information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

#1 Type of Information Collection Request: Extension of a Currently Approved Collection;

Title of Information Collection: HHS Acquisition Regulation—Solicitation and Contracts;

Form/OMB No.: OS–0990–0115;

Use: Information is needed to evaluate feasibility of contractor(s) scientific or technical approach, management plan, and cost to accomplish the program or services required by the government.

Frequency: Recordkeeping, Reporting;

Affected Public: State, local, or tribal governments and not-for-profit institutions;

Annual Number of Respondents: 5,357*;*

Total Annual Responses: 5,357;

Average Burden Per Response: 1 hour;

Total Annual Hours: 883,905

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, access the HHS Web site address at *http://www.hhs.gov/oirm/infocollect/pending/* or e-mail your request, including your address, phone number, OMB number, and OS document identifier, to *naomi.cook@hhs.gov*, or call the Reports Clearance Office on (202) 690–6162. 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 (0990–0115), Room 531–H, 200 Independence Avenue, SW., Washington, DC 20201.

Dated: July 20, 2004.

Robert E. Polson,

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

[FR Doc. 04–17115 Filed 7–27–04; 8:45 am]

BILLING CODE 4168–17–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2004N–0159]

Schering Corp. et al.; Withdrawal of Approval of 92 New Drug Applications and 49 Abbreviated New Drug Applications; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a notice that appeared in the **Federal Register** of May 5, 2004 (69 FR 25124). The document announced the withdrawal of approval of 92 new drug applications (NDAs) and 49 abbreviated new drug applications (ANDAs). The document inadvertently withdrew approval of ANDA 88–584 for DHCplus (dihydrocodeine bitartrate, acetaminophen, and caffeine) Capsules, 356.4 milligrams, held by Purdue Frederick Co., One Stamford Forum, Stamford, CT 06901–3431. FDA confirms that approval of ANDA 88–584 is still in effect.

DATES: The notice published on May 5, 2004 (69 FR 25124) as corrected by this document has a date of June 4, 2004.

FOR FURTHER INFORMATION CONTACT: Florine P. Purdie, Center for Drug Evaluation and Research (HFD–7), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–594–2041.

In FR Doc. 04–10194 appearing on page 25124 in the issue of Wednesday, May 5, 2004, the following correction is made: On page 25130, in the table, the entry for NDA 88–584 is removed.

Dated: June 2, 2004.

Steven Galson,

Acting Director, Center for Drug Evaluation and Research.

[FR Doc. 04–17110 Filed 7–27–04; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

The National Institutes of Health

Submission for OMB Review; Comment Request; Brain Power! The NIDA Junior Scientist Program and the Companion Program, Brain Power! Challenge

SUMMARY: Under the provisions of section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Institute on Drug Abuse (NIDA), the National Institutes of Health (NIH) has submitted to the Office of Management and Budget (OMB) a request for review and approval of the information collection listed below. This proposed information collection was previously published in the **Federal Register** on April 1, 2004, page 17194, and allowed 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: Brain Power! The NIDA Junior Scientist Program, for grades K–5, and the companion program for Middle School, the Brain Power! Challenge. *Type of Information Collection Request:* NEW. *Need and Use of Information Collection:* This is a request for a three-year clearance to evaluate the effectiveness of the Brain Power! Program's ability to: (1) Increase student's knowledge about the biology of the brain and the neurobiology of drug addiction, (2) increase positive attitudes toward science, careers in science, science as an enjoyable endeavor, and the use of animals in research; and stimulate interest in scientific careers; and (3) engender more realistic perceptions of scientists as being from many races, ages, and genders. The secondary goals of the evaluation are to determine the Program's impact on attitudes and intentions toward drug use. The findings will provide valuable information concerning the goals of NIDA's Science Education Program of increasing scientific literacy and stimulating interest in scientific careers. In order to test the effectiveness of the evaluation, information will be collected from students before and after exposure to the curriculum with pre- and post-test self-report measures. Surveys will also be administered to teachers after the completion of the program to examine ease and fidelity of implementation, as well as impact in knowledge and understanding of the neurobiology of addiction. Surveys will be administered to parents to obtain parental reaction and opinion on the materials and the degree to which parents find the curriculum informative and appropriate.

Frequency of Response: On occasion. *Affected Public:* Elementary and middle school students, teachers, and parents. *Type of Respondents:* Students, Teachers, and Parents. The reporting