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 18–21, 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 20004

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

expertise in various aspects of reproductive and developmental toxicology and other relevant areas of science. The expert panel meeting will be open to the public with time scheduled for oral public comment.

Ethylene glycol is a high production volume chemical used chiefly in antifreeze for heating and cooling systems. There is widespread exposure to ethylene glycol due to its use as an automotive antifreeze and as a de-icer for aircraft. The toxicology database on ethylene glycol includes recent mechanistic data and occupational exposure information. Propylene gylcol, similar in structure to ethylene glycol, is used as an antifreeze, de-icing solution, and in various paints and coatings. Unlike ethylene glycol, propylene glycol is approved for use in various food additives, drugs, and cosmetics.

Request for Public Input

The CERHR invites input from the public and other interested parties on ethylene glycol and propylene glycol including toxicology information from completed and ongoing studies, information on planned studies, as well as information about current production levels, human exposure, use patterns, and environmental occurrence. Information and comments should be forwarded to the CERHR at P.O. Box 12233, MD EC–32, Research Triangle Park, NC 27709 (mail), (919) 541–3455 (phone), (919) 316–4511 (fax), or *shelby@niehs.nih.gov* (email). Information and comments received by May 6, 2002 will be made available to the CERHR staff and the expert panel for consideration in the evaluation.

The CERHR also invites nominations of qualified scientists to serve on the expert panel for the ethylene glycol/propylene glycol evaluation. Panelists are primarily drawn from the CERHR Expert Registry and/or the nomination of other scientists who meet the criteria for listing in that registry. Criteria for listing in the CERHR Expert Registry include: formal academic training and experience in a relevant scientific field, publications in peer-reviewed journals, membership in relevant professional societies, certification by an appropriate scientific Board or other entities, and participation in similar committee activities. All panel members serve as individual experts in their specific areas of expertise, not as representatives of their employer or other organization. Scientists on the expert panel will represent a wide range of expertise including developmental toxicology, reproductive toxicology, epidemiology, general toxicology, pharmacokinetics, exposure assessment, and biostatistics. Nominations received by May 6, 2002 will be considered for the Ethylene Glycol/Propylene Glycol Expert Panel and inclusion in the CERHR Expert Registry. Nominations should be forwarded to the CERHR at the address given above.

Additional Information about CERHR

The NTP and the NIEHS established the CERHR in June 1998 [FR (Vol. 63, No. 239, p. 68782, December 1998)]. The purpose of the CERHR is to provide scientifically-based, uniform assessments of the potential for adverse effects on reproduction and development caused by agents to which humans may be exposed. The CERHR also serves as a resource for information on various environmental exposures and their potential to affect pregnancy and child development. Its Web site (*http://cerhr.niehs.nih.gov*) has information about common concerns related to fertility, pregnancy and the health of unborn children, and links to other resources for information about public health.

The CERHR follows a formal, open process for the selection and review of chemicals nominated for evaluation of potential reproductive and/or developmental hazards. This process includes an evaluation of the chemical(s) by an external scientific panel that follows specific guidelines in conducting its assessment and provides multiple opportunities for public input. As a final step in the process, the CERHR publishes a NTP–CERHR report on each chemical that includes the expert panel report, public comments, and a NTP brief. A summary of the review process was recently published in the **Federal Register** (Vol. 66, No. 136, pp. 37047–37048, July 16, 2001). The process and guidelines are posted on the CERHR Web site and are available in hard copy by contacting the CERHR (address provided above). The CERHR welcomes the nomination of chemicals to be considered for future evaluation or qualified scientists for its expert registry. These nominations can be made through the CERHR Web site or by contacting the CERHR directly (see address above).

Dated: February 8, 2002.

Samuel H. Wilson,

Deputy Director, National Institute of Environmental Health Sciences.

[FR Doc. 02–5138 Filed 3–4–02; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Current List of Laboratories Which Meet Minimum Standards To Engage in Urine Drug Testing for Federal Agencies

AGENCY: Substance Abuse and Mental Health Services Administration, HHS.

ACTION: Notice.

SUMMARY: The Department of Health and Human Services notifies Federal agencies of the laboratories currently certified to meet standards of Subpart C of Mandatory Guidelines for Federal Workplace Drug Testing Programs (59 FR 29916, 29925). A notice listing all currently certified laboratories is published in the **Federal Register** during the first week of each month. If any laboratory's certification is suspended or revoked, the laboratory will be omitted from subsequent lists until such time as it is restored to full certification under the Guidelines.

If any laboratory has withdrawn from the National Laboratory Certification Program during the past month, it will be listed at the end, and will be omitted from the monthly listing thereafter.

This notice is also available on the internet at the following websites: *http://workplace.samhsa.gov; http://www.drugfreeworkplace.gov.*

FOR FURTHER INFORMATION CONTACT: Mrs. Giselle Hersh or Dr. Walter Vogl, Division of Workplace Programs, 5600 Fishers Lane, Rockwall 2 Building, Room 815, Rockville, Maryland 20857; Tel.: (301) 443–6014, Fax: (301) 443–3031.

SUPPLEMENTARY INFORMATION: Mandatory Guidelines for Federal Workplace Drug Testing were developed in accordance with Executive Order 12564 and section 503 of Pub. L. 100–71. Subpart C of the Guidelines, ''Certification of Laboratories Engaged in Urine Drug Testing for Federal Agencies,'' sets strict standards which laboratories must meet in order to conduct urine drug testing for Federal agencies. To become certified an applicant laboratory must undergo three rounds of performance testing plus an on-site inspection.

To maintain that certification a laboratory must participate in a quarterly performance testing program plus periodic, on-site inspections.

Laboratories which claim to be in the applicant stage of certification are not to be considered as meeting the minimum requirements expressed in the HHS