

identify the record, specify the information they are contesting, state the corrective action sought and the reasons for the correction along with supporting justification showing why the record is not accurate, timely, relevant, or complete. Rules regarding amendment of Privacy Act records appear in 45 CFR part 5a. If additional information or assistance is required, contact the HHS Privacy Act Officer, Room 2221, Mary E. Switzer Building, Department of Health and Human Services, 330 "C" Street, SW., Washington, DC 20201. Write the words "Privacy Act Request" on the envelope and on the letter.

#### RECORDS SOURCE CATEGORIES:

Employee, contractor, or applicant; sponsoring agency; former sponsoring agency; other federal agencies; contract employer; former employer.

#### SYSTEMS EXEMPTED FROM CERTAIN PROVISIONS OF THE ACT:

None.

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Notice of Availability: Test Tools and Test Procedures Approved for the Office of the National Coordinator for Health Information Technology (ONC) Temporary Certification Program

**AGENCY:** Office of the National Coordinator for Health Information Technology, Office of the Secretary, Department of Health and Human Services.

**ACTION:** Notice.

**Authority:** 42 U.S.C. 300jj-11.

**SUMMARY:** This notice announces the availability of test tools and test procedures approved by the National Coordinator for Health Information Technology (the National Coordinator) for the testing of Complete EHRs and/or EHR Modules by ONC-Authorized Testing and Certification Bodies (ONC-ATCBs) under the ONC temporary certification program. The approved test tools and test procedures are identified on the ONC Web site at: <http://healthit.hhs.gov/certification>.

**FOR FURTHER INFORMATION CONTACT:** Carol Bean, Director, Certification Division, Office of the National Coordinator for Health Information Technology, 202-690-7151.

#### SUPPLEMENTARY INFORMATION:

On June 24, 2010, the Department of Health and Human Services issued a

final rule establishing a temporary certification program for the purposes of testing and certifying health information technology ("Establishment of the Temporary Certification Program for Health Information Technology," 75 FR 36158) (Temporary Certification Program final rule).<sup>1</sup> The Temporary Certification Program final rule added a new "Subpart D—Temporary Certification Program for HIT" to part 170 of title 45 of the Code of Federal Regulations (CFR). Section 170.423(e) of Subpart D requires ONC-ATCBs to "[u]se test tools and test procedures approved by the National Coordinator for the purposes of assessing Complete EHRs and/or EHR Modules compliance with the certification criteria adopted by the Secretary." The preamble of the Temporary Certification Program final rule stated that when the National Coordinator had approved test tools and/or test procedures ONC would publish a notice of availability in the **Federal Register** and identify the approved test tools and test procedures on the ONC Web site. As discussed in the Temporary Certification Program final rule, we anticipated that test tools and test procedures would not be finalized by the National Institute of Standards and Technology (NIST), and therefore unable to be considered for approval by the National Coordinator, until after the Secretary made publicly available a final rule for the initial set of standards, implementation specifications, and certification criteria for electronic health record technology.<sup>2</sup> This final rule, "Health Information Technology: Initial Set of Standards, Implementation Specifications, and Certification Criteria for Electronic Health Record Technology" (HIT Standards and Certification Criteria final rule) was made available for public inspection on July 13, 2010, and was published in the **Federal Register** on July 28, 2010.

The National Coordinator has approved, for use by ONC-ATCBs in accordance with 45 CFR 170.423(e), test tools and test procedures developed by NIST for testing Complete EHRs and/or

<sup>1</sup> The Department issued a proposed rule entitled "Proposed Establishment of Certification Programs for Health Information Technology" (75 FR 11328, March 10, 2010) that proposed the establishment of a temporary certification program and a permanent certification program and stated the Department's intentions to issue separate final rules for each program.

<sup>2</sup> The "Health Information Technology: Initial Set of Standards, Implementation Specifications, and Certification Criteria for Electronic Health Record Technology" interim final rule was made available for public inspection on December 30, 2009, and published in the **Federal Register** on January 13, 2010 (75 FR 2014).

EHR Modules to the applicable certification criterion or criteria adopted by the Secretary in the HIT Standards and Certification Criteria final rule. These approved test tools and test procedures are identified on the ONC Web site at: <http://healthit.hhs.gov/certification>.

Dated: August 2, 2010.

**David Blumenthal,**  
National Coordinator for Health Information Technology.

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Health Resources and Services Administration

#### Agency Information Collection Activities: Submission for OMB Review; Comment Request

Periodically, the Health Resources and Services Administration (HRSA) publishes abstracts of information collection requests under review by the Office of Management and Budget (OMB), in compliance with the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35). To request a copy of the clearance requests submitted to OMB for review, e-mail [paperwork@hrsa.gov](mailto:paperwork@hrsa.gov) or call the HRSA Reports Clearance Office on (301) 443-1129.

The following request has been submitted to the Office of Management and Budget for review under the Paperwork Reduction Act of 1995:

**Proposed Project: Title:** "Health Care and Other Facilities" Project Status Update Form (OMB No. 0915-0309)—[Extension].

The Health Resources and Services Administration's Health Care and Other Facilities (HCOF) program provides congressionally-directed funds to health-related facilities for construction-related activities and/or capital equipment purchases. Awarded facilities are required to provide a periodic (quarterly for construction-related projects, annually for equipment only projects) update of the status of the funded project until it is completed. The monitoring period averages about 3 years, although some projects take up to 5 years to complete. The information collected from these updates is vital to program management staff to determine whether projects are progressing according to the established timeframes, meeting deadlines established in the Notice of Grant Award (NGA), and drawing down funds appropriately. The

data collected from the updates is also shared with the Division of Grants Management Operations (DGMO) for their assistance in the overall evaluation of each project's progress.

An electronic form is currently being used for progress reporting for the

HCOF program. This form provides awardees access to directly input the required status update information in a timely, consistent, and uniform manner. The electronic form minimizes burden to respondents and informs respondents when there are missing data elements

prior to submission. We acknowledge a change in the burden estimate due to close out of old projects, and the addition of new projects for FY 2010.

The annual estimate of burden is as follows:

Instrument	Number of respondents	Responses per respondent	Total responses	Hours per response	Total Burden hours
Construction-Related .....	481	4	1,924	.5	962
Equipment Only .....	1,238	1	1,238	.5	619
<b>Total .....</b>	<b>1,719</b>	<b>.....</b>	<b>3,162</b>	<b>.....</b>	<b>1,581</b>

Written comments and recommendations concerning the proposed information collection should be sent within 30 days of this notice to the desk officer for HRSA, either by e-mail to *OIRA\_submission@omb.eop.gov* or by fax to 202-395-6974. Please direct all correspondence to the "attention of the desk officer for HRSA."

Dated: August 3, 2010.

**Sahira Rafiullah,**

*Director, Division of Policy and Information Coordination.*

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Chapter 35). To request a copy of these requests, call the CDC Reports Clearance Officer at (404) 639-5960 or send an e-mail to *omb@cdc.gov*. Send written comments to CDC Desk Officer, Office of Management and Budget, Washington, DC or by fax to (202) 395-5806. Written comments should be received within 30 days of this notice.

**Proposed Project**

Evaluation of Safe Dates Project—(OMB No. 0920-0783 exp. 6/30/2011)—Revision—National Center for Injury Prevention and Control (NCIPC), Centers for Disease Control and Prevention (CDC).

*Background and Brief Description*

Safe Dates, a dating violence prevention curriculum for 8th and 9th grade students, has been shown to be effective at preventing victimization and perpetration of teen dating violence in one rural North Carolina school district, but appropriateness of the program with urban, high-risk adolescents is unknown. The data collection will require participation from teachers at eight schools who delivered the Safe Dates program and students at one school who received the program. Qualitative data will be collected

through student focus groups and teacher interviews. Students will complete a participant profile form to capture basic demographic information. The specific aim of this study is to assess whether the Safe Dates adolescent dating violence prevention program needs modification/adaptation for urban, high-risk adolescents.

Approximately 40 students at one school will participate in focus groups. Two focus groups will consist of 8-10 boys, and two focus groups will include 8-10 girls. Informed written consent from parents for each student's participation and informed written assent from tenth graders for their own participation will be obtained. Twenty teachers will participate in interviews. Students and teachers will be asked about their experiences with the Safe Dates program and ideas they may have about adapting the program for urban schools.

Data collection will occur in July 2010. It is anticipated that study results will be used to determine whether the Safe Dates program should be modified for an urban, high-risk population. There is no cost to respondents other than their time. The total estimated annual burden hours are 849.

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Disease Control and Prevention**

[30 Day-10-0783]

**Agency Forms Undergoing Paperwork Reduction Act Review**

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C.

**ESTIMATED ANNUALIZED BURDEN HOURS**

Type of respondent	Instrument name	Number of respondents	Number of responses per respondent	Average burden per respondent (in hours)	Total response burden (hours)
Student .....	Effectiveness follow-up survey .....	1,318	1	35/60	769
	Focus group guide and participant profile form.	40	1	1.5	60
Teacher .....	Interview guide .....	20	1	1	20