

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.

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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 collection 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 research 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 estimated public burden and associated response time, should be directed to the: Office of Management 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]

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