Information and Regulatory Affairs, Attention Department of Justice Desk Officer, Washington, DC 20503. Additionally, comments may be submitted to OMB via facsimile to (202) 395–5806. Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address one or more of the following four points:

- —Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;
- —Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;
- —Enhance the quality, utility, and clarity of the information to be collected; and
- —Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses.

Overview of This Information Collection

- (1) Type of Information Collection: Extension of a currently approved collection.
- (2) Title of the Form/Collection: Firearms Transaction Record Low Volume Part I Over-the-Counter and Part II Intra-State Non-Over-the-Counter.
- (3) Agency form number, if any, and the applicable component of the Department sponsoring the collection: Form Number: ATF F 4473 (5300.24) Part I (LV) and ATF F 4473 (5300.25) Part II (LV) and ATF REC 7570/2. Bureau of Alcohol, Tobacco, Firearms and Explosives.
- (4) Affected public who will be asked or required to respond, as well as a brief abstract: Primary: Business or other forprofit. Other: Individual or households. The forms are used by low volume firearms dealers to record acquisition and disposition of firearms and to determine the eligibility of buyers to receive firearms. The forms are part of the licensee's permanent record and may be used to trace firearms.
- (5) An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond/reply: It is estimated that 1,000

respondents will complete a 20-minute form.

(6) An estimate of the total public burden (in hours) associated with the collection: There are an estimated 1,666 annual total burden hours associated with this collection.

If additional information is required contact: Brenda E. Dyer, Department Clearance Officer, United States Department of Justice, Justice Management Division, Policy and Planning Staff, Patrick Henry Building, Suite 1600, 601 D Street NW., Washington, DC 20530.

Dated: May 4, 2005.

Brenda E. Dyer,

Department Clearance Officer, Department of Justice.

[FR Doc. 05–9181 Filed 5–6–05; 8:45 am] BILLING CODE 4410–FY–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated June 28, 2004, and published in the **Federal Register** on July 13, 2004, (69 FR 42068), Eli-Elsohly Laboratories, Inc., Mahmoud A. Elsohly PhD., 5 Industrial Park Drive, Oxford, Mississippi 38655, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed in Schedule I and II:

Drug	Schedule
Codeine (9050)	

The company plans to manufacture the listed controlled substances for use in analysis and drug test standards.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Eli-Elsohly Laboratories, Inc. to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Eli-Elsohly Laboratories, Inc. to ensure that the company's registration is consistent with the public interest. The investigation has included inspection and testing of the company's physical security systems, verification

of the company's compliance with state and local laws, and a review of the company's background and history. Therefore, pursuant to 21 U.S.C. 823, and in accordance with 21 CFR 1301.33(a), the above named company is granted registration as a bulk manufacturer of the basic classes of controlled substances listed.

Dated: May 2, 2005.

William J. Walker,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 05–9177 Filed 5–6–05; 8:45 am] BILLING CODE 4410–09–P

NATIONAL FOUNDATION ON THE ARTS AND THE HUMANITIES

Federal Council on the Arts and the Humanities, Arts and Artifacts Indemnity Panel Advisory Committee; Notice of Meeting

Pursuant to the provisions of the Federal Advisory Committee Act (Pub. L. 92–463 as amended) notice is hereby given that a meeting of the Arts and Artifacts Indemnity Panel of the Federal Council on the Arts and the Humanities will be held at 1100 Pennsylvania Avenue, NW., Washington, DC 20506, in Room 714, from 9 a.m. to 5 p.m., on Monday, May 23, 2005.

The purpose of the meeting is to review applications for Certificates of Indemnity submitted to the Federal Council on the Arts and the Humanities for exhibitions beginning after July 1, 2005.

Because the proposed meeting will consider financial and commercial data and because it is important to keep values of objects, methods of transportation and security measures confidential, pursuant to the authority granted me by the Chairman's Delegation of Authority to Close Advisory Committee Meetings, dated July 19, 1993, I have determined that the meeting would fall within exemption (4) of 5 U.S.C. 552(b) and that it is essential to close the meeting to protect the free exchange of views and to avoid interference with the operations of the Committee.

It is suggested that those desiring more specific information contact Acting Advisory Committee Management Officer, Michael McDonald, 1100 Pennsylvania Avenue,