

the forms, their requests for approval are evaluated. There is no cost to

respondents other than their time to participate.

Estimated Annualized Burden Hours:

Section name	Data type	Number of respondents	Number of responses per respondent	Average burden response (in hours)	Total burden hours
84.11	Applications	43	8	86	29,584
84.33	Labeling	43	8	2	688
84.35	Modifications	43	8	66	22,704
84.41	Reporting	43	8	23	7,912
84.43	Record Keeping	43	8	46	15,824
84.257	Labeling	43	8	3	1,032
84.1103	Labeling	43	8	3	1,032
Total					78,776

Maryam I. Daneshvar,

Acting Reports Clearance Officer, Centers for Disease Control and Prevention.

[FR Doc. E7-16591 Filed 8-21-07; 8:45 am]

BILLING CODE 4163-18-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

National Institute for Occupational Safety and Health (NIOSH) Advisory Board on Radiation and Worker Health (ABRWH or Advisory Board)

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92-463), the Centers for Disease Control and Prevention announces the following committee meeting:

Name: Advisory Board on Radiation and Worker Health, National Institute for Occupational Safety and Health.

Audio Conference Call Time and Date: 11 a.m.–4 p.m., EST, Tuesday, September 4, 2007.

Place: Audio Conference Call via FTS Conferencing. The USA toll free dial in number is 1-866-643-6504 with a pass code of 9448550.

Status: Open to the public, but without a public comment period.

Background: The Advisory Board was established under the Energy Employees Occupational Illness Compensation Program Act of 2000 to advise the President on a variety of policy and technical functions required to implement and effectively manage the new compensation program. Key functions of the Advisory Board include providing advice on the development of probability of causation guidelines which have been promulgated by the Department of Health and Human Services (HHS) as a final rule, advice on methods of dose reconstruction which have also been promulgated by HHS as a final rule, advice on the scientific validity and quality of dose estimation and reconstruction efforts being performed for purposes of the compensation

program, and advice on petitions to add classes of workers to the Special Exposure Cohort (SEC).

In December 2000, the President delegated responsibility for funding, staffing, and operating the Advisory Board to HHS, which subsequently delegated this authority to the CDC. NIOSH implements this responsibility for CDC. The charter was issued on August 3, 2001, renewed at appropriate intervals, most recently, August 3, 2007, and will expire on August 3, 2009.

Purpose: This Advisory Board is charged with (a) providing advice to the Secretary, HHS, on the development of guidelines under Executive Order 13179; (b) providing advice to the Secretary, HHS, on the scientific validity and quality of dose reconstruction efforts performed for this program; and (c) upon request by the Secretary, HHS, advising the Secretary on whether there is a class of employees at any Department of Energy facility who were exposed to radiation but for whom it is not feasible to estimate their radiation dose, and on whether there is reasonable likelihood that such radiation doses may have endangered the health of members of this class.

Matters To Be Discussed: The agenda for the conference call includes: Report of Board Member Votes Recorded Since the Last Board Meeting; Update on Rocky Flats Follow-Up Actions; Update on SC&A Review of TBD 6000 and General Steel Industries Appendix; Report on SC&A Contract Talks for FY08; Discussions of Initial Steps for a Board Contractor for FY09 and Beyond; Report on Privacy Act "Clearance" Procedures; Update on Letters to DOE and DOL on Chapman Valve; Work Group Updates; Status of and Plans for Future Board Activities; and Board Working Time.

The agenda is subject to change as priorities dictate.

Recommended changes to the agenda from the Office of General Counsel resulted in the **Federal Register** notice being published less than fifteen days before the date of the meeting.

Because there is not a public comment period, written comments may be submitted. Any written comments received will be included in the official record of the meeting

and should be submitted to the contact person below well in advance of the meeting.

Contact Person for More Information: Dr. Lewis V. Wade, Executive Secretary, NIOSH, CDC, 4676 Columbia Parkway, Cincinnati, OH 45226, Telephone 513.533.6825, Fax 513.533.6826.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both CDC and the Agency for Toxic Substances and Disease Registry.

Dated: August 16, 2007.

Michael Tropauer,

Acting Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. E7-16557 Filed 8-21-07; 8:45 am]

BILLING CODE 4163-18-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers For Medicare & Medicaid Services

Privacy Act of 1974; Report of a Modified System of Records

AGENCY: Department of Health and Human Services (HHS), Center for Medicare & Medicaid Services (CMS).

ACTION: Notice of a Modified System of Records (SOR).

SUMMARY: In accordance with the requirements of the Privacy Act of 1974, we propose to modify an existing system titled, "National Emphysema Treatment Trial (NETT), System No. 09-70-0531," established at 65 **Federal Register** 47995 (August 4, 2000). We propose to modify existing routine use number 1 that permits disclosure to agency contractors and consultants to include disclosure to CMS grantees who perform a task for the agency. CMS grantees, charged with completing

projects or activities that require CMS data to carry out that activity, are classified separate from CMS contractors and/or consultants. The modified routine use will remain as routine use number 1. We will delete routine use number 4 authorizing disclosure to support constituent requests made to a congressional representative. If an authorization for the disclosure has been obtained from the data subject, then no routine use is needed. The Privacy Act allows for disclosures with the "prior written consent" of the data subject.

We will broaden the scope of routine uses number 5 and 6, authorizing disclosures to combat fraud and abuse in the Medicare and Medicaid programs to include combating "waste" which refers to specific beneficiary or recipient practices that result in unnecessary cost to all federally-funded health benefit programs. Additionally, we will broaden the scope of this system by including the section titled "Additional Circumstances Affecting Routine Use Disclosures," that addresses "Protected Health Information (PHI)" and "small cell size." The requirement for compliance with HHS regulation "Standards for Privacy of Individually Identifiable Health Information" apply when ever the system collects or maintain PHI. This system may contain PHI. In addition, our policy to prohibit release if there is a possibility that an individual can be identified through "small cell size" will apply to the data disclosed from this system.

We are modifying the language in the remaining routine uses to provide a proper explanation as to the need for the routine use and to provide clarity to CMS's intention to disclose individual-specific information contained in this system. The routine uses will then be prioritized and reordered according to their usage. We will also take the opportunity to update any sections of the system that were affected by the recent reorganization or because of the impact of the Medicare Prescription Drug, Improvement, and Modernization Act of 2003 (MMA) (Pub. L. 108-173) provisions and to update language in the administrative sections to correspond with language used in other CMS SORs.

The primary purpose of the system of records is to collect and maintain data that will allow CMS to provide secure data on participants in the randomized phase of the study, pay claims, and to monitor and evaluate the clinical trial. Information retrieved from this system of records will also be disclosed to: (1) Support regulatory, reimbursement and policy functions performed within the

agency or by a contractor, consultant, or CMS grantee; (2) assist another Federal or state agency to enable such agency to administer a Federal health benefits program, or to enable such agency to fulfill a requirement of a Federal statute or regulation that implements a health benefits program funded in whole or in part with Federal funds; (3) support research, evaluation, or epidemiological projects related to the prevention of disease or disability, or the restoration or maintenance of health, and for payment related projects; (4) support litigation involving the agency; and (5) combat fraud, waste, and abuse in certain health benefits programs. We have provided background information about the new system in the **SUPPLEMENTARY INFORMATION** section below. Although the Privacy Act requires only that CMS provide an opportunity for interested persons to comment on the proposed routine uses, CMS invites comments on all portions of this notice. See **EFFECTIVE DATE** section for comment period.

EFFECTIVE DATE: CMS filed a modified system report with the Chair of the House Committee on Government Reform and Oversight, the Chair of the Senate Committee on Homeland Security and Governmental Affairs, and the Administrator, Office of Information and Regulatory Affairs, Office of Management and Budget (OMB) on August 10, 2007. To ensure that all parties have adequate time in which to comment, the modified SOR, including routine uses, will become effective 40 days from the publication of the notice, or from the date it was submitted to OMB and the Congress, whichever is later, unless CMS receives comments that require alterations to this notice.

ADDRESSES: The public should address comments to: CMS Privacy Officer, Division of Privacy Compliance, Enterprise Architecture and Strategy Group, Office of Information Services, CMS, Room N2-04-27, 7500 Security Boulevard, Baltimore, Maryland 21244-1850. Comments received will be available for review at this location, by appointment, during regular business hours, Monday through Friday from 9 a.m.-3 p.m., Eastern Time zone.

FOR FURTHER INFORMATION CONTACT: Joanna Baldwin, Health Insurance Specialist, Division of Medical and Surgical Services, Coverage and Analysis Group, Office of Clinical Standards and Quality, CMS, Room C1-10-25, 7500 Security Boulevard, Baltimore, Maryland 21244-1850. Her telephone number is (410) 786-7205. She can be reached by telephone at 410-

786-7205 or e-mail Joanna.Baldwin@cms.hhs.gov.

SUPPLEMENTARY INFORMATION:

I. Description of the Proposed System of Records

A. Statutory and Regulatory Basis for SOR

The statutory authority for this system is given under the provisions of Section 1862(a) (1) (A) of the Social Security Act, and 42 U.S.C. 1395, which states that Medicare must provide coverage for items and services that are "reasonable and necessary for the diagnosis or treatment of illness or injury or to improve the functioning of a malformed body member."

B. Collection and Maintenance of Data in the System

This system will collect and maintain individually identifiable and other data collected on Medicare beneficiaries and providers participating in the study. Data will be collected from Medicare administrative and claims records, patient medical charts, physician records, and via survey instruments administered to beneficiaries and providers. The collected information will include, but is not limited to Medicare claims and eligibility data, name, address, telephone number, health insurance claims number, race/ethnicity, gender, date of birth, provider name, unique provider identification number, medical record number, as well as clinical, demographic, health/well-being, family and/or caregiver contact information, and background information relating to Medicare issues.

II. Agency Policies, Procedures, and Restrictions on Routine Uses

A. The Privacy Act permits us to disclose information without an individual's consent if the information is to be used for a purpose that is compatible with the purpose(s) for which the information was collected. Any such disclosure of data is known as a "routine use." The Government will only release NETT information that can be associated with an individual as provided for under "Section III. Proposed Routine Use Disclosures of Data in the System." Both identifiable and non-identifiable data may be disclosed under a routine use. We will only collect the minimum personal data necessary to achieve the purpose of NETT.

CMS has the following policies and procedures concerning disclosures of information that will be maintained in the system. Disclosure of information from the system will be approved only

to the extent necessary to accomplish the purpose of the disclosure and only after CMS:

1. Determines that the use or disclosure is consistent with the reason that the data is being collected; e.g., to collect and maintain data that will allow CMS to provide secure data on participants in the randomized phase of the study, pay claims, and to monitor and evaluate the clinical trial.

2. Determines that:

- a. The purpose for which the disclosure is to be made can only be accomplished if the record is provided in individually identifiable form;

- b. The purpose for which the disclosure is to be made is of sufficient importance to warrant the effect and/or risk on the privacy of the individual that additional exposure of the record might bring; and

- c. There is a strong probability that the proposed use of the data would in fact accomplish the stated purpose(s).

3. Requires the information recipient to:

- a. Establish administrative, technical, and physical safeguards to prevent unauthorized use of disclosure of the record;

- b. Remove or destroy, at the earliest time, all patient-identifiable information; and

- c. Agree to not use or disclose the information for any purpose other than the stated purpose under which the information was disclosed.

- d. Determines that the data are valid and reliable.

III. Proposed Routine Use Disclosures of Data in the System

A. The Privacy Act allows us to disclose information without an individual's consent if the information is to be used for a purpose that is compatible with the purpose(s) for which the information was collected. Any such compatible use of data is known as a "routine use." The proposed routine uses in this system meet the compatibility requirement of the Privacy Act. We are proposing to establish the following routine use disclosures of information maintained in the system:

1. To agency contractors, consultants or CMS grantees, who have been engaged by the agency to assist in the performance of a service related to this collection and who need to have access to the records in order to perform the activity.

We contemplate disclosing information under this routine use only in situations in which CMS may enter into a contractual or similar agreement with a third party to assist in

accomplishing CMS function relating to purposes for this system.

CMS occasionally contracts out certain of its functions when doing so would contribute to effective and efficient operations. CMS must be able to give a contractor, consultant or CMS grantees whatever information is necessary for the contractor or consultant to fulfill its duties. In these situations, safeguards are provided in the contract prohibiting the contractor, consultant or CMS grantees from using or disclosing the information for any purpose other than that described in the contract and requires the contractor, consultant or CMS grantees to return or destroy all information at the completion of the contract.

2. To another Federal or state agency to:

- a. contribute to the accuracy of CMS's proper payment of Medicare benefits;

- b. enable such agency to administer a Federal health benefits program, or, as necessary, to enable such agency to fulfill a requirement of a Federal statute or regulation that implements a health benefits program funded in whole or in part with Federal funds; and/or

- c. assist Federal/state Medicaid programs within the state.

Other Federal or state agencies, in their administration of a Federal health program, may require NETT information in order to support evaluations and monitoring of Medicare claims information of beneficiaries, including proper reimbursement for services provided.

3. To an individual or organization for a research project or in support of an evaluation project related to the prevention of disease or disability, the restoration or maintenance of health, or payment related projects.

The NETT data will provide for research or support of evaluation projects and a broader, longitudinal, national perspective of the status of Medicare beneficiaries. CMS anticipates that many researchers will have legitimate requests to use these data in projects that could ultimately improve the care provided to Medicare beneficiaries and the policies that govern their care.

4. To the Department of Justice (DOJ), court or adjudicatory body when:

- a. the agency or any component thereof, or

- b. any employee of the agency in his or her official capacity, or

- c. any employee of the agency in his or her individual capacity where the DOJ has agreed to represent the employee, or

- d. the United States Government, is a party to litigation or has an interest in

such litigation, and, by careful review, CMS determines that the records are both relevant and necessary to the litigation and that the use of such records by the DOJ, court or adjudicatory body is compatible with the purpose for which the agency collected the records.

Whenever CMS is involved in litigation, and occasionally when another party is involved in litigation and CMS policies or operations could be affected by the outcome of the litigation, CMS would be able to disclose information to the DOJ, court or adjudicatory body involved.

5. To a CMS contractor (including, but not necessarily limited to, fiscal intermediaries and carriers) that assists in the administration of a CMS-administered health benefits program, or to a grantee of a CMS-administered grant program, when disclosure is deemed reasonably necessary by CMS to prevent, deter, discover, detect, investigate, examine, prosecute, sue with respect to, defend against, correct, remedy, or otherwise combat fraud, waste, and abuse in such program.

We contemplate disclosing information under this routine use only in situations in which CMS may enter into a contractual, grantee, cooperative agreement or consultant relationship with a third party to assist in accomplishing CMS functions relating to the purpose of combating fraud, waste, and abuse.

CMS occasionally contracts out certain of its functions or makes grants or cooperative agreements when doing so would contribute to effective and efficient operations. CMS must be able to give a contractor, grantee, consultant or other legal agent whatever information is necessary for the agent to fulfill its duties. In these situations, safeguards are provided in the contract prohibiting the agent from using or disclosing the information for any purpose other than that described in the contract and requiring the agent to return or destroy all information.

6. To another Federal agency or to an instrumentality of any governmental jurisdiction within or under the control of the United States (including any State or local governmental agency), that administers, or that has the authority to investigate potential fraud, waste, or abuse in, a health benefits program funded in whole or in part by Federal funds, when disclosure is deemed reasonably necessary by CMS to prevent, deter, discover, detect, investigate, examine, prosecute, sue with respect to, defend against, correct, remedy, or otherwise combat fraud, waste, and abuse in such programs.

Other agencies may require NETT information for the purpose of combating fraud, waste, and abuse in such Federally-funded programs.

B. Additional Provisions Affecting Routine Use Disclosures

To the extent this system contains Protected Health Information (PHI) as defined by HHS regulation "Standards for Privacy of Individually Identifiable Health Information" (45 CFR Parts 160 and 164, Subparts A and E) 65 FR 82462 (12-28-00). Disclosures of such PHI that are otherwise authorized by these routine uses may only be made if, and as, permitted or required by the "Standards for Privacy of Individually Identifiable Health Information." (See 45 CFR 164-512 (a) (1).)

In addition, our policy will be to prohibit release even of data not directly identifiable, except pursuant to one of the routine uses or if required by law, if we determine there is a possibility that an individual can be identified through implicit deduction based on small cell sizes (instances where the patient population is so small that individuals could, because of the small size, use this information to deduce the identity of the beneficiary).

IV. Safeguards

CMS has safeguards in place for authorized users and monitors of such users to ensure against unauthorized use. Personnel having access to the system have been trained in the Privacy Act and information security requirements. Employees who maintain records in this system are instructed not to release data until the intended recipient agrees to implement appropriate management, operational and technical safeguards sufficient to protect the confidentiality, integrity and availability of the information and information systems and to prevent unauthorized access.

This system will conform to all applicable Federal laws and regulations and Federal, HHS, and CMS policies and standards as they relate to information security and data privacy. These laws and regulations may apply but are not limited to: The Privacy Act of 1974; the Federal Information Security Management Act of 2002; the Computer Fraud and Abuse Act of 1986; the Health Insurance Portability and Accountability Act of 1996; the E-Government Act of 2002, the Clinger-Cohen Act of 1996; the Medicare Modernization Act of 2003, and the corresponding implementing regulations. OMB Circular A-130, Management of Federal Resources, Appendix III, Security of Federal

Automated Information Resources also applies. Federal, HHS, and CMS policies and standards include but are not limited to: All pertinent National Institute of Standards and Technology publications; the HHS Information Systems Program Handbook and the CMS Information Security Handbook.

V. Effects of the Modified System of Records on Individual Rights

CMS proposes to establish this system in accordance with the principles and requirements of the Privacy Act and will collect, use, and disseminate information only as prescribed therein. Data in this system will be subject to the authorized releases in accordance with the routine uses identified in this system of records.

CMS will take precautionary measures to minimize the risks of unauthorized access to the records and the potential harm to individual privacy or other personal or property rights of patients whose data are maintained in this system. CMS will collect only that information necessary to perform the system's functions. In addition, CMS will make disclosure from the proposed system only with consent of the subject individual, or his/her legal representative, or in accordance with an applicable exception provision of the Privacy Act. CMS, therefore, does not anticipate an unfavorable effect on individual privacy as a result of information relating to individuals.

Dated: August 7, 2007.

Charlene Frizzera,

Chief Operating Officer, Centers for Medicare & Medicaid Services.

SYSTEM NO.:

09-70-0531.

SYSTEM NAME:

"National Emphysema Treatment Trial (NETT)," HHS/CMS/OCSQ.

SECURITY CLASSIFICATION:

Level Three Privacy Act Sensitive Data.

SYSTEM LOCATION:

CMS Data Center, 7500 Security Boulevard, North Building, First Floor, Baltimore, Maryland 21244-1850 and at various other contractor locations.

CATEGORIES OF INDIVIDUALS COVERED BY THE SYSTEM:

This system will collect and maintain individually identifiable and other data collected on Medicare beneficiaries and providers participating in the study.

CATEGORIES OF RECORDS IN THE SYSTEM:

Data will be collected from Medicare administrative and claims records,

patient medical charts, physician records, and via survey instruments administered to beneficiaries and providers. The collected information will include, but is not limited to Medicare claims and eligibility data, name, address, telephone number, health insurance claims number, race/ethnicity, gender, date of birth, provider name, unique provider identification number, medical record number, as well as clinical, demographic, health/well-being, family and/or caregiver contact information, and background information relating to Medicare issues.

AUTHORITY FOR MAINTENANCE OF THE SYSTEM:

The statutory authority for this system is given under the provisions of Section 1862(a)(1)(A) of the Social Security Act, and 42 U.S.C. 1395, which states that Medicare must provide coverage for items and services that are "reasonable and necessary for the diagnosis or treatment of illness or injury or to improve the functioning of a malformed body member."

PURPOSE(S) OF THE SYSTEM:

The primary purpose of the system of records is to collect and maintain data that will allow CMS to provide secure data on participants in the randomized phase of the study, pay claims, and to monitor and evaluate the clinical trial. Information retrieved from this system of records will also be disclosed to: (1) Support regulatory, reimbursement and policy functions performed within the agency or by a contractor, consultant, or CMS grantee; (2) assist another Federal or state agency to enable such agency to administer a Federal health benefits program, or to enable such agency to fulfill a requirement of a Federal statute or regulation that implements a health benefits program funded in whole or in part with Federal funds; (3) support research, evaluation, or epidemiological projects related to the prevention of disease or disability, or the restoration or maintenance of health, and for payment related projects; (4) support litigation involving the agency; and (5) combat fraud, waste, and abuse in certain health benefits programs.

ROUTINE USES OF RECORDS MAINTAINED IN THE SYSTEM, INCLUDING CATEGORIES OR USERS AND THE PURPOSES OF SUCH USES:

A. The Privacy Act allows us to disclose information without an individual's consent if the information is to be used for a purpose that is compatible with the purpose(s) for which the information was collected. Any such compatible use of data is known as a "routine use." The proposed routine uses in this system meet the

compatibility requirement of the Privacy Act. We are proposing to establish the following routine use disclosures of information maintained in the system:

1. To agency contractors, consultants or CMS grantees, who have been engaged by the agency to assist in the performance of a service related to this collection and who need to have access to the records in order to perform the activity.

2. To another Federal or state agency to:

a. contribute to the accuracy of CMS's proper payment of Medicare benefits;

b. enable such agency to administer a Federal health benefits program, or, as necessary, to enable such agency to fulfill a requirement of a Federal statute or regulation that implements a health benefits program funded in whole or in part with Federal funds; and/or

c. assist Federal/state Medicaid programs within the state.

3. To an individual or organization for a research project or in support of an evaluation project related to the prevention of disease or disability, the restoration or maintenance of health, or payment related projects.

4. To the Department of Justice (DOJ), court or adjudicatory body when:

a. the agency or any component thereof, or

b. any employee of the agency in his or her official capacity, or

c. any employee of the agency in his or her individual capacity where the DOJ has agreed to represent the employee, or

d. the United States Government, is a party to litigation or has an interest in such litigation, and, by careful review, CMS determines that the records are both relevant and necessary to the litigation and that the use of such records by the DOJ, court or adjudicatory body is compatible with the purpose for which the agency collected the records.

5. To a CMS contractor (including, but not necessarily limited to, fiscal intermediaries and carriers) that assists in the administration of a CMS-administered health benefits program, or to a grantee of a CMS-administered grant program, when disclosure is deemed reasonably necessary by CMS to prevent, deter, discover, detect, investigate, examine, prosecute, sue with respect to, defend against, correct, remedy, or otherwise combat fraud, waste, and abuse in such program.

6. To another Federal agency or to an instrumentality of any governmental jurisdiction within or under the control of the United States (including any State or local governmental agency), that administers, or that has the authority to

investigate potential fraud or abuse in, a health benefits program funded in whole or in part by Federal funds, when disclosure is deemed reasonably necessary by CMS to prevent, deter, discover, detect, investigate, examine, prosecute, sue with respect to, defend against, correct, remedy, or otherwise combat fraud, waste, and abuse in such programs.

B. Additional Provisions Affecting Routine Use Disclosures

To the extent this system contains Protected Health Information (PHI) as defined by HHS regulation "Standards for Privacy of Individually Identifiable Health Information" (45 CFR Parts 160 and 164, Subparts A and E) 65 FR 82462 (12-28-00). Disclosures of such PHI that are otherwise authorized by these routine uses may only be made if, and as, permitted or required by the "Standards for Privacy of Individually Identifiable Health Information." (See 45 CFR 164-512 (a) (1).)

In addition, our policy will be to prohibit release even of data not directly identifiable, except pursuant to one of the routine uses or if required by law, if we determine there is a possibility that an individual can be identified through implicit deduction based on small cell sizes (instances where the patient population is so small that individuals could, because of the small size, use this information to deduce the identity of the beneficiary).

POLICIES AND PRACTICES FOR STORING, RETRIEVING, ACCESSING, RETAINING, AND DISPOSING OF RECORDS IN THE SYSTEM:

STORAGE:

All records are stored on electronic media.

RETRIEVABILITY:

The collected data are retrieved by an individual identifier; e.g., beneficiary name or HICN, and unique provider identification number.

SAFEGUARDS:

CMS has safeguards in place for authorized users and monitors such users to ensure against unauthorized use. Personnel having access to the system have been trained in the Privacy Act and information security requirements. Employees who maintain records in this system are instructed not to release data until the intended recipient agrees to implement appropriate management, operational and technical safeguards sufficient to protect the confidentiality, integrity and availability of the information and information systems and to prevent unauthorized access.

This system will conform to all applicable Federal laws and regulations

and Federal, HHS, and CMS policies and standards as they relate to information security and data privacy. These laws and regulations may apply but are not limited to: The Privacy Act of 1974; the Federal Information Security Management Act of 2002; the Computer Fraud and Abuse Act of 1986; the Health Insurance Portability and Accountability Act of 1996; the E-Government Act of 2002, the Clinger-Cohen Act of 1996; the Medicare Modernization Act of 2003, and the corresponding implementing regulations. OMB Circular A-130, Management of Federal Resources, Appendix III, Security of Federal Automated Information Resources also applies. Federal, HHS, and CMS policies and standards include but are not limited to: All pertinent National Institute of Standards and Technology publications; the HHS Information Systems Program Handbook and the CMS Information Security Handbook.

RETENTION AND DISPOSAL:

CMS will retain information for a total period not to exceed 25 years. All claims-related records are encompassed by the document preservation order and will be retained until notification is received from DOJ.

SYSTEM MANAGER AND ADDRESS:

Director, Division of Medical and Surgical Services, Coverage and Analysis Group, Office of Clinical Standards and Quality, CMS, Room C4-10-10, 7500 Security Boulevard, Baltimore, Maryland 21244-1850.

NOTIFICATION PROCEDURE:

For purpose of access, the subject individual should write to the system manager who will require the system name, employee identification number, tax identification number, national provider number, and for verification purposes, the subject individual's name (woman's maiden name, if applicable), HICN, and/or SSN (furnishing the SSN is voluntary, but it may make searching for a record easier and prevent delay).

RECORD ACCESS PROCEDURE:

For purpose of access, use the same procedures outlined in Notification Procedures above. Requestors should also reasonably specify the record contents being sought. (These procedures are in accordance with Department regulation 45 CFR 5b.5 (a) (2).)

CONTESTING RECORD PROCEDURES:

The subject individual should contact the system manager named above, and reasonably identify the record and specify the information to be contested.

State the corrective action sought and the reasons for the correction with supporting justification. (These procedures are in accordance with Department regulation 45 CFR 5b.7.)

RECORDS SOURCE CATEGORIES:

The data collected and maintained in this system are retrieved from Medicare enrollment records, Medicare beneficiaries or proxies, and medical providers (such as physicians, medical facilities, home health care providers) for a sample of enrollees.

SYSTEMS EXEMPTED FROM CERTAIN PROVISIONS OF THE ACT:

None.

[FR Doc. 07-4076 Filed 8-21-07; 8:45 am]

BILLING CODE 4120-03-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects:

Title: National Survey of Child and Adolescent Well-Being—Second Cohort (NSCAW II).

OMB No.: 0970-0202.

Description: The Department of Health and Human Services (HHS)

intends to collect data on a new sample of children and families for the National Survey of Child and Adolescent Well-Being (NSCAW). The NSCAW was authorized under Section 427 of the Personal Responsibility and Work Opportunities Reconciliation Act of 1996. The original survey began in November 1999 with a national sample of 5,501 children, ages 0–14, who had been the subject of investigation by Child Protective Services during the baseline data collection period, which extended from November 1999 through April 2000. Direct assessments and interviews were conducted with the children themselves, their primary caregivers, their caseworkers, and, for school-aged children, their teachers; agency directors also were interviewed at baseline. Follow-up data collections were conducted 12 months, 18 months, and 36 months post-baseline, and a fifth data collection is currently under way.

The NSCAW is the only source of nationally representative, firsthand information about the functioning and well-being, service needs, and service utilization of children and families who come to the attention of the child welfare system. Information is collected about children's cognitive, social, emotional, behavioral, and adaptive functioning, as well as family and community factors that are likely to influence their functioning. Family

service needs and service utilization also are addressed in the data collection.

The current data collection plan calls for selecting a new cohort of 5,700 children and families and repeating the data collection procedures used in the original study. Selection of a new cohort will allow the comparison of characteristics of children who are entering the child welfare system today with those who entered prior to the implementation of the Adoption and Safe Families Act and prior to the advent of the Child and Family Services Review process. The data collection will follow the same format as that used in previous rounds of data collection, and will employ, with only modest revisions, the same instruments that have been used in previous rounds.

Currently, HHS intends to collect baseline data and one follow-up 18 months later, with future follow-up rounds contingent on funding availability. Data from NSCAW are made available to the research community through licensing arrangements from the National Data Archive on Child Abuse and Neglect at Cornell University.

Respondents: 5,700 Children and their associated permanent or foster caregivers, caseworkers, and teachers; in addition, an administrator will be interviewed in each location from which children are sampled.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Child Interview	5,700	1	1.2	6,840
Permanent Caregiver Interview	3,800	1	2.0	7,600
Foster Caregiver Interview	1,990	1	1.5	2,985
Caseworker Interview	5,700	1	1.0	5,700
Teacher Questionnaire	3,000	1	.75	2,250
Agency Questionnaire	97	1	1.0	97

Estimated Total Annual Burden Hours: 25,472.

In compliance with the requirements of Section 506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Administration, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. E-mail address:

infocollection@acf.hhs.gov. All requests should be identified by the title of the information collection.

The Department specifically requests comments on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or

other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Dated: August 15, 2007.

Brendan C. Kelly,

Reports Clearance Officer.

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