Federal Communications Commission. William F. Caton,

Acting Secretary.

[FR Doc. 02-5255 Filed 3-4-02; 8:45 am]

BILLING CODE 6712-01-P

FEDERAL EMERGENCY **MANAGEMENT AGENCY**

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

ACTION: Notice and request for comments.

SUMMARY: The Federal Emergency Management Agency has submitted the following proposed information collection to the Office of Management and Budget for review and clearance in accordance with the requirements of the Paperwork Reduction Act of 1995 (44 U.S.C. 3507).

Title: Application Form for Single Lot or Structural Amendments to National Flood Insurance Program Maps.

Type of Information Collection: Revision of a Currently Approved Collection.

OMB Number: 3067-0257. Abstract: FEMA Form 81-92 is designed to assist requesters in gathering information that FEMA needs to determine whether a certain single-lot property or structure is likely to be flooded during a flood event that has a one-percent annual change of being equaled or exceeded in any given year (base flood). FEMA Form 81-92A is a Spanish version of FEMA Form 81–92 and, as such, only one of the two formas would be required for any one application.

Affected Public: Individual or Households, Business or Other For-Profit, and State, Local or Tribal Government.

Number of Respondents: 9,000. Estimated Time per Respondent: FEMA Form 81–92, 2.4 hours and FEMA Form 81-92A, 2.4 hours.

Estimated Total Annual Burden Hours: 21,600 hours.

Frequency of Response: On Occasion. Comments: Interested persons are invited to submit written comments on the proposed information collection to the Desk Officer for the Federal Emergency Management Agency, Office of Information and Regulatory Affairs, Office of Management and Budget, Washington, DC 20503 within 30 days of the date of this notice.

FOR FURTHER INFORMATION CONTACT:

Requests for additional information or copies of the information collection should be made to Muriel B. Anderson, Chief, Records Management Section, Program Services and Systems Branch, Facilities Management and Services Division, Administration and Resource Planning Directorate, Federal Emergency Management Agency, 500 C Street, SW, Room 316, Washington, DC 20472, telephone number (202) 646-2625 or facsimile number (202) 646-3347, or e-mail muriel.anderson@fema.gov.

Dated: February 27, 2002.

Reginald Trujillo,

Branch Chief, Program Services and Systems Branch, Facilities and Services Management Division, Administration and Resource Planning Directorate.

[FR Doc. 02-5159 Filed 3-4-02; 8:45 am] BILLING CODE 6718-01-P

FEDERAL RESERVE SYSTEM

Formations of, Acquisitions by, and Mergers of Bank Holding Companies

The companies listed in this notice have applied to the Board for approval, pursuant to the Bank Holding Company Act of 1956 (12 U.S.C. 1841 et seq.) (BHC Act), Regulation Y (12 CFR Part 225), and all other applicable statutes and regulations to become a bank holding company and/or to acquire the assets or the ownership of, control of, or the power to vote shares of a bank or bank holding company and all of the banks and nonbanking companies owned by the bank holding company, including the companies listed below.

The applications listed below, as well as other related filings required by the Board, are available for immediate inspection at the Federal Reserve Bank indicated. The application also will be available for inspection at the offices of the Board of Governors. Interested persons may express their views in writing on the standards enumerated in the BHC Act (12 U.S.C. 1842(c)). If the proposal also involves the acquisition of a nonbanking company, the review also includes whether the acquisition of the nonbanking company complies with the standards in section 4 of the BHC Act (12 U.S.C. 1843). Unless otherwise noted, nonbanking activities will be conducted throughout the United States. Additional information on all bank holding companies may be obtained from the National Information Center website at www.ffiec.gov/nic/.

Unless otherwise noted, comments regarding each of these applications must be received at the Reserve Bank indicated or the offices of the Board of Governors not later than April 1, 2002.

A. Federal Reserve Bank of Chicago (Phillip Jackson, Applications Officer)

230 South LaSalle Street, Chicago, Illinois 60690-1414:

1. The Baraboo Bancorporation, Inc., Baraboo, Wisconsin; to acquire 100 percent of the voting shares of The Bancorp, Inc., Cedarburg, Wisconsin, and thereby indirectly acquire Northwoods State Bank, Elcho, Wisconsin.

Board of Governors of the Federal Reserve System, February 28, 2002.

Robert deV. Frierson,

Deputy Secretary of the Board. [FR Doc. 02-5197 Filed 3-4-02; 8:45 am] BILLING CODE 6210-01-S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

Privacy Act of 1974; Report of New System

AGENCY: Department of Health and Human Services (HHS), Centers for Medicare & Medicaid Services (CMS) (formerly the Health Care Financing Administration).

ACTION: Notice of 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 of records. The proposed system is titled "Claims Payment System For Medicare's "Healthy Aging" Demonstration Project (CPS-HA), HHS/ CMS/CBC, System No. 09-70-0539." CMS proposes to establish a new system of records containing enrollment and claims payment information plus research-related survey data, in support of a short-term demonstration project testing new potential benefits in the Medicare program.

The primary purpose of the system of records is to manage and maintain information needed to pay Medicare claims under the research demonstration program known as the Healthy Aging project (HA) including its component known as the Medicare Stop Smoking Program (MSSP). The system of records will enable CMS to: enroll and communicate with eligible Medicare beneficiaries who volunteer to participate in HA initiatives, communicate with clinicians and other providers and suppliers who submit claims payable under HA demonstrations, review submitted claims and pay those conforming to applicable payment criteria and federal law, and develop, maintain, and analyze research information showing the

potential impact of HA interventions on the quality and cost of health care services in Medicare. Information retrieved from this system of records will also be disclosed to support regulatory, reimbursement, and policy functions performed within the agency or by a contractor or consultant; assist another Federal or State agency with information 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; support constituent requests made to a Congressional representative; support litigation involving the agency; facilitate research on the quality and effectiveness of care provided; and, combat fraud and abuse in certain health benefits programs. We have provided background information about the proposed system in the "Supplementary Information" section below. Although the Privacy Act requires only that the "routine use" portion of the system be published for comment, CMS invites comments on all portions of this notice. See **EFFECTIVE DATES** section for comment period.

EFFECTIVE DATES: CMS filed a new system report with the Chair of the House Committee on Government Reform and Oversight, the Chair of the Senate Committee on Governmental Affairs, and the Administrator, Office of Information and Regulatory Affairs, Office of Management and Budget (OMB) on February 25, 2002. In any event, we will not disclose any information under a routine use until 40 days after publication. We may defer implementation of this system of records 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: Director, Division of Data Liaison and Distribution (DDLD), 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:

James Coan, Division of Health Promotion and Disease Prevention, Center for Beneficiary Choices, CMS, Mailstop S3–02–01, 7500 Security Boulevard, Baltimore, Maryland 21244– 1850. The telephone number is (410) 786–9168.

SUPPLEMENTARY INFORMATION:

I. Description of the New System of Records

Statutory and Regulatory Basis For System of Records

The authority to conduct the demonstration project for which the system of records is needed is section 402(a) of Public Law 90–248, as amended by section 222(b)(2) of Public Law 92–603, 42 U.S.C. 1395b–1(a)

II. Collection and Maintenance of Data in the System

A. Scope of the Data Collected

The CPS—HA includes the Medicare Health Insurance Claim (HIC) Number, sex, race, age, zip code, state and county. It also includes claims information related to HA claims, answers to enrollment questionnaires and other information needed to confirm a beneficiary's eligibility for enrollment and ongoing participation in the demonstration, and other survey and research information needed to pay claims, administer the HA program, and develop research reports on the study's findings.

B. Agency Policies, Procedures, and Restrictions on the Routine Use

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 CPS-HA information that can be associated with an individual patient as provided for under "Section III. Entities Who May Receive Disclosures Under Routine Use." Both identifiable and nonidentifiable data may be disclosed under a routine use. Identifiable data includes individual records with CPS-HA information and identifiers. Nonidentifiable data includes individual records with CPS-HA information and masked identifiers or CPS-HA information with identifiers stripped out of the file.

We will only disclose the minimum personal data necessary to achieve the purpose of the CPS—HA system. CMS has the following policies and procedures concerning disclosures of information that will be maintained in the system. In general, disclosure of information from the SOR will be approved only for the minimum information necessary to accomplish the purpose of the disclosure after CMS:

1. Determines that the use or disclosure is consistent with the reason that the data is being collected; *e.g.*,

developing and refining payment systems and monitoring the quality of care provided to patients.

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;
- 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. Entities Who May Receive Disclosures Under Routine Use

These routine uses specify circumstances, in addition to those provided by statute in the Privacy Act of 1974, under which CMS may release information from the CPS-HA without the consent of the individual to whom such information pertains. Each proposed disclosure of information under these routine uses will be evaluated to ensure that the disclosure is legally permissible, including but not limited to ensuring that the purpose of the disclosure is compatible with the purpose for which the information was collected. We are proposing to establish the following routine use disclosures of information maintained in the system:

1. To agency contractors, or consultants who have been contracted by the agency to assist in the performance of a service related to this system of records 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 agency business functions relating to purposes for this system of records.

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 whatever information is necessary for the contractor to fulfill its duties. In these situations, safeguards are provided in the contract prohibiting the contractor from using or disclosing the information for any purpose other than that described in the contract and requires the contractor to return or destroy all information at the completion of the contract.

2. To a Peer Review Organization (PRO) in order to assist the PRO to perform Title XI and Title XVIII functions relating to assessing and improving HA quality of care. PROs will work to implement quality improvement programs, provide consultation to CMS, its contractors, and to State agencies.

The PROs may use these data to support quality improvement activities and other PRO responsibilities as detailed in Title XI, sections 1151–1164.

- 3. To another Federal or State agency:
- a. To contribute to the accuracy of CMS's proper payment of Medicare benefits,
- b. To 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, or
- c. To improve the state survey process for investigation of complaints related to health and safety or quality of care and to implement a more outcome oriented survey and certification program.

Other Federal or State agencies in their administration of a Federal health program may require CPS—HA information in order to support evaluations and monitoring of quality of care for special populations or special care areas, including proper payment for services provided. Releases of information would be allowed if the proposed use(s) for the information proved compatible with the purpose for which CMS collects the information.

4. To an individual or organization for research on the utilization of inpatient rehabilitation services as well as evaluation or epidemiological projects related to the prevention of disease or disability, the restoration or maintenance of health, or for understanding and improving payment projects.

The CPS—HA data will provide an opportunity for comprehensive research, evaluation and

epidemiological projects regarding HA patients. CMS anticipates that many researchers will have legitimate requests to use these data in projects that could ultimately improve the care provided to HA patients and the policy that governs the care.

5. To a Member of Congress or to a congressional staff member in response to an inquiry of the Congressional Office made at the written request of the constituent about whom the record is maintained.

Beneficiaries sometimes request the help of a Member of Congress in resolving some issue relating to a matter before CMS. The Member of Congress then writes CMS, and CMS must be able to give sufficient information to be responsive to the inquiry.

- 6. 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.

Whenever CMS is involved in litigation, or occasionally when another party is involved in litigation and CMS's 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. A determination would be made in each instance that, under the circumstances involved, the purposes served by the use of the information in the particular litigation is compatible with a purpose for which CMS collects the information.

7. 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 or abuse in such program.

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

accomplishing CMS functions relating to the purpose of combating fraud and abuse.

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

8. 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 or abuse in such programs.

Other agencies may require CPS—HA information for the purpose of combating fraud and abuse in such Federally funded programs. Releases of information would be allowed if the proposed use(s) for the information proved compatible with the purposes of collecting the information.

9. To insurance companies, third party administrators (TPA), employers, self-insurers, managed care organizations, other supplemental insurers, non-coordinating insurers, multiple employer trusts, group health plans (i.e., health maintenance organizations (HMO) or a competitive medical plan (CMP)) with a Medicare contract, or a Medicare-approved health care prepayment plan (HCPP), directly or through a contractor, and other groups providing protection for their enrollees. Information to be disclosed shall be limited to Medicare entitlement data. In order to receive the information, they must agree to:

a. Certify that the individual about whom the information is being provided is one of its insured or employees, or is insured and/or employed by another entity for whom they serve as a third party administrator; utilize the information solely for the purpose of processing the individual's insurance claims; and

b. Safeguard the confidentiality of the data and prevent unauthorized access.

Other insurers, CMP, HMO, and HCPP may require CPS—HA information in order to support evaluations and monitoring of Medicare claims information of beneficiaries, including proper payment for services provided.

B. Additional Provisions Affecting Routine Use Disclosures

In addition, our policy will be to prohibit release even of non-identifiable data, except pursuant to one of the routine uses, if 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 who are familiar with the enrollees could, because of the small size, use this information to deduce the identity of the beneficiary).

This System of Records contains
Protected Health Information as defined
by the Department of Health and Human
Services' regulation "Standards for
Privacy of Individually Identifiable
Health Information" (45 CFR parts 160
and 164, 65 FR 82462 as amended by 66
FR 12434). Disclosures of Protected
Health Information 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."

IV. Safeguards

The HHS CPS-HA system will conform to applicable law and policy governing the privacy and security of Federal automated information systems. These include but are not limited to: the Privacy Act of 1984, Computer Security Act of 1987, the Paperwork Reduction Act of 1995, the Clinger-Cohen Act of 1996, and OMB Circular A-130, Appendix III, "Security of Federal Automated Information Resources." CMS has prepared a comprehensive system security plan as required by OMB Circular A–130, Appendix III. This plan conforms fully to guidance issued by the National Institute for Standards and Technology (NIST) in NIST Special Publication 800–18, "Guide for Developing Security Plans for Information Technology Systems." Paragraphs A–C of this section highlight some of the specific methods that CMS is using to ensure the security of this system and the information within it.

A. Authorized Users

Personnel having access to the system have been trained in Privacy Act requirements. Employees who maintain records in the system are instructed not to release any data until the intended recipient agrees to implement appropriate administrative, technical, procedural, and physical safeguards sufficient to protect the confidentiality of the data and to prevent unauthorized access to the data. Records are used in a designated work area and system location is attended at all times during working hours.

To ensure security of the data, the proper level of class user is assigned for each individual user level. This prevents unauthorized users from accessing and modifying critical data. The system database configuration includes five classes of database users:

- Database Administrator class owns the database objects (e.g., tables, triggers, indexes, stored procedures, packages) and has database administration privileges to these objects.
- Quality Control Administrator class has read and write access to key fields in the database;
- Quality Index Report Generator class has read-only access to all fields and tables:
- Policy Research class has query access to tables, but are not allowed to access confidential patient identification information; and
- Submitter class has read and write access to database objects, but no database administration privileges.

B. Physical Safeguards

All server sites will implement the following minimum requirements to assist in reducing the exposure of computer equipment and thus achieve an optimum level of protection and security for the CMS system:

Access to all servers is to be controlled, with access limited to only those support personnel with a demonstrated need for access. Servers are to be kept in a locked room accessible only by specified management and system support personnel. Each server is to require a specific log-on process. All entrance doors are identified and marked. A log is kept of all personnel who were issued a security card, key and/or combination, which grants access to the room housing the server, and all visitors are escorted while in this room. All servers are housed in an area where appropriate environmental security controls are implemented, which include measures implemented to mitigate damage to Automated Information Systems (AIS) resources caused by fire, electricity, water and inadequate climate controls.

Protection applied to the workstations, servers and databases include:

• User Log-on—Authentication is to be performed by the Primary Domain

Controller/Backup Domain Controller of the log-on domain.

- Workstation Names—Workstation naming conventions may be defined and implemented at the agency level.
- Hours of Operation—May be restricted by Windows NT. When activated all applicable processes will automatically shut down at a specific time and not be permitted to resume until the predetermined time. The appropriate hours of operation are to be determined and implemented at the agency level.
- Inactivity Lockout—Access to the NT workstation is to be automatically locked after a specified period of inactivity.
- Warnings—Legal notices and security warnings are to be displayed on all servers and workstations.
- Remote Access Security—Windows NT Remote Access Service (RAS) security handles resource access control. Access to NT resources is to be controlled for remote users in the same manner as local users, by utilizing Windows NT file and sharing permissions. Dial-in access can be granted or restricted on a user-by-user basis through the Windows NT RAS administration tool.

C. Procedural Safeguards

All automated systems must comply with Federal laws, guidance, and policies for information systems security. These include, but are not limited to: the Privacy Act of 1974; the Computer Security Act of 1987; OMB Circular A–130, revised; Information Resource Management (IRM) Circular #10; HHS Automated Information Systems Security Program; the CMS Information Systems Security Policy, Standards, and Guidelines Handbook; and other CMS systems security policies. Each automated information system should ensure a level of security commensurate with the level of sensitivity of the data, risk, and magnitude of the harm that may result from the loss, misuse, disclosure, or modification of the information contained in the system.

V. Effects of the New System 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 monitor the collection and reporting of CPS–HA data. CPS–HA

information on patients is submitted to CMS through standard systems. Accuracy of the data is important since incorrect information could result in the wrong payment for services and a less effective process for assuring quality of services. CMS will utilize a variety of onsite and offsite edits and audits to increase the accuracy of CPS–HA data.

CMS will take precautionary measures (see item IV. above) 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 is maintained in the 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 maintaining this system of records.

Dated: February 20, 2002.

Thomas A. Scully,

Administrator, Centers for Medicare & Medicaid Services.

09-70-0539

SYSTEM NAME:

Claims Payment System For Medicare's "Healthy Aging" Demonstration Project (CPS–HA).

SECURITY CLASSIFICATION:

Level 3, Privacy Act Sensitive.

SYSTEM LOCATION:

CMS Data Center, 7500 Security Boulevard, North Building, First Floor, Baltimore, Maryland 21244–1850 and CMS contractors and agents at various locations.

CATEGORIES OF INDIVIDUALS COVERED BY THE SYSTEM:

This system will contain claims and demographic information on Medicare beneficiaries who have volunteered to participate in Medicare's Healthy Aging program including specific demonstration projects such as the MSSP, as well as claims-related information for submissions from providers and suppliers providing services that are covered under Medicare exclusively within the HA program and its demonstration projects. The system will also retain research information such as enrollment questionnaires and survey data from participants in the program.

CATEGORIES OF RECORDS IN THE SYSTEM:

This system of records will contain demographic and claims-related information on Medicare beneficiaries who have elected to participate, as well as eligibility and enrollment data collected through voluntary surveys, payment information for providers and vendors submitting claims, and other information designed to support the enrollment, claims payment, and research reporting functions of the CPS—HA program.

AUTHORITY FOR MAINTENANCE OF THE SYSTEM:

The authority to conduct the demonstration project for which the system of records is needed is section 402(a)(1)(G) and (a)(2) of Public Law 90–248, as amended by section 222(b)(2) of Public Law 92–603, 42 U.S.C. 1395b–1(a)(1)(G) and (a)(2).

PURPOSE(S):

The primary purpose of the system of records is to manage and maintain information needed to pay Medicare claims under the research demonstration program known as the Healthy Aging project (HA) including its component known as the Medicare Stop Smoking Program (MSSP). The system of records will enable CMS to: Enroll and communicate with eligible Medicare beneficiaries who volunteer to participate in CPS-HA initiatives, communicate with clinicians and other providers and suppliers who submit claims payable under CPS-HA demonstrations, review submitted claims and pay those conforming to applicable payment criteria and federal law, and develop, maintain, and analyze research information showing the potential impact of CPS-HA interventions on the quality and cost of health care services in Medicare. Information retrieved from this system of records will also be disclosed to support regulatory, reimbursement, and policy functions performed within the agency or by a contractor or consultant; assist another Federal or State agency with information 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; support constituent requests made to a Congressional representative; support litigation involving the agency; facilitate research on the quality and effectiveness of care provided; and, combat fraud 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:

These routine uses specify circumstances, in addition to those provided by statute in the Privacy Act of 1974, under which CMS may release information from the CPS-HA without the consent of the individual to whom such information pertains. Each proposed disclosure of information under these routine uses will be evaluated to ensure that the disclosure is legally permissible, including but not limited to ensuring that the purpose of the disclosure is compatible with the purpose for which the information was collected. In addition, our policy will be to prohibit release even of nonidentifiable data, except pursuant to one of the routine uses, if 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 who are familiar with the enrollees could, because of the small size, use this information to deduce the identity of the beneficiary). Be advised, this System of Records contains Protected Health Information as defined by the Department of Health and Human Services' regulation "Standards for Privacy of Individually Identifiable Health Information" (45 CFR parts 160 and 164, 65 FR 8462 as amended by 66 FR 12434). Disclosures of Protected Health Information 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."

- 1. To agency contractors or consultants who have been contracted by the agency to assist in the performance of a service related to this system of records and who need to have access to the records in order to perform the activity.
- 2. To a Peer Review Organization (PRO) in order to assist the PRO to perform Title XI and Title XVIII functions relating to assessing and improving quality of care. PROs will work to implement quality improvement programs, provide consultation to CMS, its contractors, and to State agencies.
 - 3. To another Federal or State agency:
- a. To contribute to the accuracy of CMS's proper payment of Medicare benefits,
- b. To 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, or

c. To improve the state survey process for investigation of complaints related to health and safety or quality of care and to implement a more outcome oriented survey and certification program.

4. To an individual or organization for research on the utilization of inpatient rehabilitation services as well as evaluation or epidemiological projects related to the prevention of disease or disability, or the restoration or maintenance of health epidemiological, or for understanding and improving payment projects.

5. To a member of Congress or to a congressional staff member in response to an inquiry of the Congressional Office made at the written request of the constituent about whom the record is

maintained.

6. 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 the use of such records by the DOJ, court or adjudicatory body is compatible with the purpose for which the agency collected the records.
- 7. 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 or abuse in such program.
- 8. 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 or abuse in such programs.

- 9. To insurance companies, third party administrators (TPA), employers, self-insurers, managed care organizations, other supplemental insurers, non-coordinating insurers, multiple employer trusts, group health plans (i.e., health maintenance organizations (HMO) or a competitive medical plan (CMP)) with a Medicare contract, or a Medicare-approved health care prepayment plan (HCPP), directly or through a contractor, and other groups providing protection for their enrollees. Information to be disclosed shall be limited to Medicare entitlement data. In order to receive the information, they must agree to:
- a. Certify that the individual about whom the information is being provided is one of its insured or employees, or is insured and/or employed by another entity for whom they serve as a third party administrator;
- b. Utilize the information solely for the purpose of processing the individual's insurance claims; and
- c. Safeguard the confidentiality of the data and prevent unauthorized access

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

STORAGE:

All records are stored on magnetic media. Input data arrives as paper claims in the case of provider or supplier claims, and eligibility and enrollment information such as the enrollment survey and follow-up monitoring surveys are recorded in hard copy before transcription to magnetic media.

RETRIEVABILITY:

The Medicare records are retrieved by health insurance claim (HIC) number of the beneficiary. Provider IDs and supplier registration numbers are used to facilitate inquiries into specific claims as needed.

SAFEGUARDS:

CMS has safeguards 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 systems security requirements. Employees who maintain records in the system are instructed not to release any data until the intended recipient agrees to implement appropriate administrative, technical, procedural, and physical safeguards sufficient to protect the confidentiality of the data and to prevent unauthorized access to the data.

In addition, CMS has physical safeguards in place to reduce the

exposure of computer equipment and thus achieve an optimum level of protection and security for the CMS system. For computerized records, safeguards have been established in accordance with HHS standards and National Institute of Standards and Technology guidelines; e.g., security codes will be used, limiting access to authorized personnel. System securities are established in accordance with HHS, Information Resource Management (IRM) Circular #10, Automated Information Systems Security Program; CMS Information Systems Security, Standards Guidelines Handbook and OMB Circular No. A-130 (revised) Appendix III.

RETENTION AND DISPOSAL:

CMS will retain identifiable CPS—HA data for a total period of 25 years. Data residing with the designated claims payment contractor shall be returned to CMS at the end of the third project year, with all data then being the responsibility of CMS for adequate storage and security.

SYSTEM MANAGER(S) AND ADDRESS:

Director, Center for Beneficiary Choices, CMS, Mailstop C5–19–16, 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, health insurance claim number, and for verification purposes, the subject individual's name (woman's maiden name, if applicable), address, age, and sex, and social security number (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.)

RECORD SOURCE CATEGORIES:

- 1. Enrollment data on Medicare beneficiaries volunteering to participate in CPS—HA projects will come from beneficiaries who report the information to CMS officials or contractors, pursuant to information collection activities approved at the Office of Management and Budget and through an Institutional Review Board as required by law. Follow-up surveys and questionnaires for participants will also come directly from beneficiaries' voluntary reporting.
- 2. Claims data will come through voluntary submissions of providers, suppliers, and others seeking reimbursement for covered services provided to a Medicare beneficiary, in accordance with the provisions of the demonstration and the conditions of participation in the Medicare program.
- 3. Research analysis and reporting will come from the enrollment data, surveys and questionnaires provided by beneficiaries, as well as the analysis and compilations of this information developed by CMS officials, contractors, research collaborators, and others supporting the CPS–HA project and fulfilling the conditions of confidentiality, privacy and security outlined in this Notice.
- 4. Eligibility information as well as financial or quality reporting related to program integrity or other matters may also interact with existing CMS registries such as those relating to Medicare claims, provider registries, beneficiary enrollment databases, national claims histories.
- 5. Provider information to document the eligibility of a provider, supplier, or other person or entity to submit Medicare claims under the CPS—HA program, receive continuing medical education within the scope of the CPS—HA program, or for other uses will come from existing Medicare records of eligible providers and suppliers (as may be modified according to the needs of the CPS—HA program).

SYSTEMS EXEMPTED FROM CERTAIN PROVISIONS OF THE ACT:

None.

[FR Doc. 02–5140 Filed 3–4–02; 8:45 am] BILLING CODE 4120–03–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Advisory Committee Meetings; Correction

In FR Doc. 01–28108 appearing on pages 56689–56690 in the issue for

Friday, November 9, 2001, the dates and location of some Health Professions and Nurse Education Special Emphasis Panel meetings have changed. The meeting scheduled on April 22–25, 2002, has changed to March 18–21, 2002; the meeting on April 29–May 2, 2002, has changed to April 2–5, 2002; the meeting on May 6–9, 2002, has changed to April 15–18, 2002; and the location of these meetings has changed to the Hilton Silver Spring. The correct information is as follows:

Name: Health Careers Opportunity Program Peer Review Group.

Date and Time: March 1â–21, 2002. Place: Hilton Silver Spring, 8727 Colesville Road, Silver Spring, MD 20910.

Open on: March 18, 2002, 8 a.m. to 10 a.m. Closed on: March 18, 2002, 10 a.m. to adjournment (approximately 6 p.m.), March 19–21, 2002, 8 a.m. to adjournment (approximately 6 p.m.).

Name: Basic Nurse Education and Practice Grants Program Peer Review Group I. Date and Time: April 2–5, 2002. Place: Hilton Silver Spring, 8727 Colesville Road, Silver Spring, MD 20910.

Open on: April 2, 2002, 8 a.m. to 10 a.m. Closed on: April 2, 2002, 10 a.m. to adjournment (approximately 6 p.m.), April 3–5, 2002, 8 a.m. to adjournment (approx. 6 p.m.).

Name: Basic Nurse Education and Practice Grants Program Peer Review Group II. Date and Time: April 15–18, 2002. Place: Hilton Silver Spring, 8727 Colesville Road, Silver Spring, MD 20910.

Open on: April 15, 2002, 8 a.m. to 10 a.m. Closed on: April 15, 2002, 10 a.m. to adjournment (approximately 6 p.m.), April 16–18, 2002, 8 a.m. to adjournment (approximately 6 p.m.).

Dated: February 26, 2002.

Jane M. Harrison,

Director, Division of Policy Review and Coordination.

[FR Doc. 02–5132 Filed 3–4–02; 8:45 am] BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

National Advisory Council on Nurse Education and Practice; Notice of Meeting

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), announcement is made of the following National Advisory Council on Nurse Education and Practice scheduled to meet during the month of April 2002.

Name: National Advisory Council on Nurse Education and Practice Date and Time: April 11, 2002, 8:30 a.m.-5 p.m., April 12, 2002, 8:30 a.m.-3 p.m.

Place: Hotel Washington, Pennsylvania
Avenue, NW., at 15th St., Washington, DC

The meeting is open to the public. Agenda: Department, Agency, Bureau, and Division administrative updates; introduction of new members; discussion of Council administrative procedures; and presentations of national and regional nursing workforce issues with special emphasis on nursing faculty shortage to be followed with workgroup sessions on nursing workforce and education for practice improvement to address strategies for intervention and recommendations impacting Title VIII legislation.

Anyone interested in obtaining a roster of members, minutes of the meeting, or other relevant information should write or contact Ms. Elaine G. Cohen, Executive Secretary, National Advisory Council on Nurse Education and Practice, Parklawn Building, Room 9–35, 5600 Fishers Lane, Rockville, Maryland 20857, telephone (301) 443–1405.

Dated: February 27, 2002.

Jane M. Harrison,

Director, Division of Policy Review and Coordination.

[FR Doc. 02–5161 Filed 3–4–02; 8:45 am] BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Public Health Service

National Toxicology Program (NTP); National Institute of Environmental Health Sciences (NIEHS)

The NTP Center for the Evaluation of Risks to Human Reproduction (CERHR) 1. Announces a Future Evaluation of Ethylene Glycol (CASRN: 107–21–1) and Propylene Glycol (CASRN: 57–55–6), 2. Requests Public Input on These Chemicals, and 3. Solicits the Nomination of Individuals Qualified to Serve on an Expert Panel.

Evaluation of Ethylene Glycol and Propylene Glycol

The CERHR plans to hold an expert panel evaluation of ethylene glycol (CASRN: 107–21–1) and propylene glycol (CASRN: 57–55–6). The exact date for this expert panel meeting is not yet set, but is tentatively planned for the fall of 2002. Additional details about the meeting, including the date and location, will be published in a future **Federal Register** notice.

The CERHR will convene an expert panel to evaluate the reproductive and developmental toxicity of ethylene glycol and propylene glycol. The expert panel will consist of approximately 12 scientists, selected for their scientific