

Health Information Technology Oversight Council

Consent Implementation Subcommittee

Formed: October 2011

Meeting schedule: 1-2 times per month, for 2-4 months (to be adjusted as needed)

Purpose: Operationalize the opt-out consent policy

Objectives:

1. To help bring clarity and understanding to the operational questions and issues raised during the public comment process regarding the former proposed administrative rules to implement the opt-out consent policy
2. To help identify any additional operational questions or issues that need to be addressed
3. To help in re-drafting language to be included in future administrative rules to implement the opt-out consent policy

Membership: Representation from the Legal & Policy Workgroup, the Consumer Advisory Panel, and targeted community organizations and expertise

Some of the specific questions this group will discuss and provide input on:

1. What kinds of health information exchange should be subject to the opt-out consent policy regulations?
 - a. How should “health information exchange” be defined for the purposes of these rules? (for example, Direct messaging or “push” HIE, versus query or repository-based HIE)
 - b. Which types of data sharing relationships or scenarios should be subject to the rules? (for example, within versus between health systems or affiliated organizations)
 - c. Which types of health care providers should be subject to the rules?
2. What are the most efficient and effective methods for providing a meaningful and informed choice to a patient to opt-out of health information exchange?
3. What are the technical limitations that providers face in implementing an opt-out? (for example, which EHR/HIE systems are designed to share patient data automatically between a patient’s care providers?)