



Electronic records mandates may clash with privacy laws

Patchwork of state laws adds complexity.

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President Obama has declared that electronic medical records will “reduce error rates, reduce our long-term cost of health care and create jobs.” “Obama’s Prime Time Press Briefing,” N.Y. Times, Feb. 9, 2009. Congress has authorized \$19 billion to implement provisions of the American Reinvestment and Recovery Act of 2009 intended to accelerate the adoption and use of “certified electronic health record [EHR] technology” during the next several years by hospitals and physicians that provide services to Medicare and Medicaid beneficiaries. H.R. 1, 111th Cong. §§ 4001-4201 (2009). Professionals and hospitals that fail to implement EHR technology by 2014 stand to suffer reductions in Medicare reimbursements.

The goal of this campaign is to adopt EHR technology to replace the current paper and fragmented computer files maintained by the vast majority of hospitals and physicians. Imagine a health information technology (HIT) system that includes all of a patient’s diagnoses, medical history, laboratory and test results, medications prescribed, payor claims data, hospital records and other pertinent data. That system would be available to a patient’s

health plan, hospital, pharmacy and doctors. Payors and regulators also can use this type of system to reduce fraud, waste and duplication, as well as control processing costs and improve disease-state management programs. EHR technology promises to reduce medication and other medical errors and streamline clinical decision-making and communication.

That “holy grail” has been envisioned by many participants in the health care industry today, but unfortunately it is not achievable under the current patchwork of federal and state laws and most existing HIT systems. In its ambitious effort to hasten the advent of EHR for the 21st century, the federal government actually may be working at cross-purposes with privacy protections established under federal and state law. Here’s why.

CONFIDENTIALITY MANDATES

First, existing law mandates the daunting task of obtaining individual patient consents. Health information of a particularly sensitive nature, such as records concerning an individual’s treatment for mental illness, drug addiction or alcohol abuse, creates uniquely complicated legal and practical problems with respect to

interoperable EHR technology. For example, federal law protects the confidentiality of records regarding the identity, diagnosis and treatment of any patient if such records are maintained in connection with an alcohol or drug abuse treatment program that is regulated or directly or indirectly assisted by a federal program. 42 U.S.C. 290dd-2. With few exceptions, such health records cannot be disclosed, even among health care providers for purposes concerning medical treatment, without the patient’s prior written consent. § 290dd-2(b).

Consequently, absent a patient’s prior authorization, clinical laboratory test results produced by a federally funded hospital program that indicate or reveal the patient’s treatment for drug addiction apparently have to be segregated or omitted from any other information entered by the hospital in the patient’s EHR, or otherwise shielded from disclosure to other health care providers. Of course, the issue of segregating and omitting such information from the patient’s health record predated the advent of EHR, but it is precisely the integrated and cumulative nature of EHR that necessitates additional security and privacy measures to prevent the unauthorized disclosure of sensitive health records.

