Written comments and recommendations for the proposed information collections must be mailed within 60 days of this notice directly to the OS Paperwork Clearance Officer designated at the following address: Department of Health and Human Services, Office of the Secretary, Assistant Secretary for Budget, Technology, and Finance, Office of Information and Resource Management, Attention: Naomi Cook (0990–0128), Room 531–H, 200 Independence Avenue, SW., Washington, DC 20201.

Dated: April 19, 2004.

**Robert E. Polson,**

*Office of the Secretary, Paperwork Reduction Act Reports Clearance Officer.*

[FR Doc. 04–9669 Filed 4–28–04; 8:45 am]

**BILLING CODE 4168–17–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

### National Task Force on Fetal Alcohol Syndrome and Fetal Alcohol Effect: Conference Call Meeting

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

*Name:* National Task Force on Fetal Alcohol Syndrome and Fetal Alcohol Effect (NTFFASFAE).

*Time and Date:* 2 p.m.–3 p.m., e.s.t., May 13, 2004.

*Place:* The conference call will originate at the National Center on Birth Defects and Developmental Disabilities (NCBDDD), in Atlanta, Georgia. Please *see* **SUPPLEMENTARY INFORMATION** for details on accessing the conference call.

*Status:* Open to the public, limited only by the availability of telephone ports.

*Purpose:* The Secretary is authorized by the Public Health Service Act, section 399G, (42 U.S.C. 280f, as added by Public Law 105–392) to establish a National Task Force on Fetal Alcohol Syndrome and Fetal Alcohol Effect to: (1) Foster coordination among all governmental agencies, academic bodies and community groups that conduct or support Fetal Alcohol Syndrome (FAS) and Fetal Alcohol Effect (FAE) research, programs and surveillance; and (2) to otherwise meet the general needs of populations actually or potentially impacted by FAS and FAE.

*Matters to be Discussed:* The Task Force will convene via conference call to: (1) Discuss and approve the recent revisions made to the Recommendations on Diagnostic and Referral Criteria for Fetal Alcohol Syndrome, (2) develop new Task Force working groups, and (3) obtain updates on recent motions passed by the Task Force related to the IDEA (Individuals with Disabilities Education Act) reauthorization and an FAS education requirement for teachers.

Agenda items are subject to change as priorities dictate.

*Supplementary Information:* This conference call is scheduled to begin at 2 p.m., eastern standard time. To participate in the conference call, please dial 1–866–453–4348 and enter conference code 602246. You will then be automatically connected to the call.

*Contact Person for More Information:* R. Louise Floyd, DSN, RN, Designated Federal Official, National Center on Birth Defects and Developmental Disabilities, CDC, 1600 Clifton Road, NE., (E–86), Atlanta, Georgia 30333, telephone 404/498–3923, fax 404/498–3550.

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

Dated: April 23, 2004.

**Bill J. Atkinson,**

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

[FR Doc. 04–9694 Filed 4–28–04; 8:45 am]

**BILLING CODE 4163–18–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES (DHHS)

### Centers for Disease Control and Prevention

### Advisory Committee for Injury Prevention and Control

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

*Name:* Advisory Committee for Injury Prevention and Control, and its subcommittees, the Science and Program Review Subcommittee and the Subcommittee on Intimate Partner Violence and Sexual Assault Science and Program Review Subcommittee (SPRS).

*Times and Dates:* 6:30 p.m.–9 p.m., May 17, 2004; 8 a.m.–5:30 p.m., May 18, 2004; 8 a.m.–10 a.m., May 19, 2004.

*Place:* Hyatt Regency Atlanta, 265 Peachtree Street, NE., Atlanta, Georgia 30303.

*Status:*

Open: 6:30 p.m.–7 p.m., May 17, 2004.

Closed: 7 p.m.–9 p.m., May 17, 2004.

Closed: 8 a.m.–5:30 p.m., May 18, 2004.

Open: 8 a.m.–10 a.m., May 19, 2004.

*Purpose:* The SPRS provides advice on the needs, structure, progress and performance of programs of the National Center for Injury Prevention and Control (NCIPC), as well as second-level scientific and programmatic review for applications for research grants, cooperative agreements, and training grants related to injury control and violence prevention, and recommends approval of projects that merit further consideration for funding support. The SPRS also advises on priorities for research to be supported by contracts, grants, and cooperative agreements and provides concept review of program proposals and announcements.

*Matters To Be Discussed:* The SPRS will be discussing the results of the NCIPC Initial Review Group's review and vote on grant applications submitted in response to 1 program announcement for Injury Control Research Centers and research grant applications submitted in response to 12 program announcements for individual research grant and cooperative agreement applications This portion of the meeting (7 p.m.–9 p.m., May 17, 2004, and 8 a.m.–5:30 p.m., May 18, 2004), will be closed to the public in accordance with provisions set forth in section 552b(c)(4) and (6), title 5, U.S.C., and the Determination of the Director, Management Analysis and Services Office, CDC, pursuant to Public Law 92–463.

Agenda items are subject to change as priorities dictate.

*Name:* Subcommittee on Intimate Partner Violence and Sexual Assault (SIPVSA).

*Time and Date:* 9 a.m.–11:30 a.m., May 19, 2004.

*Place:* Hyatt Regency Atlanta, 265 Peachtree Street, NE., Atlanta, Georgia 30303.

*Status:* Open to the public, limited only by the space available.

*Purpose:* To advise and make recommendations to the full advisory committee and the Director, NCIPC, regarding feasible goals for prevention and control of domestic and sexual violence. The SIPVSA will make recommendations regarding strategies, objectives, and priorities in programs, policies and research.

*Matters To Be Discussed:* The SIPVSA will review the NCIPC research agenda priorities and implementation related to intimate partner violence and sexual assault and discuss strategies for examining models for integration of intimate partner violence and sexual assault prevention into broader public health infrastructure and strategies.

*Name:* Advisory Committee for Injury Prevention and Control.

*Time and Dates:* 1 p.m.–5:30 p.m., May 19, 2004; 8:30 a.m.–2:30 p.m., May 20, 2004.

*Place:* Hyatt Regency Atlanta, 265 Peachtree Street, NE., Atlanta, Georgia 30303.

*Status:*

Closed: 1 p.m.–1:45 p.m., May 19, 2004.

Open: 1:45 p.m.–5:30 p.m., May 19, 2004.

Open: 8:30 a.m.–2:30 p.m., May 20, 2004.

*Purpose:* The Committee advises and makes recommendations to the Secretary, Health and Human Services, the Director, CDC, and the Director, NCIPC, regarding feasible goals for the prevention and control of injury. The Committee makes recommendations regarding policies, strategies, objectives, and priorities, and reviews progress toward injury prevention and control.

*Matters To Be Discussed:* Prior to the full committee meeting, there will be a brief meeting conducted by conference call of the Working Group on Injury Control and

Infrastructure Enhancement, a group formed to report to the full committee identifying gaps and suggesting ways to enhance injury prevention efforts. The working group will focus on defining injury infrastructure and developing a simple mechanism to assess current efforts underway throughout the injury field to enhance that infrastructure. Starting at 1 p.m., May 19, through 1:45 p.m., the full committee will vote on the results of secondary review. This portion of the meeting will be closed to the public in accordance with provisions set forth in section 552(b)(4) and (6), title 5 U.S.C., and the Determination of the Director, Management Analysis and Services Office, CDC, pursuant to Pub L. 92–463. Following the closed session, the meeting will open to the public for an update on Center activities from the Director, NCIPC; reports from the Subcommittees and Working Group; state infrastructure development; and discussion on how NCIPC can support the recommendations of CDC's Futures Initiative.

Agenda items are subject to change as priorities dictate.

*Contact Person for More Information:* Ms. Louise Galaska, Executive Secretary, ACIPC, NCIPC, CDC, 4770 Buford Highway, NE., M/S K02, Atlanta, Georgia 30341–3724, telephone (770) 488–4694.

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

Dated: April 23, 2004.

**Bill J. Atkinson,**

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

[FR Doc. 04–9693 Filed 4–28–04; 8:45 am]

**BILLING CODE 4163–18–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

### Submission for OMB Review; Comment Request; Application for the Pharmacology Research Associate Program

**SUMMARY:** Under the provisions of section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Institute of General Medical Sciences (NIGMS), the National Institutes of Health (NIH) has submitted to the Office of Management and Budget (OMB) a request to review and approve the information colleciton listed below. This proposed information collection was previously published in the Federal Register on February 13, 2004, pages 7236–7237, and allow 60 days for public comment. No public comments were received. The purpose of this notice is to allow an additional 30 days for public comment. The National Institutes of Health may not conduct or sponsor, and the respondent is not required to respond to, an information collection that has been extended, revised, or implemented on or after October 1, 1995, unless it displays a currently valid OMB control number.

*Proposed Collection: Title:* Application for the Pharmacology Research Associate Program. *Type of Information Collection Request:* Extension of a currently approved collection. *Need and Use of Information Collection:* The Pharmacology Research Associate (PRAT) Program will use the applicant and referee information to award opportunities for training and experience in laboratory or clinical investigation to individuals with a Ph.D. degree in pharmacology or a related science, M.D., or other professional degree through appointments as PRAT Fellows at the National Institutes of Health or the Food and Drug Administration. The goal of the program is to develop leaders in pharmacological reserach for key positions in academic, industrial, and Federal research laboratories. *Frequency of Response:* Once a year. *Affected Public:* Individuals or households; Businesses or other for-profit.

The annual reporting burden is as follows:

| Type and Number of Respondents | Estimated Number of Responses Per Respondent | Estimated Total Responses | Average Burden Hours per Responses | Estimated Total Annual Burden Hours Requested |
|---|---|---|---|---|
| Applicants 50 | 1 | 50 | 2.00 | 100 |
| Referees 150 | 1 | 150 | 0.167 | 25 |

*Total Number of Respondents:* 200.

*Total Number of Responses:* 200.

*Total Hours:* 125.

The annualized cost to respondents is estimated at:

*Applicants:* $5,500.00

*Referees:* $1,250.00

There are no Capital Costs, Operating Costs, and/or Maintenance Costs to report.

*Request for Comments:* Written comments and/or suggestions from the public and affected agencies should address one or more of the following points: (1) Evaluate whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility; (2) Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) Enhance the quality, utility, and clarity of the information to be collected; and (4) Minimize the burden of the collection of information on those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

*Direct Comments to OMB:* Written comments and/or suggestions regarding the item(s) contained in this notice, especially regarding the estiamted public burden and associated response time, should be directed to the: Office of Manageament and Budget, Office of Regulatory Affairs, New Executive Office Building, Room 10235, Washington, DC 20503, Attention: Desk Officer for NIH. To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact: Ms. Sally Lee, NIGMS, NIH, Natcher Building, Room 2AN–18H, 45 Center Drive, MSC 6200, Bethesda, MD 20892–6200, or call non-toll-free number 301–594–2755 or e-mail your request, including your address to *LeeS@nigms.nih.gov*.

*Comments Due Date:* Comments regarding this information collection are best assured of having their full effect if received within 30 days of the date of this publication.

Dated: April 21, 2004.

**Sally Lee,**

*Deputy Executive Officer, National Institute of General Medical Sciences.*

[FR Doc. 04–9683 Filed 4–28–04; 8:45 am]

**BILLING CODE 4140–01–M**