

burden of the proposed collection of the information, including the validity of the methodology and assumptions used;

(3) 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 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

1. Type of Information Collection:

Approval of a Revised Collection

2. Title of the Forms:

FBI National Academy: End-of-Session Student Course Questionnaire

FBI National Academy: General Remarks Questionnaire

3. Agency Form Number, if any, and the applicable component of the department sponsoring the collection:

Form Number: 1110-0050

Sponsor: Training Division, Federal Bureau of Investigation (FBI), Department of Justice (DOJ)

4. Affected Public who will be asked or required to respond, as well as a brief abstract:

Primary: FBI National Academy students that represent state and local police and sheriffs' departments, military police organizations, and federal law enforcement agencies from the United States and over 150 foreign nations.

Brief Abstract: This collection is requested by FBI National Academy. These questionnaires have been designed to collect feedback from National Academy students regarding their courses and instructors. The results are used to help determine if the National Academy program is functioning as intended and meeting its goals and objectives. We will utilize the students' comments to improve the current curriculum.

5. An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:

Approximately 1,000 FBI National Academy students per year will respond to two types of questionnaires. (1) FBI National Academy: End-of-Session Student Course Questionnaire and (2) FBI National Academy: General Remarks Questionnaire. It is predicted we will receive a 75% response rate for both

questionnaires. Each student will respond to seven Student Course questionnaires—one for each course they completed. The average time for reading the questionnaire directions is estimated to be two (2) minutes; the time to complete each questionnaire is estimated to be approximately 13 minutes. Thus the total time to complete one Student Course questionnaire is 15 minutes and 105 minutes for all seven questionnaires.

For the FBI National Academy: General Remarks Questionnaire, students will respond to one questionnaire. The average time for reading the questionnaire directions is estimated to be two (2) minutes; the time to complete the questionnaire is estimated to be approximately 10 minutes. Thus the total time to complete the General Remarks Questionnaire is 12 minutes.

The total estimated time for both questionnaires per respondent is approximately 117 minutes or about 2 hours.

6. An estimate of the total public burden (in hours) associated with the collection:

Given that approximately 75% of those surveyed (or 750) will respond, the total public burden for completing all questionnaires is 1462.5 hours.

For additional information, contact: Jerri Murray, Department Clearance Officer, U.S. Department of Justice, Policy and Planning Staff, Justice Management Division, Two Constitution Square, 145 N Street NE., Room 3W0-1407B, Washington, DC 20530.

Dated: January 30, 2014.

Jerri Murray,
Department Clearance Officer for PRA, U.S. Department of Justice.

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Application; S & B Pharma, Inc.

Pursuant to 21 CFR 1301.34(a), this is notice that on March 18, 2013, S & B Pharma, Inc., DBA Norac Pharma, 405 S. Motor Avenue, Azusa, California 91702-3232, made application to the Drug Enforcement Administration (DEA) for registration as an importer of the following basic classes of controlled substances:

Drug	Schedule
4-Anilino-N-phenethyl-4-piperidine (8333).	II
Tapentadol (9780)	II
Fentanyl (9801)	II

The company plans to import the listed controlled substances for internal use, and to manufacture bulk intermediates for sale to its customers.

Any bulk manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic classes of controlled substances listed in schedules I or II, which fall under the authority of section 1002(a)(2)(B) of the Act (21 U.S.C. 952(a)(2)(B)) may, in the circumstances set forth in 21 U.S.C. 958(i), file comments or objections to the issuance of the proposed registration and may, at the same time, file a written request for a hearing on such application pursuant to 21 CFR 1301.43, and in such form as prescribed by 21 CFR 1316.47.

Any such written comments or objections should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODW), 8701 Morrisette Drive, Springfield, Virginia 22152; and must be filed no later than March 6, 2014.

This procedure is to be conducted simultaneously with and independent of the procedures described in 21 CFR 1301.34(b), (c), (d), (e), and (f). As noted in a previous notice published in the **Federal Register** on September 23, 1975, 40 FR 43745-46, all applicants for registration to import a basic class of any controlled substance in schedules I or II are, and will continue to be, required to demonstrate to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, that the requirements for such registration pursuant to 21 U.S.C. 958(a); 21 U.S.C. 823(a); and 21 CFR 1301.34(b), (c), (d), (e), and (f) are satisfied.

Dated: January 15, 2014.

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

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