

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Medicare & Medicaid Services

#### Privacy Act of 1974; Report of a New System of Records

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

**ACTION:** Notice of a New System of Records (SOR).

**SUMMARY:** In accordance with the requirements of the Privacy Act of 1974, we are proposing to establish a new system titled, "Senior Risk Reduction Demonstration and Evaluation (SRRDE), System No. 09-70-0592." The program is authorized under provisions of the Social Security Act (42 U.S.C. 1395b-1(a)), which gives the Secretary the broad authority to, "develop and engage in experiments and demonstration projects." The goal of the SRRDE is to determine whether risk reduction programs that have been developed and tested in the private sector can also be tailored to and work well with Medicare beneficiaries to improve their health and reduce avoidable health care utilization. The specific aims of the demonstration and evaluation are to: (1) Determine whether a senior risk reduction service provided by Medicare will be accepted by beneficiaries, achieve high participation rates, and be viewed positively by beneficiaries; (2) reduce health risk factors, improve health behaviors, improve functioning, and prevent disability; and (3) save money for Medicare.

The purpose of this system is to collect and maintain demographic and health related data on the target population of non-institutionalized Medicare beneficiaries between the ages of 67 and 74 who are potential participants in the SRRDE program. We will also collect certain identifying information on Medicare providers who provide services to such beneficiaries. Information retrieved from this system may be disclosed to: (1) Support regulatory, reimbursement, and policy functions performed within the agency or by a contractor, grantee, consultant or other legal agent; (2) assist another Federal or state agency with information to contribute to the accuracy of CMS's proper payment of Medicare benefits, enable such agency to administer a Federal health benefits program, or to enable such agency to fulfill a requirement of Federal statute or regulation that implements a health benefits program funded in whole or in

part with Federal funds; (3) support 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) support litigation involving the agency; and (5) combat fraud, waste, and abuse in certain Federally-funded 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 DATES** section for comment period.

**DATES: Effective Date:** CMS filed a new SOR report with the Chair of the House Committee on Government Reform and Oversight, the Chair of the Senate Committee on Homeland Security & Governmental Affairs, and the Administrator, Office of Information and Regulatory Affairs, Office of Management and Budget (OMB) on October 6, 2006. To ensure that all parties have adequate time in which to comment, the new system will become effective 30 days from the publication of the notice, or 40 days from the date it was submitted to OMB and the Congress, whichever is later. We may defer implementation of this system or one or more of the routine use statements listed below if we receive comments that persuade us to defer implementation.

**ADDRESSES:** The public should address comments to the CMS Privacy Officer, Division of Privacy Compliance, Enterprise Architecture and Strategy Group, Office of Information Services, Mail-stop 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.

**FOR FURTHER INFORMATION CONTACT:** Pauline Lapin, Division of Health Promotion & Disease Prevention Demonstrations, Medicare Demonstrations Program Group, Office of Research Development & Information, Mail Stop S3-06-24, Centers for Medicare & Medicaid Services, 7500 Security Boulevard, Baltimore, MD 21244-1849. She can be reached by telephone at 410-786-6883, or via e-mail at [Pauline.Lapin@cms.hhs.gov](mailto:Pauline.Lapin@cms.hhs.gov).

**SUPPLEMENTARY INFORMATION:** Behavioral lifestyle choices with respect to diet, physical activity, alcohol, and tobacco

use are associated with the leading causes of morbidity and mortality in the United States. Recent research suggests that well-structured risk reduction programs can achieve significant improvements in a population's risk profile. Successful programs are founded on solid behavior change theory, use tailored interventions, are personalized and sufficiently intensive, and are delivered with adequate social supports. The SRRDE program will be tailored to the needs, concerns, and learning styles of seniors. The goal is to develop personalized materials and instruments, followed by interventions tailored to the risks presented by the participants.

CMS will offer risk reduction services to non-institutionalized Medicare beneficiaries between the ages of 67 and 74. The demonstration requires random selection of beneficiaries from across the United States as well as from communities that have exemplary Information and Referral/Assistance programs for seniors. Medicare will assign approximately 15,000-17,000 randomly selected beneficiaries to each site for recruitment. Medicare's inclusion criteria for beneficiaries eligible for the demonstration and evaluation are as follows: they must be a Medicare fee-for-service beneficiary enrolled in both Parts A and B, they may be dual eligible for both Medicare and Medicaid, and Medicare must be their primary payer. Medicare's exclusion criteria for beneficiaries to participate in the demonstration and evaluation are as follows: they cannot be currently enrolled in a Medicare Health Plan; they cannot be enrolled in a hospice or End State Renal Disease (ESRD) program; cannot currently be participating in another CMS demonstration; cannot have residence in an institution for 100 days or the past 12 months; cannot have the inability to participate in self-care activities due to severe dementia or other serious mental illness; and cannot have had initial enrollment into Medicare before the age of 65.

#### I. Description of the Proposed System of Records

##### A. Statutory and Regulatory Basis for SOR

The statutory authority for this system is given under section 402(a)(1)(B) and (a)(2) of the Social Security Amendments of 1967, Public Law No. 90-248, as amended, 42 United States Code § 1395b-1(a)(1)(B) and (a)(2).

### *B. Collection and Maintenance of Data in the System*

This system will collect and maintain individually identifiable and other data collected on non-institutionalized Medicare beneficiaries between the ages of 67 and 74 who are potential participants in the SRRDE program. 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. Data will be collected from Medicare administrative and claims records, SRRDE site administrative data systems, patient medical charts, and physician records.

### **II. Agency Policies, Procedures, and Restrictions on the Routine Use**

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 SRRDE 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 SRRDE.

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 demographic and health related data on the target population of non-institutionalized Medicare beneficiaries between the ages of 67 and 74 who are potential participants in the SRRDE program.

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.

4. 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 support agency contractors, consultants or 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 grantee 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 grantee from using or disclosing the information for any purpose other than that described in the contract and requires the contractor, consultant or grantee to return or destroy all information at the completion of the contract.

2. To assist 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 SRRDE information in order to support evaluations and monitoring of Medicare claims information of beneficiaries, including proper reimbursement for services provided.

3. To support 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 SRRDE 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 researchers may 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 support 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 assist 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, or 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 assist 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, or abuse in such programs.

Other agencies may require SRRDE 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 such users to ensure against excessive or 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 Proposed 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: October 4, 2006.

**Charlene Frizzera,**

*Acting Chief Operating Officer, Centers for Medicare & Medicaid Services.*

#### **SYSTEM NO. 09-70-0592**

##### **SYSTEM NAME:**

"Senior Risk Reduction Demonstration and Evaluation (SRRDE)," HHS/CMS/ORDI.

##### **SECURITY CLASSIFICATION:**

Level Three Privacy Act Sensitive Data.

##### **SYSTEM LOCATION:**

Centers for Medicare & Medicaid Services (CMS) Data Center, 7500 Security Boulevard, North Building, First Floor, Baltimore, Maryland 21244-1850 and at various co-locations of CMS agents.

##### **CATEGORIES OF INDIVIDUALS COVERED BY THE SYSTEM:**

This system will collect and maintain individually identifiable and other data collected on non-institutionalized Medicare beneficiaries between the ages of 67 and 74 who are potential participants in the SRRDE program.

##### **CATEGORIES OF RECORDS IN THE SYSTEM:**

The collected information will include, but is not limited to: Medicare claims and eligibility data, name, address, telephone number, health insurance claims number (HICN), 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. Data will be collected from Medicare administrative and claims records, SRRDE site administrative data systems,

patient medical charts, and physician records.

**AUTHORITY FOR MAINTENANCE OF THE SYSTEM:**

The statutory authority for this system is given under section 402 (a)(1)(B) and (a)(2) of the Social Security Amendments of 1967, Public Law 90–248, as amended, 42 United States Code 1395b–1(a)(1)(B) and (a)(2).

**PURPOSE(S) OF THE SYSTEM:**

The purpose of this system is to collect and maintain demographic and health related data on the target population of non-institutionalized Medicare beneficiaries between the ages of 67 and 74 who are potential participants in the SRRDE program. We will also collect certain identifying information on Medicare providers who provide services to such beneficiaries. Information retrieved from this system may be disclosed to: (1) Support regulatory, reimbursement, and policy functions performed within the agency or by a contractor, grantee, consultant or other legal agent; (2) assist another Federal or state agency with information to contribute to the accuracy of CMS's proper payment of Medicare benefits, enable such agency to administer a Federal health benefits program, or to enable such agency to fulfill a requirement of Federal statute or regulation that implements a health benefits program funded in whole or in part with Federal funds; (3) support 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) support litigation involving the agency; and (5) combat fraud, waste, and abuse in certain Federally-funded 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 support agency contractors, consultants or 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 assist 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 support 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 support 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 assist 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, or abuse in such program.

6. To assist 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, or 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.

**SAFEGUARDS:**

CMS has safeguards in place for authorized users and monitors such users to ensure against excessive or 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 10 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 Office of Research Development & Information, Mail Stop S3-06-24, Centers for Medicare & Medicaid Services, 7500 Security Boulevard, Baltimore, MD 21244-1849.

#### 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:

Data will be collected from Medicare administrative and claims records, SRRDE site administrative data systems, patient medical charts, and physician records.

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

None.

[FR Doc. E6-17055 Filed 10-13-06; 8:45 am]

BILLING CODE 4120-03-P

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Medicare & Medicaid Services

#### Privacy Act of 1974; Report of a Modified or Altered System of Records

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

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

**SUMMARY:** In accordance with the Privacy Act of 1974, we are proposing to modify or alter an existing SOR, "Medicare Current Beneficiary Survey (MCBS)," System No. 09-70-6002, last published at 66 *Federal Register* 15496 (March 19, 2001). We propose to assign a new CMS identification number to this system to simplify the obsolete and confusing numbering system originally designed to identify the Bureau, Office, or Center of the Health Care Financing Administration that maintained the system of records. The new assigned identifying number for this system should read: System No. 09-70-0519.

We propose to modify existing routine use number 2 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 be renumbered 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 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 this modified system is to collect and maintain a research database for CMS and other researchers that is capable of producing data sets suitable for both longitudinal and cross-sectional analysis to be used to: (1) Produce projections for current programs and proposed program changes, (2) produce national level estimates of health care expenditures by the aged and disabled, and (3) provide a research database that can be used to provide guidance to program management and policies. The 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 a CMS grantee; (2) assist another Federal or State agency with information to contribute to the accuracy of CMS's proper payment of Medicare benefits, enable such agency to administer a Federal health benefits program, or to enable such agency to fulfill a requirement of Federal statute or regulation that implements a health benefits program funded in whole or in part with Federal funds; (3) assist 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; and (4) support litigation involving the agency. We have provided background information about the modified 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 modified or altered routine uses, CMS invites comments on all portions of this notice. See **EFFECTIVE DATE** section for comment period.

**EFFECTIVE DATE:** CMS filed a modified or altered SOR report with the Chair of the