

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

[Document Identifier: HHS-OS-17657-30D]

Agency Information Collection Activities; Submission to OMB for Review and Approval; Public Comment Request

AGENCY: Office of the Secretary, HHS.

ACTION: Notice.

SUMMARY: In compliance with section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, will submit an Information Collection Request (ICR), described below, to the Office of Management and Budget (OMB) for review and approval. The ICR is for revision of the approved information collection assigned OMB control number 0955-0002, scheduled to expire on October 31, 2012. Comments submitted during the first public review of this ICR will be provided to OMB. OMB will accept further comments from the public on this ICR during the review and approval period.

DATES: Comments on the ICR must be received on or before November 19, 2012.

ADDRESSES: Submit your comments to OIRA_submission@omb.eop.gov or via facsimile to (202) 395-5806. See

SUPPLEMENTARY INFORMATION for other information about submitting comments.

FOR FURTHER INFORMATION CONTACT: Information Collection Clearance staff, Information.Collection.Clearance@hhs.gov, (202) 690-6162.

SUPPLEMENTARY INFORMATION: When submitting comments or requesting information, please include the OMB control number 0955-0002 and document identifier HHS-OS-17657-30D for reference.

Information Collection Request Title: Facts for Consumers about Health IT Service Providers.

OMB No.: 0955-0002.

Abstract: ONC is proposing to revise current OMB approved Facts for Consumers about Health IT Service Providers. The current OMB approval is applicable through October 31, 2012. It includes iterative rounds of in-depth consumer testing to assess and analyze consumer understanding and input about a model privacy notice for personal health records (PHRs). ONC intends to revise the project to use the same focus group and cognitive usability interview testing process for the development of a model notice of privacy practices (NPP).

Need and Proposed Use of the Information: 45 CFR 164.520 requires covered entities to make available a NPP for protected health information to their patients or health plan members. The notice must, among other things, outline the purposes for which the covered entity is permitted to use and disclose health information, the rights of individuals with respect to their health information, the entities' duties to protect that information, and the process for filing a complaint concerning possible violations of the HIPAA Privacy Rule, such as an improper use or disclosure of information. 45 CFR 164.520 requires that the notice be written in plain language, but studies have shown that these notices are often difficult for

patients to understand due to their length and complexity.

The Federal Health IT Strategic Plan identifies the Fair Information Practice Principles (FIPPS) an important guidepost in the development of privacy policies and programs. Openness and Transparency is a key principle of fair information practices. The NPP is an important component of fulfilling this principle. If patients cannot adequately understand the notice because of its length or complexity, then the use and disclosure of their health information is not open and transparent.

In addition, each participant will have been recruited through a 15-minute screening interview. The participants will be recruited according to U.S. census statistics for race/ethnicity, age, marital status, gender, and income.

Likely Respondents: Respondents to this information collection are members of the General Public.

Burden Statement: Burden in this context means the time expended by persons to generate, maintain, retain, disclose or provide the information requested. This includes the time needed to review instructions, to develop, acquire, install and utilize technology and systems for the purpose of collecting, validating and verifying information, processing and maintaining information, and disclosing and providing information, to train personnel and to be able to respond to a collection of information, to search data sources, to complete and review the collection of information, and to transmit or otherwise disclose the information. The total annual burden hours estimated for this ICR are summarized in the table below.

TOTAL ESTIMATED ANNUALIZED BURDEN—HOURS

Forms (if necessary)	Type of respondent	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Cognitive Testing Screening	General Public	84	1	15/60	21
Cognitive Testing	General Public	42	1	90/60	63
Total	126	84

Keith A. Tucker,

Information Collection Clearance Officer.

[FR Doc. 2012-25692 Filed 10-18-12; 8:45 am]

BILLING CODE 4150-45-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

[Document Identifier: HHS-ONC-17577-30D]

Agency Information Collection Activities; Submission to OMB for Review and Approval; Public Comment Request

AGENCY: Office of the National Coordinator for Health IT, HHS.

ACTION: Notice.

SUMMARY: In compliance with section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the Office of the National Coordinator for Health IT (ONC), Department of Health and Human Services, has submitted an Information Collection Request (ICR), described below, to the Office of Management and Budget (OMB) for review and approval. The ICR is for a new collection. Comments submitted during the first public review of this ICR will be provided to OMB. OMB will accept further comments from the public on this ICR during the review and approval period.

DATES: Comments on the ICR must be received on or before November 19, 2012.

ADDRESSES: Submit your comments to OIRA_submission@omb.eop.gov or via facsimile to (202) 395-5806.

FOR FURTHER INFORMATION CONTACT: Information Collection Clearance staff, Information.CollectionClearance@hhs.gov or (202) 690-6162.

SUPPLEMENTARY INFORMATION: When submitting comments or requesting information, please include the Information Collection Request Title and document identifier HHS-ONC-17577-30D for reference.

Information Collection Request Title: National Survey on Health Information Exchange in Clinical Laboratories.

Abstract: ONC seeks approval to collect key data from a relatively small sample of clinical laboratories nationwide for the Evaluation of the State Health Information Exchange Cooperative Agreement Program. The *National Survey on Health Information Exchange in Clinical Laboratories* will assess and evaluate the electronic transfer of health information from clinical laboratories to ordering physicians. It will focus on two key measures: (1) Percentage of laboratory facilities that are able to send structured lab results electronically to ordering physicians and (2) percentage of lab results that are currently being sent electronically in coded format to ordering physicians.

Need and Proposed Use of the Information: A key goal of the State Health Information Exchange Cooperative Agreement Program is to

promote the electronic exchange of structured test results from clinical laboratories to healthcare providers. ONC will use these survey findings to develop a comprehensive understanding of the baseline level of laboratory information exchange in order to inform program activities to promote laboratory information exchange and provide more targeted assistance to states and territories in developing their laboratory information exchange strategies.

Likely Respondents: There will be two similar versions of the questionnaire—one for hospital-based labs and one for independent labs.

Burden Statement: Burden in this context means the time expended by persons to generate, maintain, retain, disclose or provide the information requested. This includes the time needed to review instructions, to develop, acquire, install and utilize technology and systems for the purpose of collecting, validating and verifying information, processing and maintaining information, and disclosing and providing information, to train personnel and to be able to respond to a collection of information, to search data sources, to complete and review the collection of information, and to transmit or otherwise disclose the information. The total annual burden hours estimated for this ICR are summarized in the table below.

TOTAL ESTIMATED ANNUALIZED BURDEN—HOURS

Form name	Type of respondent	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hours
Hospital-Based Laboratory Survey on Health Information Exchange.	Hospital-Based Laboratories	2,729	1	20/60	910
	Independent Laboratories	1,963	1	17.70/60	579
Total	4,692	1	19.04/60	1,489

Keith A. Tucker,
Information Collection Clearance Officer.
[FR Doc. 2012-25737 Filed 10-18-12; 8:45 am]
BILLING CODE 4150-45-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Announcement of Requirements and Registration for Cancer Care Video Challenge

AGENCY: Office of the National Coordinator for Health Information Technology, HHS.

ACTION: Notice.

SUMMARY: The *Cancer Care Video Challenge* is an opportunity for members of the public to create short, <2 minute videos sharing a personal story of how they use technology to help meet a goal related to an experience with cancer. Cash prizes are available to winning videos.

DATES: Effective on October 17, 2012.

FOR FURTHER INFORMATION CONTACT: Erin Poetter, Policy Analyst, Office of Consumer eHealth, ONC erin.poetter@hhs.gov; 202.205.3310.

SUPPLEMENTARY INFORMATION:

Subject of Challenge Competition: The Office of the National Coordinator for Health Information Technology (ONC), seeks to motivate and inspire patients and their families to get access to their health information and to leverage health IT and other consumer eHealth tools to be empowered to better manage their health and the health of loved ones. Patients and their families today have access to an unprecedented number of tools and resources to enhance their ability manage care. The *Cancer Care Video Challenge* is an