Barriers	General solutions
CDS capabilities of	Determine core capabilities
existing eRx products	Publish practical recommendations and design concepts to
	reduce vendor rework
Usability of systems	Sharing of best practices and lessons learned
and of CDS modules	Bibliography/reading list provided by industry thought
	leadership groups and/or certification organizations
Access to patient data	Increased data availability with appropriate protections
needed to support CDS	Integration of eRx with EHR
	Clear pathway from eRx to EHR – through same or different
	vendors
Access to best CDS	Accessible published/stored knowledge
knowledge for all	Practical standard representations of knowledge and content
products	Knowledge acquisition and execution tools
Local management and	Practical organizational models for development, selection, and
maintenance of	updating of rules, content, and interventions
knowledge	Tools to select / extract / customize knowledge

Table 5. Barriers to widespread effective use of CDS in eRx.

Lack of standards for	Creation and acceptance of practical standards
Lack of standards for	Creation and acceptance of practical standards
dictionaries, data, sigs,	Endorsement of standards by government agencies and key
etc., increases cost,	stakeholder organizations
variability, and error	Industry collaboration and financial support for standards
	programs
Cost and difficulty of	Financial support programs
implementation	Revolving loans
	Removing barriers on support programs (e.g., Stark)
	Development of systems that are easier to implement and
	configure
	Implementation guides, templates, and toolkits
Cost of use	Ongoing reimbursement differential
	Pay-for-performance programs
Difficulty in	Standard classifications and common definitions for CDS
recognizing value	elements, to improve generalizability of research on CDS
	methods
	Educational forums, references, and websites
	Increased publication of results
Perception of	Clearly-stated liability considerations
increased liability if	Appropriate liability protections and safe harbors
CDS advice is rejected	Education

Appendix- Whitepaper People and Process

During the summer of 2004, ONCHIT expressed interest in obtaining expert input to help guide Federal Government activities concerning CDS in electronic prescribing and related domains. ONCHIT approached the Health Information and Management Systems Society (HIMSS) with the request to collect this expert input and produce a whitepaper. The HIMSS CDS Workgroup had recently published a detailed guide for CDS implementers, and was actively working on a second edition of that resource and related initiatives. Concurrently, the American Medical Informatics Association (AMIA) had struck an agreement with the Agency for Healthcare Research and Quality to produce a series of whitepapers on various topics; CDS was already slated to be one of the topics covered in that series. All members of the HIMSS workgroup are active members of AMIA as well. It was agreed by all participants that the CDS workgroup would produce this whitepaper under the auspices of both AMIA and HIMSS, using support by AMIA through the AHRQ whitepaper grant, and additional support from HIMSS for the CDS workgroup activities. The workgroup also includes the chair of the eHealth Initiative's e-Prescribing Project (which had recently published a comprehensive whitepaper containing recommendations for improving value and adoption of eRx, including CDS issues) and the chair of the HL7 Clinical Decision Support technical committee. Some of the material in this whitepaper reflects work done by those two organizations as well.

HHS units designated as primary recipients for the whitepaper include ONCHIT and AHRQ, as well as NCVHS and CMS (due to their related responsibilities under the

Medicare Modernization Act). Because the newly-created Commission on Certification of Health Information Technology (CCHIT) will play an important role in driving CDS features in electronic health records, and because the models developed here have direct applicability to CCHIT's work, two members from that commission were also included as primary recipients of the recommendations.

The workgroup developed drafts of the recommendation tables, and discussed these in detail during a half-day meeting with invited experts at the Medinfo Conference on September 9, 2004 in San Francisco. The initial panelists were selected based on stakeholder representation and expertise concerning the issues at hand. Based on feedback obtained during this meeting, the tables were revised and circulated to a broader group for feedback. After a number of further rounds of input, the tables were revised into the final versions presented in this whitepaper.

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