

Research and Production Act of 1993, 15 U.S.C. 4301 *et seq.* (“the Act”), Title Council of America (“TCA”) has filed written notifications simultaneously with the Attorney General and the Federal Trade Commission disclosing (1) the name and principal place of business of the standards development organization and (2) the nature and scope of its standards development activities. The notifications were filed for the purpose of invoking the Act’s provisions limiting the recovery of antitrust plaintiffs to actual damages under specified circumstances.

Pursuant to section 6(b) of the Act, the name and principal place of business of the standards development organization is: Title Council of America, Anderson, SC. The nature and scope of TCA’s standards development activities are: Standard specifications for the installation of ceramic tile, for ceramic tile installation materials, and for ceramic tile including tile, porcelain tile, glass tile, and special purpose tile.

Dorothy B. Fountain,

Deputy Director of Operations, Antitrust Division.

[FR Doc. 04–25076 Filed 11–9–04; 8:45 am]

BILLING CODE 4410–11–M

DEPARTMENT OF JUSTICE

Antitrust Division

Notice Pursuant to the National Cooperative Research and Production Act of 1993—Wild Bird Feeding Industry

Notice is hereby given that, on September 23, 2004, pursuant to section 6(a) of the National Cooperative Research and Production Act of 1993, 15 U.S.C. 4301 *et seq.* (“the Act”), Wild Bird Feeding Industry (“WBF”) has filed written notifications simultaneously with the Attorney General and the Federal Trade Commission disclosing (1) the name and principal place of business of the standards development organization and (2) the nature and scope of its standards development activities. The notifications were filed for the purpose of invoking the Act’s provisions limiting the recovery of antitrust plaintiffs to actual damages under specified circumstances.

Pursuant to section 6(b) of the Act, the name and principal place of business of the standards development organization is: Wild Bird Feeding Industry, Sioux Falls, SD. The nature and scope of WBF’s standards development activities are: Investigating and developing standards of identity and

quality for products sold for wild birds. These products include feeders, houses, baths and accessories, and seed and other food.

Dorothy B. Fountain,

Deputy Director of Operations, Antitrust Division.

[FR Doc. 04–25084 Filed 11–9–04; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated May 21, 2004, and published in the **Federal Register** on June 3, 2004, (69 FR 31411), American Radiolabeled Chemicals, Inc., 101 Arc Drive, St. Louis, Missouri 63146, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Gamma hydroxybutyric acid (2010).	I
Dimethyltryptamine (7435)	I
Dihydromorphine (9145)	I
Cocaine (9041)	II
Codeine (9050)	II
Hydromorphone (9150)	II
Benzoyllecgonine (9180)	II
Ecgonine (9180)	II
Meperidine (9230)	II
Metazocine (9240)	II
Morphine (9300)	II
Oxymorphone (9652)	II

The company plans to manufacture in bulk, small quantities of the listed controlled substances as radiolabeled compounds.

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 American Radiolabeled Chemicals, Inc. to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated American Radiolabeled Chemicals, 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, the above named company is granted registration as a bulk manufacturer of

the basic classes of controlled substances listed.

Dated: November 1, 2004.

William J. Walker,

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

[FR Doc. 04–25103 Filed 11–9–04; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to 21 CFR 1301.33(a), Title 21 of the Code of Federal Regulations (CFR), this is notice that on July 28, 2004, Guilford Pharmaceuticals, Inc., 6611 Tributary Street, Baltimore, Maryland 21224, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of Cocaine (9041), a basic class of controlled substance in Schedule II.

The company plans to manufacture a cocaine derivative to be used as an intermediate for the production of Dopascan Injection. Cocaine derivatives are a Schedule II controlled substance in the cocaine basic class.

Any other such applicant and any person who is presently registered with DEA to manufacture such a substance may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

Any such comments or objections may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, DC 20537, Attention: DEA **Federal Register** Representative Office of Liaison and Policy (ODLR) and must be filed no later than January 10, 2005.

Dated: November 1, 2004.

William J. Walker,

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

[FR Doc. 04–25104 Filed 11–9–04; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on August 16,

2004, Noramco Inc., 1440 Olympic Drive, Athens, Georgia 30601, made application by letter to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of Codeine (9050), a basic class of controlled substance in Schedule II.

The company plans to utilize codeine to produce small quantities of naturally occurring codeine impurities for use in quality assurance and internal testing of the finished products.

Any other such applicant and any person who is presently registered with DEA to manufacture such a substance may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

Any such comments or objections may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, DC 20537, Attention: Federal Register Representative, Office of Liaison and Policy (ODLR) and must be filed no later than January 10, 2005.

Dated: November 1, 2004.

William J. Walker,

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

[FR Doc. 04-25102 Filed 11-9-04; 8:45 am]

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DEPARTMENT OF LABOR

Office of the Secretary

Proposed Collection; Comment Request

ACTION: Notice.

SUMMARY: The Department of Labor, as part of its continuing effort to reduce paperwork and respondent burden, conducts a pre-clearance consultation program to provide the general public and Federal agencies an opportunity to comment on proposed and/or continuing collections of information in accordance with the Paperwork Reduction Act of 1995 (PRA95) (44 U.S.C. 3506(c)(2)(A)). This program helps to ensure that requested data can be provided in the desired format, reporting burden (time and financial resources) is minimized, collection instruments are clearly understood, and the impact of collection requirements can be properly assessed. Currently, Departmental Management is soliciting comments concerning the proposed Information Collection Request (ICR) for the Assessment of Compliance Assistance Activities Generic Clearance.

A copy of the ICR can be obtained by contacting the individual listed below in the **ADDRESSES** section of this notice.

DATES: Written comments must be submitted to the office listed in the **ADDRESSES** section below on or before January 10, 2005.

ADDRESSES: Send comments to Barbara Bingham, Office of the Assistant Secretary for Policy, 200 Constitution Avenue, NW., Room S-2312, Washington, DC 20210. Ms. Bingham can be reached on 202-693-5080 (this is not a toll-free number) or by e-mail at bingham-barbara@dol.gov.

SUPPLEMENTARY INFORMATION:

I. Background

The Department of Labor (DOL) proposes to assess and measure self-reported changes in behavior through surveys of workers, employers and other stakeholders. These surveys will provide feedback on compliance assistance documents and materials, onsite consultation visits, telephone and technical assistance, Web sites, partnerships and alliances, and compliance assistance seminars and workshops delivered by DOL across the country to the regulated community. This feedback will help DOL agencies improve the future quality and delivery of compliance assistance tools and services. This generic clearance allows agencies to gather information from both Federal and non-Federal users.

II. Desired Focus of Comments

The Department of Labor is particularly interested in comments which:

- 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 agency's 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., permitted electronic submissions of response.

III. Current Actions

DOL agencies have conducted few surveys designed to assess changes in

worker, employer and stakeholder behavior as a result of the compliance assistance received. DOL proposes to seek approval of this collection of information for a three year period.

Type of Review: New collection of information.

Agency: Office of the Assistant Secretary for Policy, Office of Compliance Assistance.

Title: Information Collection Request for the Assessment of Compliance Assistance Activities Generic Clearance.

OMB Number: 1225-0NEW.

Affected Public: Individuals and households; business or other for-profit; not-for-profit institutions; farms; Federal Government; and State, Local, or Tribal Government.

Frequency: On occasion.

Number of Respondents: 29,995.

Annual Responses: 9,998.

Average Time Per Response: Varies by survey/evaluation with an average of 13 minutes per survey.

Total Annual Burden Hours: 2,202.

Total Burden Cost (capital/startup): \$0.

Total Burden Cost (operating/maintenance): \$0.

Comments submitted in response to this notice will be summarized and/or included in the request for Office of Management and Budget approval of the information collection request; they also will become a matter of public record.

Signed at Washington, DC this 4th day of November, 2004.

David Gray,

Acting Assistant Secretary, Office of the Assistant Secretary for Policy.

[FR Doc. 04-25048 Filed 11-9-04; 8:45 am]

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DEPARTMENT OF LABOR

Employee Benefits Security Administration

[Prohibited Transaction Exemption 2004-17; (Exemption Application No. D-11223) et al.]

Grant of Individual Exemptions; Linda Ann Smith, M.D. Profit Sharing Plan and Trust (the Plan)

AGENCY: Employee Benefits Security Administration, Labor.

ACTION: Grant of individual exemptions.

SUMMARY: This document contains exemptions issued by the Department of Labor (the Department) from certain of the prohibited transaction restrictions of the Employee Retirement Income Security Act of 1974 (the Act) and/or the Internal Revenue Code of 1986 (the Code).