investors that lead to their belief that they will be successful and that the success will last. In other words, why do investors and owners of managed anesthesia practices believe that there is profit to be derived from what they do and where is the profit derived? Simply put, what is the nature of competition in anesthesia currently, what is strategic versus operational effectiveness, and what is necessary to be competitive today? Practices and management companies succeed today not with their ideas but with their execution.

Sustaining vs. Disruptive Innovation

At their core, essentially none of the competitive competencies being sold by anesthesia management companies are in the category of disruptive innovations. Instead, successful companies within the anesthesia space are still focused on implementing and executing sustaining innovations. These product offerings include “advances” such as increasing the adoption of the anesthesia care team model, better management practices built upon fiscal and behavioral discipline, increased alignment with institutional goals, embracing a culture of transparent outcomes measurement, and quality and process improvement that most other industries have long taken for granted.

These product attributes, while still, sadly, radical to many in anesthesia, are merely sustaining innovations and not disruptive.

The activities themselves are not even strategic in the business sense. Even the smallest of practices in the anesthesia marketplace have the ability to adopt and implement all of the aforementioned product offerings. In fact, these activities are more appropriately described as improvements in operational effectiveness. One view on the concept of operational effectiveness was developed by Michael Porter, a thought leader in the area of strategy and competitiveness, to differentiate activities that can be done by many (competing by doing things we both do, only better) from activities that can only be done by a few (strategic differentiation) (Porter, 1996).

At its essence Porter argues that few companies should be able to successfully compete solely on the basis of continual improvements in operational effectiveness because of the rapid dissemination of best practices. This is particularly true for process-driven specialties such as anesthesia. The fact that large regional and national companies are successfully competing for practices is less about their strategic differentiation than it is about the reticence of incumbent groups to adapt to changing times and make the difficult decisions needed when dealing with improving operational effectiveness.

What then, as the residents in the audience asked me, should they—and existing groups—do today to remain competitive in the current marketplace?

Strategies to Remain Competitive

First, actively manage the performance of the practice and its members with an emphasis on the latter. Unfortunately, many practices like to believe, and try to function as if, they offer a homogenous level of quality and service to their patients, surgeons, and facilities. They generally do not offer this, and most of the practices deep down know it. Ironically, most everyone else in the building knows it as well. Differing skill sets, attitudes, and motivations among clinicians mean that without some form of measurement and management being brought to bear, the care and service delivered will be uneven and predictably so. The ability to provide leadership, manage people and situations, and make difficult staffing decisions is
a large part of the product offering by successful practices.

One of the fundamental problems that groups face is an inability to make management decisions for the future that might cause some immediate pain. Some of this is due to generational issues within the field. Like me, many of the leaders of groups are older and have a relatively short timeline to retirement. The idea of investing in the future of the practice means incurring some financial pain now for a return to be delivered later. If your professional life timeline is measured in months or even a few years, as in the case of the leadership of many groups, there is no possibility of return down the road. In fact, this is a significant reason that many groups solicit acquisition offers now to the potential detriment of younger members of the practice.

Second, seek ways to lower the cost of your product before someone does it for you. Large regional practices and national companies get an audience for just a few reasons. Poor quality of anesthesia care is almost never one of them. If your practice receives subsidization I promise you that it is viewed as a cost on the hospital’s income statement and will be treated like one. Provide high quality services and the hospital may well be willing to pay you for them. However, you should still actively seek ways to reduce the financial burden on the hospital, if possible. If nothing else the exercise itself will educate the practice for the next difficult contract negotiating session and send a powerful message that you aren’t seeking a handout. This will involve developing familiarity with facility operations and workflow including activities outside of the operating rooms. Many processes and people have an effect on surgical patient flow and are drivers of cost for both the facility and your practice. The science behind these investigations is called operations management and, as an added benefit, knowledge of this branch of management will be part of the future of our specialty anyway. A useful beginning reference with practical tools is the work by Ronen (Ronen, Pliskin, & Pass, 2006).

Third, embrace the concept of non-punitive measurement, accountability, and the virtues of the quality improvement process as put forth by Deming, Berwick, James, and others (Berwick, 1989; Deming, 2000; James, 1989). Not only is it inevitable that increased individual transparency in terms of outcomes is coming to anesthesia, it is desirable. More important, it is already here. Unfortunately some of you don’t know it yet. If your hospital has an electronic health record, a surgical information management tool, or a pharmacy management system, then it has a record of many of your activities as well. For example, do your patients need more or less opioids than those of your associates? Is it possible that your nerve blocks are not as effective as your peers? Are pain scores for your patients recorded? Think there could be a correlation? Fortunately, I think, clinicians and institutions have been living in an era where health care has been data rich and knowledge poor. One of the current big ideas in the business world involves the concept of “big data” (McAfee & Brynjolfsson, 2012). Increasingly, technology is allowing the connecting of the myriad number of seemingly disparate data points and creating opportunities for evaluation, correlation and possibly even causation.

Instead of resisting it or putting it off until forced to accept it, offer to take ownership of the measurement and the management of the findings. Yes, it will be initially awkward, even painful, to learn objectively that not everyone in the practice has the same proficiency in the administration of regional anesthesia or engenders the same level of confidence from nursing and surgical colleagues. In reality, however, it is likely just a validation of what everyone including the hospital administration, surgeons, and nursing staff already know. There are a number of easily read resources to gain a working knowledge of these concepts (Carey & Lloyd, 1995; Provost & Murray, 2011).

Fourth, learn how to communicate and most importantly negotiate well. Every interaction that we have in the perioperative process is at its essence a negotiation. Conversations with surgeons and nursing staff about patient prepared-
SUCCESSFULLY COMPETING IN ANESTHESIA SERVICES TODAY

Continued from page 5

Note that the negotiation skills to which I refer are not positional tactics based on zero-sum scenarios with winner and loser outcomes, but the daily exchanges requiring integrative approaches to solving mutual problems that have risks, benefits, and tradeoffs. Here, desired outcomes are decisions based on collective needs and respect. Fortunately there are a number of resources available to gain some basic understanding of these concepts (Marcus, Dorn, & McNulty, 2011; Shell & Moussa, 2007).

Readers should note that those who excel at this form of communication also excel at both listening and understanding the perspective of those on the other end of the particular issue at hand.

Fifth, the role of the anesthesiologist is going to evolve over the next several years. Changing delivery and payment paradigms as well as the realities of disruptive innovations in perioperative medicine will create unique challenges as well as unique opportunities for the profession. My best advice for those in training or newly in practice, given from the perspective of someone long in the business as well as a potential employer, is to continue developing skills that cannot be easily replaced or are needed regardless of the delivery system changes that ensue. Leadership and communication skills, expertise with quality and process improvement, and maybe the most important, the skill and comfort managing the perioperative care of clinically challenging patients will be needed regardless of delivery system changes or disruptive innovations.

Finally, realize that knowing and doing are fundamentally different aspects of competition but both provide the fundamental answer to the questions posed by the residents on the nature of competition in the anesthesia business. If you don’t know what to do then execution is irrelevant. However, practices fail, even those run by sophisticated health systems or management companies, not for a lack of knowledge but for a lack of execution. When both practice leadership and line practitioners understand that their failures are for lack of execution and not for lack of direction the path forward becomes much more clear. In this case, and at this particular point, execution can trump strategy. How long this remains true, of course, is subject for another discussion.


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DAILY PROBLEM-SOLVING THROUGH NEGOTIATION

Readers should note that those who excel at this form of communication also excel at both listening and understanding the perspective of those on the other end of the particular issue at hand.

ness, scheduling of cases, or even the discussions of anesthetic options with patients and families are at their heart exercises in negotiation. Sadly, while business education should be a fundamental part of medical school and residency training, it remains largely unavailable to most medical students and residents. Furthermore, these skills are frequently poorly modeled for residents during training.

On a fundamental level this is a disservice to our specialty and to us. On a practical and competitive level excellence in negotiation is a fundamental skill for anesthesia practitioners.
Are You Making This Mistake Concerning Competition?

Continued from page 3

That same point applies to your anesthesia group.

Yet, unfortunately, protecting against competition from within is a weak point for most anesthesia groups. The reality is that many groups don’t fail simply because of competition from an outside competitor; they fail from within due to the actions of, and sometimes competition by, members of their own group who break off to directly compete with their former group or who facilitate an outside group’s ability to displace it.

Protecting against competition from within requires much more effort and detail than most groups incorporate within their “owner” documents (their shareholders agreement or partnership agreement) and within their employment agreements and subcontractor agreements.

Although in some states, the direct approach to preventing competition, covenants not to compete, is unenforceable, the correct approach is to build a series of protective measures against competition around the relationship between the group and each of its physicians, whether owners or employees/subcontractors. You can conceive of these protective measures as a series of interlocking spines or spears, each designed to provide protection. One particular measure standing alone might be compromised, but together, as a systematic structure, they provide far more potent protection.

By adopting a wide range of protective measures, groups reduce, and if possible, prevent, competition from within, whether it’s actual direct competition by group members or their facilitation of direct competition by a third party to whom those group members hand the key to the group’s economic engine.

Focusing on threats from without is hardwired into most anesthesiology group leaders. Group leaders must be just as diligent in focusing on threats from within.

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On January 25, 2013, the US Department of Health and Human Services (HHS) Office of Civil Rights (OCR) issued its long-awaited Health Insurance Portability and Accountability Act of 1996 (HIPAA) final omnibus regulations (Final Rule). The Final Rule modified the HIPAA Privacy, Security, Enforcement and Breach Notification Rules (HIPAA Rules) and is comprised of four sub-rules:

1. Final modifications to the HIPAA Privacy, Security, and Enforcement Rules mandated by the Health Information Technology for Economic and Clinical Health (HITECH) Act;
2. A final rule adopting changes to the HIPAA Enforcement Rule to incorporate the increased and tiered civil money penalty structure as set forth by HITECH;
3. A final Breach Notification rule;
4. A final rule modifying the Privacy Rule as required by the Genetic Information Nondiscrimination Act (GINA).

While the Final Rule is effective March 26, 2013, compliance with the provisions of the Final Rule is not required until September 23, 2013. This eight month window between the release date and the compliance date allows covered entities, including anesthesiologists, time to understand their roles under the Final Rule, and take action where necessary to ensure compliance by September 23. This article summarizes some of the key elements of the Final Rule applicable to anesthesiologists and their practices.

**BUSINESS ASSOCIATES**

The Final Rule renews a focus on business associates and their subcontractors, beginning with revising the definition of “business associate.” Business associates are now defined as those persons (other than members of the covered entity’s work force) or entities that perform certain functions or activities that involve the creating, receiving, maintaining or transmitting of protected health information (PHI) for a specified function or activity (e.g., claims processing or administration, data analysis, processing, or administration utilization review, quality assurance, patient safety activities, billing, benefit management, practice management, re-pricing). Additionally, the Final Regulations specify that a business associate includes the following:

- A health information organization, e-prescribing gateway, or other person or entity that provides data transmission services with respect to PHI to a covered entity and that requires access on a routine basis to such PHI;
- A person offering a personal health record to one or more individuals on behalf of a covered entity; and
- A subcontractor that creates, receives, maintains or transmits PHI on behalf of a covered entity.

Specifically excluded are healthcare providers to whom a covered entity discloses PHI for purposes of treating the individual, plan sponsors to whom a group health plan makes disclosures, a government agency determining eligibility for or enrollment in a government health plan, and a covered entity participating in an organized health care arrangement that performs certain functions.

Consistent with previous regulations and practice, covered entities must enter into Business Associate Agreements with business associates. The Business Associate Agreements must meet specific requirements, and set forth the parameters within which business associates may use and disclose PHI. Covered entities are not required to enter into direct agreements with subcontractors of their business associates. The responsibility has been placed on the business associates to ensure a contractual relationship exists between them and subcontractors that ensure compliance with the HIPAA Rules.

In line with the new focus on business associates and subcontractors, the Final Rule specifies that business associates and their subcontractors may be directly liable for certain Privacy and Security Rule violations. Therefore, busi-
ness associates and their subcontractors must ensure full compliance with HIPAA.

**Notice of Privacy Practices**

Covered entities must modify their Notice of Privacy Practices (Notice) to comply with changes in the Final Rule. Notice is required to communicate to the individual the ways in which the covered entity may use and disclose PHI, the covered entity's duties with respect to protection of the PHI and the individual's rights relative to his/her PHI. Typically, Notices must be delivered to patients not later than on their first encounter, and must be posted in a clear and prominent place on the covered entity's website.

The Final Rule does not require the Notice to include a list of all situations requiring authorization. Rather, the Notice must contain a statement indicating that most uses and disclosures of psychotherapy notes (where appropriate), uses and disclosures of PHI for marketing purposes, and disclosures that constitute a sale of PHI require authorization, as well as a statement that other uses and disclosures not described in the Notice will be made only with authorization from the individual. Moreover, if the covered entity intends to contact an individual for the purpose of fundraising for the covered entity, the Notice must contain a statement regarding fundraising communications and the individual’s right to opt out of receiving such communications. Finally, the Notice must contain a statement that affected individuals have the right to be notified following a breach of their unsecured PHI.

Pursuant to the changes in the Final Rule, anesthesiologists should review their Notices and update them as necessary to reflect the new requirements—or at least verify that the facility in whose Notice anesthesia is included is updated. Notices should also be updated on any websites that the practice may have. While existing patients do not need to receive a copy of the updated Notice, they must be made available upon request. As always, it is prudent to document in the patient's file when Notices are given to them.

**Individual Access to PHI**

Except for limited circumstances, individuals have the right to receive and review a copy of their PHI in a designated record set. With certain exceptions, a designated record set is made up of the records maintained by or for the covered entity that is used, in whole or part, to make decisions about that individual, or that is a provider's medical and billing records about that individual.

The Final Rule requires, for PHI maintained electronically, upon an individual’s request for an electronic copy of his/her PHI, the covered entity must provide that individual with access to the electronic information, in the electronic form and format requested by the individual, if it is readily producible. If the information is not readily producible, it must be delivered in a readable electronic format (e.g., MS Word or Excel, text, HTML, or text-based PDF) that is agreed to by the covered entity and the individual. Individuals must be given access to their records within 30 days of the request, regardless of whether the records are in paper or electronic format and whether paper records are stored off-site. Notwithstanding this timeframe, covered entities will have an opportunity for a one-time extension of 30 days.

Anesthesiologists maintaining electronic records should be reviewing their HIPAA policies to ensure they reflect this change in the HIPAA Rules. Moreover, anesthesiologists must ensure that they can grant patients’ access to their PHI within 30 days of the request.

**Requesting Restrictions on Uses and Disclosures**

Individuals have the right to request restrictions on the use and disclosure of their PHI for treatment, payment or operations (reasons for which a covered entity is generally not required to obtain authorization for the use and disclosure of an individual’s PHI), disclosures to those who are involved in the individual’s care or payment for care, or disclosures to family members. A covered entity is not under an obligation to grant this request; however, those covered entities agreeing to comply must abide by the restrictions.

The Final Rule expands the individual's right to request restrictions without the covered entity's right to deny the request. Specifically, for individuals who have paid the healthcare provider in full out-of-pocket, healthcare providers must grant requests to restrict disclosures to the individual’s health plan.

While certain uses and disclosures are required by law and thus cannot be circumvented by an individual requesting restrictions, anesthesiologists should review and revise their policies and procedures with respect to individuals’ access to their own PHI. Moreover, any forms used to process such requests must also be reviewed and revised, as necessary. For those anesthesiologists who have not been in the practice of granting restrictions, they must develop a process to comply with requests by private pay patients requesting restrictions on information disclosed to their health plans.
Breach Notification

In addition to the modifications listed above, the rules pertaining to breach notification were considerably amended. Prior to the Final Rule, “breach” was defined as a use or disclosure of PHI that posed a significant risk of financial, reputational, or other harm to the individual. A breach was presumed if the impermissible use or disclosure resulted in harm to the individual.

However, the standard by which “breach” is measured significantly changed under the Final Rule from a “risk of harm” standard to a “low probability that PHI has been compromised” standard. In other words, an impermissible use or disclosure of PHI is presumed to be a breach unless it has been demonstrated that there is a low probability that the PHI has been compromised. Therefore, breach notification is necessary in all situations, unless it is demonstrated that there is a low probability that the PHI has been compromised or an exception applies.

To determine whether the low probability standard has been met, the OCR set four factors that must be considered when performing a risk assessment:

a. The nature and extent of the PHI involved, including the types of identifiers and the likelihood of reidentification of the information;

b. The unauthorized person who impermissibly used the PHI or to whom the disclosure was made;

c. Whether the PHI was actually acquired or viewed or, alternatively, if only the opportunity existed for the information to be acquired or viewed; and

d. The extent to which the risk to the PHI has been mitigated.

Following consideration of the factors, the risk assessment must evaluate the overall probability that the PHI has been compromised.

In addition to revising the definition of breach, the requirement that the Secretary be notified of breaches involving fewer than 500 individuals was revised. Because some breaches may go undetected for long periods of time, notification must be made to the Secretary within 60 calendar days after the end of the year in which the breach was discovered.

Investigations and Penalties

The Final Rule requires the OCR to investigate any complaint of a HIPAA violation when a preliminary review of the facts indicates that there may be a violation due to willful neglect. Willful neglect is defined as conscious, intentional failure or reckless indifference to the obligation to comply with the provision violated. The OCR may exercise its discretion in conducting a compliance review or complaint investigation in instances where culpability may be less than a willful neglect.

Importantly, the Final Rule increases the Secretary’s discretion to choose between an informal and formal resolution of investigations or compliance reviews. This change allows the Secretary to impose civil monetary penalties without pursuing an informal resolution process (previously, an informal process was required to attempt to resolve issues involving noncompliance).

In 2009, the HIPAA tiered penalties were incorporated into the HIPAA Rules pursuant to HITECH. Violations of the HIPAA Rules could result in penalties of up to $1.5 million. In determining the amount of any civil monetary penalty, the Final Rule sets forth the following four factors to be considered:

1. The nature of the violation;
2. The nature and extent of the resulting harm;
3. The history of prior compliance with HIPAA; and
4. The financial condition of the covered entity or business associate.

Reality Check – HIPAA Enforcement is On the Rise

If the increased penalties and flexibility by the Secretary to impose the

| TABLE 1 |
|-----------------|-----------------|-----------------|
| Violation Category | Each Violation | All Such Violations of an Identical Provision in a Calendar Year |
| For violations in which it is established that the covered entity or business associate did not know and, by exercising reasonable diligence, would not have known that the covered entity violated a provision | $100-$50,000 | $1,500,000 |
| For a violation in which it is established that the violation was due to reasonable cause and not to willful neglect | $1,000-$50,000 | $1,500,000 |
| For a violation in which it is established that the violation was due to willful neglect and was timely corrected | $10,000-$50,000 | $1,500,000 |
| For a violation in which it is established that the violation was due to willful neglect and was NOT timely corrected | $50,000 | $1,500,000 |
penalties is not alarming enough, enforcement is a very real issue that many covered entities face. A sampling of some of the settlements that have occurred in the last year include the following:

- January 2, 2013 – A $50,000 settlement with a hospice, arising out of a stolen laptop containing over 400 patients’ PHI (notably, this settlement represents the first HIPAA breach settlement involving less than 500 individuals)
- June 26, 2012 – A $1.7 million settlement with Alaska Medicaid arising out of a report to HHS involving a stolen thumb drive containing PHI of more than 500 Alaska Medicaid beneficiaries
- May 24, 2012 – A $750,000 settlement with a Massachusetts hospital arising out of a report of disclosures made to the Attorney General regarding 473 unencrypted data tapes that were sent to a third party to be erased, but only one of the three boxes of tapes arrived
- April 17, 2012 – A $100,000 settlement with a physician group arising out of a report to HHS that the physician practice was posting clinical appointments for its patients on publicly accessible website calendar
- March 13, 2012 – A $1.5 million settlement with a private insurance company arising out of disclosures made under the Breach Notification Rule involving the theft of 57 unencrypted computer hard drives from a data closet

An in-depth review of the facts of each of these cases revealed that the main cause for the civil penalties was what the OCR found during the investigation. Most of the time, the investigation revealed significant deficiencies in compliance that may not have been directly related to the initial complaint.

In light of its recent pilot audit program and continuous press releases regarding settlements and penalties, the OCR is ramping up its HIPAA enforcement and no covered entity is immune from scrutiny.

**What You Can Do Now**

While the Final Rule does not significantly alter the way anesthesiologists have been operating under HIPAA in recent years, it does signal a need for groups to revisit their HIPAA policies and procedures, update them as necessary, and educate their workforce on those updates. The following are some specific steps anesthesia practices should be taking now:

- Review, revise, and update HIPAA policies and procedures as more specifically described in this article;
- Identify which relationships will fall under the definition of “business associate” and ensure that there is a Business Associate Agreement with that entity;
- For those relationships previously identified as being that of a business associate relationship, ensure the Business Associate Agreement complies with the updated regulations;
- Review and update, as necessary, the Notice to properly reflect new requirements of the Final Rule; and
- Ensure all members of your group and workforce are educated on the new requirements and policies, and be sure to document the date of the education and who attended.

While these recommendations are specific to the revisions in the Final Rule, all anesthesia groups should regularly engage in self-audits of their compliance with their HIPAA policies and procedures, update the HIPAA policies and procedures to reflect deficiencies discovered in audits, and regularly educate the group and its workforce on the HIPAA policies and procedures and any updates that have been made. Taking these steps and documenting them will best position a group if and when it is audited or investigated by the OCR.

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The creation of the Medicare/Medicaid Electronic Health Record (EHR) Incentive Program (commonly known as the “Meaningful Use Program”) gave physicians and hospitals a strong incentive to integrate EHRs into their practices. (For more information regarding Meaningful Use, see “Proposed Meaningful Use Stage 2 – What it Means to the Anesthesia and Pain Communities” published in the Spring 2012 issue of the Communiqué.) As part of their EHR system, many anesthesiologists have started using mobile devices such as laptops, tablets and smartphones. If used properly, these devices allow access to patients’ EHRs from anywhere that a WiFi connection (or cell phone signal) is available. This often results in quicker responses to questions from patients, families, and other providers. While the use of mobile technology has benefits, anesthesiologists choosing to utilize this technology must pay special attention to making sure they do so in a manner that conforms to their group’s or facility’s security policy and protects the privacy of the information.

This article will outline some of the various mobile security tools anesthesiologists can implement to aid in protecting their patient’s EHRs.

**Draft a Mobile Use Policy**

Anesthesia groups should develop and implement a mobile use policy, or include specific provisions in their security policy regarding mobile use. To develop a mobile use policy, the group must first decide whether it will allow its employees to access EHRs via mobile devices at all. Assuming this will be permitted in some fashion, the group must consider whether anesthesiologists will be permitted to use their personal mobile devices, or whether only “company owned” devices will be permitted to access secure information. Groups should also contemplate whether all mobile devices are permitted to access EHRs or whether access will be restricted to certain types of technology. For example, a group may decide that laptop computers are permitted to access EHRs, but tablets and mobile phones are not. Groups may also want to implement some of the various specific suggestions contained in this article. After an effective policy is drafted, the group should train its employees on the provisions of the policy and how they can achieve compliance with the same.

**Follow Your Organization’s Policy**

Reading and complying with the group’s or facility’s policy is the number one step anesthesiologists should take when implementing mobile technology and choosing which mobile security techniques to utilize. A group’s or facility’s policy may contain specific requirements that are not discussed or that differ from the items outlined in this article. Questions concerning a group’s or facility’s policy, or how to best secure a mobile device, should be directed to the group’s or facility’s Security Officer. Depending on the type of mobile device the anesthesiologist intends to use, the manner in which EHR is accessed, and the software the group or facility uses to store the EHRs, some of the items outlined below may not be applicable to all anesthesiologists. The Security Officer will assist the anesthesiologist in making sure he or she is using mobile technology in a manner that is compliant not only with the HIPAA Security Rule, but with the laws applicable in their specific jurisdiction.

**Physical Security**

Keeping mobile devices physically secure is the most obvious type of mobile security. Because mobile devices are, by definition, “mobile,” they are easily stolen or misplaced. While nobody can completely prevent their mobile devices from being stolen, everyone can take steps to decrease the likelihood of a theft. Instead of leaving a laptop on the back seat of a car, providers should consider locking it in the trunk or not leaving it in a car at all. Do not leave a tablet sitting on the table at the coffee shop; instead, bring it with you when you get a refill of your
coffee. If an anesthesiologist uses his or her cell phone to access patient information, he or she should not let their child borrow it on the weekend. Finally, if it is utilized in public areas, anesthesiologists should consider protecting the screen of their mobile device from being viewed by unauthorized individuals by using a privacy filter.

**Passwords**

Simply having a password to gain access to mobile devices is not enough. Providers need to make sure that they choose unique passwords that are not easy to guess. Studies have suggested that the most common passwords include “123456,” “password” and “iloveyou.” Common categories of passwords include using your telephone number, spouse’s name or pet’s name. These common passwords should be avoided because they are relatively easy to guess. Instead, anesthesiologists should use a password that is easy for them to remember, but hard for unauthorized users to guess. Generally, passwords should be at least six characters in length, and should include upper and lower case letters, one or more numbers, and one or more characters such as “!” , “#” or “@”.

Anesthesiologists should also remember that using the same password for all accounts means that if someone gains access to one account, he or she gains access to all accounts. Therefore, anesthesiologists should use unique passwords for each piece of software that allows them to access EHRs, change their passwords frequently, and never store passwords in insecure locations. For example, placing a sticky note on a laptop that says, “Password: ComMun!que2013ABC” renders an otherwise strong password virtually meaningless.

**Auto-Logoff or Timeout**

Most, if not all, mobile devices have built-in features that automatically log the user off (or lock the device) after a set amount of time of inactivity. Anesthesiologists should turn this feature on, and they should require a password to be entered in order to “wake” the device.

**Saving Information Locally**

Information may be stored on the mobile device itself, or it may be accessed remotely. The benefits of storing information remotely (i.e., not storing information on the device itself) is that the information is more likely to be up-to-date and require additional authentication to access the information beyond simply having access to the device. Some organizations may choose to allow anesthesiologists to store information locally on the device so that it can be accessed at any time without a connection to the internet. Having locally stored information means that if the anesthesiologist’s mobile device is lost or stolen, an unauthorized user may be able to obtain patient information with greater ease. (See “Remote Wipe” below). If information is stored locally, anesthesiologists should be sure to frequently back the information up to a secure server. Doing so means that if your device is misplaced or stolen, the information will not be lost.

**Remote Wipe**

Many mobile devices contain a feature that allows the owner to erase the memory or hard drive of the mobile device remotely in the event it is misplaced or stolen. Check with the manufacturer of your device to learn more about whether your device contains this feature, and if it does, make sure it is set up and ready to be activated. If it does not, talk to your Security Officer and consider investing in software that allows this capability.

**Firewall/Virus Scan**

A firewall is a tool that monitors incoming and outgoing activity and blocks certain transmissions according to the user’s specifications. For example, a firewall may be programed to prevent file sharing. Virus scanning software is designed to identify potentially harmful files and quarantine or delete them as necessary. Both of these tools should be utilized by anesthesiologists, and importantly, must be kept up to date.

**Where to Go for More Information**

Utilizing mobile devices in a medical setting improves patient care by allowing anesthesiologists to quickly access patient information from anywhere. In the event a mobile device is stolen or misplaced, or if an anesthesiologist feels his or her mobile device’s security may have been compromised, they should immediately contact their organization’s Security Officer. Providers can also visit www.healthit.gov for more information about implementing health information technology, or contact a qualified attorney.

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Benjamin Franklin said, “An ounce of prevention is worth a pound of cure.” It is certain that Mr. Franklin was not speaking about the value of preemptive compliance work, yet the old adage aptly applies to the work done by physician groups to prevent allegations of fraud or abuse.

The Office of Inspector General for the Department of Health and Human Services (“OIG”) recently reported that the government expected to set a record of $6.9 billion in recoveries from its investigations and enforcement actions for its fiscal year 2012. As the chart in Figure 1 shows, this $6.9 billion is part of a trend of continuously increasing recoveries.

For this reason, many physician groups have implemented compliance programs designed to minimize the chances that the group will commit what the government perceives to be fraud or abuse. One key to effective compliance is an understanding of those issues of particular importance to the government.

There are many ways that the government signals areas of interest for particular specialties. This article will focus on the areas identified for review that are relevant to anesthesia and chronic pain practices in the OIG Work Plan and the Recovery Audit Contractor Program.

**THE OIG WORK PLAN**

Each year the OIG publishes a Work Plan that is the culmination of work done throughout the previous year to: (1) assess relative risks in the programs for which the OIG has oversight authority; (2) identify the areas most in need of attention; and (3) set priorities for the sequence and proportion of resources to be allocated in the upcoming year(s). Compliance-savvy groups view the Work Plan as a roadmap to ongoing OIG focus areas for fraud and abuse.

**Anesthesia Focus Area**

Traditionally anesthesia has not been prominently included in the Work Plan. However, the 2013 Work Plan includes OIG plans to review anesthesia modifiers reported for the level of service provided.

We will review Medicare Part B claims for personally performed anesthesia services to determine whether they were supported in accordance with Medicare requirements. We will also determine whether Medicare payments for anesthesiologist services reported on a claim with the “AA” service code modifier met Medicare requirements. Physicians report the appropriate anesthesia modifier to denote whether the service was

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personally performed or medically directed. The service code “AA” modifier is used for anesthesia services personally performed by an anesthesiologist, and the “QK” modifier is used for medical direction of two, three, or four concurrent anesthesia procedures by an anesthesiologist. The QK modifier limits payment at 50 percent of the Medicare-allowed amount for personally performed services claimed with the AA modifier. Payments to any service provider are precluded unless the provider has furnished the information necessary to determine the amounts due.\(^4\)

**Chronic Pain Practice Focus Areas**

Unlike anesthesia, issues relating to chronic pain practices have been included almost yearly in the Work Plan. For example, in the 2008 Work Plan the OIG directly targeted pain practices:

Interventional pain management procedures consist of minimally invasive procedures, such as needle placement of drugs in targeted areas, ablation of targeted nerves, and some surgical techniques. Many clinicians believe that these procedures are useful in diagnosing and treating chronic, localized pain that does not respond well to other treatments. Interventional pain management is a relatively new and growing medical specialty. In 2005, Medicare paid nearly $2 billion for these procedures. We will determine the appropriateness of Medicare payments for interventional pain management procedures and assess the oversight of these procedures.\(^5\)

Likewise, in 2010 the OIG Work Plan specifically singled out payment for transforaminal epidural injections:

Transforaminal epidural injections are used as an interventional technique to diagnose or treat back problems, such as pain that starts in the back and radiates down the leg. … Medicare Part B physician claims for transforaminal epidural injections increased by 130 percent between 2003 and 2007.

Based on these statistics, the OIG indicated it would review Medicare claims to determine the appropriateness of Medicare Part B payments for transforaminal epidural injections and determine whether there were policies and safeguards to prevent inappropriate payments for transforaminal epidural injections.\(^6\)

While issues raised in the 2013 Work Plan are not specific to pain alone, there are a number of issues that will impact pain practices including:

- **Questionable Billing for Electrodiagnostic Testing:** The OIG will determine the extent to which Medicare utilization rates differ by provider specialty, diagnosis, and geographic area for needle electromyogram and nerve conduction testing.

- **Place-of-Service Coding Errors:** The OIG will review physician coding for services performed in ambulatory surgical centers and hospital outpatient departments to determine whether they properly coded the place of service. Medicare pays a physician a higher amount when a service is performed in a nonfacility setting, such as a physician’s office, than it does when the service is performed in a hospital outpatient department or, with certain exceptions, in an ambulatory surgical center.

- **Evaluation and Management Services/EHR issues:** The OIG will review multiple E/M services for the same providers and beneficiaries to identify electronic health records (EHR) documentation practices associated with potentially improper payments. Medicare contractors have noted an increased frequency of medical records with identical documentation across services. Medicare requires providers to select the code for the service on the basis of the content of the service and have documentation to support the level of service reported.

- **Evaluation and Management Services — Use of Modifiers During the Global Surgery Period:** The OIG will review the appropriateness of the use of certain claims modifier codes during the global surgery period and determine whether Medicare payments for claims with modifiers used during such a period were in accordance with Medicare requirements. Prior OIG work found that improper use of modifiers during the global surgery period resulted in inappropriate payments. The global surgery payment HHS includes a surgical


Continued on page 16
service and related preoperative and postoperative E/M services provided during the global surgery period.7

Practical Advice

Compliance Officers for anesthesia and pain practices should carefully review the OIG Work Plan for 2013 to determine if their practices are providing services included in the OIG focus areas. For practices that are providing services in an OIG focus area, the chance of being the subject of a Medicare audit or other administrative review is increased. Therefore, Compliance Officers should conduct audits of services provided within the OIG focus areas to ensure that the medical record documentation is complete and accurate, the medical record documentation supports the billed claim, and the services were provided in a manner consistent with Medicare policy.

Recovery Audit Program (RAC) Audits

The RAC program’s mission is to identify and reduce improper Medicare payments through detection and collection of overpayments coupled with the implementation of actions to prevent future improper payments.8 The program is carried by four (4) private companies under contract with the government to conduct post-payment audits. The companies are:

- **Region A: Performant Recovery** (CT, DE, DC, ME, MD, MA, NH, NJ, NY, PA, RI and VT)9
- **Region B: CGI Federal Inc.** (IL, IN, KY, MI, MN, OH and WI)10
- **Region C: Connolly, Inc.** (AL, AR, CO, FL, GA, LA, MS, NM, NC, OK, SC, TN, TX, VA, WV, Puerto Rico and U.S. Virgin Islands)11
- **Region D: HealthDataInsights, Inc.** (AK, AZ, CA, HI, ID, IA, KS, MO, MT, ND, NE, NV, OR, SD, UT, WA, WY, Guam, American Samoa and Northern Marianas)12

These RAC audit contractors must submit issues to the government for approval and must then identify issues that will be subject to audit on their websites.

Anesthesia Focus Areas

Connolly, Performant and HealthDataInsights have published the following anesthesia issues for audit:

- Anesthesia–CRNA or anesthesiologist overpaid: “Anesthesia provided by a CRNA (Certified Registered Nurse Anesthetist) and Anesthesiologist (physician) without a 50% cutback as per Medicare guidelines involving CRNA’s supervised by anesthesiologists.” (Connolly)
- Anesthesia care package E/M Services: “Under the NCCI Edit rules, the anesthesia care package consists of preoperative evaluation, standard preparation and monitoring services, administration of anesthesia, and post-anesthesia care. Anesthesia CPT codes 00100 to 01999 include Evaluation and Management (E&M) services rendered on the same day of the anesthesia procedure. If the only service provided in management of epidural/subarachnoid drug administration, then an E&M service should not be reported in addition to CPT code 01996.” (HealthDataInsights and Performant)

Pain Focus Areas

The RAC auditors have published numerous issues relevant to chronic pain practices:

- Inappropriate payments for transforaminal epidural injections: “Local Coverage Determination policy has indicated specific conditions or diagnoses that are covered for Transformational Epidural Injections. Carrier claims have been identified where the first-listed and/or other diagnosis codes do not match to the covered diagnosis codes in the LCD policies.” (Connolly)
- Trigger point injections – excessive units: “Only one Trigger Point Injection CPT Code can be billed per date of service.” (CGI Federal)
- Excessive units of facet joint blocks: “CPT Codes 64492 and 64495 should only be billed once per date of service.” (CGI Federal)
- Facet joints denervation billed without guidance: “In accordance with [local coverage determinations], Facet Joint Denervation requires placement of a needle in the facet joint under fluoroscopic or CT guidance.” (CGI Federal)
- Transforaminal epidural injection billed with guidance: “Per CPT Manual 2011, CPT Codes 77001 – 77003 and 77012, are not to be reported with CPT Codes 64479, 64480, 64483 and 64484.” (CGI Federal)
• Place of service errors for physician claims for services performed in an ASC or outpatient hospital: “We will review physician coding the place of service on claims for services performed in ambulatory surgical centers (ASC) and hospital outpatient departments. Federal regulations provide for different levels of payments to physicians depending on where the services are performed. Medicare pays a physician a higher amount when a service is performed in a non-facility setting, such as a physician’s office (POS 11) than it does when the service is performed in a hospital outpatient department (POS 22) or, with certain exceptions, in an ASC (POS 24).” (Connolly, CGI Federal, and HealthDataInsights)

• New patient visits: “Identification of overpayments relating to the same provider group and specialty billing more than one new patient Evaluation and Management service within a 3 year period of time.” (Performant, Connolly, CGI Federal, and HealthDataInsights)

Practical Advice

Compliance Officers for anesthesia and pain practices should carefully review the website for their RAC auditing company to determine if their practices are providing services in identified audit areas. For practices that are providing services under review, the chance of being the subject of a RAC or other Medicare audit is increased. Therefore, Compliance Officers should conduct focused audits of services provided within the RAC audit areas to ensure that the medical record documentation is complete and accurate, the medical record documentation supports the billed claim, and the services were provided in a manner consistent with Medicare policy. Moreover, Compliance Officers should check the website for their RAC contractor on a regular basis to identify additional relevant review areas that may arise.

Conclusion

Francis Bacon first stated that “knowledge is power.” While it is certain that Bacon was not speaking about the value of knowing about government fraud and abuse focus areas, the statement aptly applies to the preemptive compliance work that can be done by physician groups that understand government concerns. Compliance Officers for anesthesia and pain groups would do well to live by the advice of Mr. Franklin and Bacon: know which areas the government is interested in and take practical steps to ensure that the group prevents negative government audit audit results and inquiries through proper documentation and billing for services.

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There is no doubt that the emergence of the Electronic Health Record (EHR) program is changing the way providers capture documentation on the front end. According to the 2012 NCHS Data Brief, 55 percent of physician groups have already adopted an EHR. Among the 45 percent that have yet to implement an EHR system, nearly half plan to purchase or use a system already purchased this year. Hospitals are also purchasing and installing EHRs at a rapid rate.

EHR templates are rapidly gaining footholds despite some growing pains. CMS has issued advice on the use of checkboxes and drop-down menus accommodating discrete data capture. Despite access to such “documentation tools” via point-and-click templates, most physicians are complaining that it takes longer to document an encounter in an EHR than to previously dictate it. The RAND Corporation released a paper describing the phenomenon that occurs when an industry’s technological capabilities improve at such a dramatic pace that end-user productivity actually drops. Known as the IT productivity paradox, it demonstrates a link between poor design and usability that undermines productivity gains.1

To increase productivity with EHRs, many vendors are now offering speech options within their systems. Other physicians unhappy with the technology have given up and moved to medical scribes or documentation assistants to deal with “feeding” the EHR.

Regardless of how the information gets into the EHR, once it is there it can present challenges to health information management (HIM) specialists, including coding and auditing personnel. Universally, HIM departments and coding/auditing specialists, along with the provider community, have noted the increase in volume of computer-generated records. A previous encounter in an outpatient setting that may have required a half-page report to capture the pertinent clinical data now is five or more pages of information. With the volume of information now flooding HIM departments, coders and auditors are faced with some serious issues regarding how to make sense of it all. The key question they face from a coding and auditing perspective is, “What is relevant to this particular patient’s encounter?”

Many coding consultants and trainers have long taught physicians it is not the volume of the note but the quality of it that counts. Ever since the 1995 Evaluation and Management Services (E & M) Documentation Guidelines came out, coders, and payers too, knew that physicians could over document by providing unnecessary information to pad the note. Using an EHR in some respects makes it easier; however, such padding often occurred when dictation and transcript services were state-of-the-art. Physicians often relied on “canned” text that their transcriptions would insert based on a few words, e.g. “normal adult male exam” now they merely double-click the option. Perhaps the CMS advisement regarding this type of over documentation is rooted not in the use of templates by the physician per se, but in how the coder, or the billing system if an auto-coder is imbedded with the EHR, uses the information to bill the service.

In working with coders and auditors, it is often best to have them first focus on the presenting problem and the management or treatment option presented in the documentation. This advice is in agreement with CMS and many other payers that have gone on record regarding the importance of the medical decision making guidelines in the selection of the E & M code. Medicare’s definition of medical necessity requires that paid services meet but not exceed the patient’s medical needs and be provided in accordance with accepted standards of medical practice. Accordingly, Medicare carriers state that the patient’s condition

1 NCHS Data Brief, No.98, National Center for Health Statistics, 2012
(e.g., severity, acuity, number of medical problems) is the key determinant for the frequency and intensity of E & M services for which Medicare pays. Coding E & M services first on the basis of medical necessity followed by verification of documentation of required key work components for the selected code allows coders and clinicians to avoid several common pitfalls of E & M coding. WPS Medicare’s website states, “Providers can ensure accurate Medicare payments with correct documentation of MDM for E/M services. Medical decision making is generally easier for an already diagnosed problem than for an undiagnosed one. In addition, problems which are improving or resolving are less complex than those which are worsening or failing to change. Keep in mind that MDM should reflect the nature of the presenting problem. Treatment for a common ailment, such as an ordinary cold, will not usually warrant a comprehensive level exam."

Other challenges regarding EHR usage for HIM professional include:

- Identifying and correcting inconsistencies within the same note. For example, the physician would indicate the patient’s pain level was at a level 6 in the history of present illness and in the exam documentation that the patient was in no pain.
- Weeding out errors attributed to default functions. One large New York health system has numerous patients with family members without a heart. An ongoing erroneous default in the EHR design results in this condition when the physician does not add family history during the encounter.
- Resolving through physician queries and purging outdated or “perpetuating” data such as a medical problem long since resolved as still being unresolved or a medication as being current long after it was stopped. These errors are often due to the cut-and-paste or carry forward function. While challenging to the coder, a greater concern is the risk of harm to the patient by including outdated information in the encounter documentation. This type of error also can contribute to a “got you” malpractice issue when the physician did or did not address or treat the condition.

One final challenge on the HIM department’s plate now that will rapidly grab the attention of all physicians is the upcoming implementation of ICD-10. Known as a super-granulated coding system, ICD-10 will require some massive rewriting of EHR templates to ensure sufficient documentation is present to code the service. While point and click technology is presenting challenges today to EHR productivity, some consultants are anticipating the detailed documentation required to select an ICD-10 code may decrease productivity by up to 50 percent for both physicians and HIM professionals come October 1, 2014.

Watch for future articles in the Communiqué addressing how physicians may best prepare for the upcoming ICD-10 challenges.

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Join the anesthesiologists who have already fulfilled the first reporting period of Meaningful Use and have collected over $5,000,000 from CMS; and the thousands more who stand to collect a possible $25,000,000 during 2013. **FIRSTUse** is the first complete electronic health record (EHR) platform built exclusively for anesthesiologists and pain management specialists to easily satisfy Stage 1 of Meaningful Use as required to earn the Medicare EHR incentive payment. **FIRSTUse** is entirely web-based; you don’t even need to have an existing EMR in place. **FIRSTUse** combines EHR technology with a Personal Health Record (PHR) system. All management activities are handled by ABC, all you have to do is enroll in the meaningful use program and then contact us. Email meaningful.use@anesthesiallc.com today.

### Professional Events

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<th>Date</th>
<th>Event</th>
<th>Location</th>
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<td>May 11, 2013</td>
<td>Idaho Society of Anesthesiologists Annual Meeting</td>
<td>Hampton Inn Boise, ID</td>
<td>Sheri Sass <a href="mailto:shersass@gmail.com">shersass@gmail.com</a></td>
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<td>May 17 – 18, 2013</td>
<td>Advance Techniques for Acute and Chronic Pain Management</td>
<td>Motor City Casino Hotel Detroit, MI</td>
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<td>June 7 – 9, 2013</td>
<td>Florida Society of Anesthesiologists 2013 Annual Meeting</td>
<td>The Breakers Resort &amp; Spa Palm Beach, FL</td>
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