factors in 21 U.S.C. 823(a) and 952(a), and determined that the registration of SA INTL GMBH C/O., Sigma Aldrich Co. LLC., to import the basic classes of controlled substances is consistent with the public interest, and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971, at this time. DEA has investigated SA INTL GMBH C/O., Sigma Aldrich Co. LLC., 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. 952(a) and 958(a), and in accordance with 21 CFR 1301.34, the above named company is granted registration as an importer of the basic classes of controlled substances listed.

Dated: November 5, 2012.

Joseph T. Rannazzisi,

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

[FR Doc. 2012–27571 Filed 11–9–12; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application; Cedarburg Pharmaceuticals, Inc.

Pursuant to § 1301.33(a), of Title 21 of the Code of Federal Regulations (CFR), this is notice that on September 21, 2012, Cedarburg Pharmaceuticals, Inc., 870 Badger Circle, Grafton, Wisconsin 53024, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the following basic classes of controlled substances:

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

The company plans to manufacture the listed controlled substances in bulk for distribution to its customers. Regarding the drug code (8333), the company plans to use this controlled substance to manufacture another controlled substance.

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

Any such written comments or objections should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 8701 Morrissette Drive, Springfield, Virginia 22152; and must be filed no later than January 14, 2013.

Dated: November 5, 2012.

Joseph T. Rannazzisi,

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

[FR Doc. 2012–27572 Filed 11–9–12; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application; Johnson Matthey Inc.

Pursuant to § 1301.33(a), Title 21 of the Code of Federal Regulations (CFR), this is notice that on September 10, 2012, Johnson Matthey Inc., Custom Pharmaceuticals Department, 2003 Nolte Drive, West Deptford, New Jersey 08066–1742, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Drug	Schedule
Gamma Hydroxybutyric Acid (2010).	I
Tetrahydrocannabinols (7370)	I
Dihydromorphine (9145)	I
Difenoxin (9168)	I
Propiram (9649)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Lisdexamfetamine (1205)	II
Methylphenidate (1724)	II
Nabilone (7379)	II
Cocaine (9041)	II
Codeine (9050)	II
Dihydrocodeine (9120)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Diphenoxylate (9170)	II
Ecgonine (9180)	II
Hydrocodone (9193)	II
Meperidine (9230)	II
Methadone (9250)	II
Methadone intermediate (9254)	II
Morphine (9300)	II
Thebaine (9333)	II
Oxymorphone (9652)	II
Noroxymorphone (9668)	II
Alfentanil (9737)	II
Remifentanil (9739)	II
Sufentanil (9740)	II
Fentanyl (9801)	II

The company plans to manufacture the listed controlled substances in bulk for sale to its customers.

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

Any such written comments or objections should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 8701 Morrissette Drive, Springfield, Virginia 22152; and must be filed no later than January 14, 2013.

Dated: November 5, 2012.

Joseph T. Rannazzisi,

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

[FR Doc. 2012–27565 Filed 11–9–12; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration; Wildlife Laboratories, Inc.

By Notice dated April 17, 2012, and published in the **Federal Register** on July 31, 2012, 77 FR 45378, Wildlife Laboratories, Inc., 1230 W. Ash Street, Suite D, Windsor, Colorado 80550, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Carfentanil (9743), a basic class of controlled substance listed in schedule II.

The company plans to manufacture the above listed controlled substance for sale to veterinary pharmacies, zoos, and for other animal and wildlife applications.

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 Wildlife Laboratories, Inc., to manufacture the listed basic class of controlled substance is consistent with the public interest at this time. DEA has investigated Wildlife 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(a), and in accordance with 21 CFR

1301.33, the above named company is granted registration as a bulk manufacturer of the basic class of controlled substance listed.

Dated: November 5, 2012.

Joseph T. Rannazzisi,

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

[FR Doc. 2012–27568 Filed 11–9–12; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration; Nektar Therapeutics

By Notice dated July 17, 2012, and published in the **Federal Register** on July 26, 2012, 77 FR 43862, Nektar Therapeutics, 1112 Church Street, Huntsville, Alabama 35801, made application by letter to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Fentanyl (9801), a basic class of controlled substance listed in schedule II.

The company plans to manufacture the listed controlled substance in support of product development.

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 Nektar Therapeutics to manufacture the listed basic class of controlled substance is consistent with the public interest at this time. DEA has investigated Nektar Therapeutics 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 review of the company's background and history. Therefore, pursuant to 21 U.S.C. 823(a), and in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of the basic class of controlled substance listed.

Dated: November 5, 2012.

Joseph T. Rannazzisi,

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

[FR Doc. 2012–27567 Filed 11–9–12; 8:45 am]

BILLING CODE 4410–09–P

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

[Notice 12–095]

NASA Advisory Council; Audit, Finance and Analysis Committee; Meeting

AGENCY: National Aeronautics and Space Administration.

ACTION: Notice of Meeting.

SUMMARY: In accordance with the Federal Advisory Committee Act, Public Law 92–463, as amended, the National Aeronautics and Space Administration announces a meeting of the Audit, Finance and Analysis Committee of the NASA Advisory Council.

DATES: Monday, November 26, 2012, 9:00 a.m.–5:15 p.m., Local Time.

ADDRESS: NASA Headquarters. 300 E Street SW., Conference Room 8E40, Washington DC 20546.

FOR FURTHER INFORMATION CONTACT: Ms. Charlene Williams, Office of the Chief Financial Officer, National Aeronautics and Space Administration Headquarters, Washington, DC 20546. Phone: 202–358–2183.

SUPPLEMENTARY INFORMATION: The agenda for the meeting includes briefings on the following topics:

- FY 2012 Financial Statement Audit
- FY 2013 Financial Management Initiatives
- Administrative Savings
- NASA Budget
- Government Accounting Office High Risk List
- Financial System Initiative

It is imperative that the meeting be held on these dates to accommodate the scheduling priorities of the key participants. Attendees will be requested to sign a register and to comply with NASA security requirements, including the presentation of a valid picture ID to NASA Security before access to NASA Headquarters. Foreign Nationals attending this meeting will be required to provide a copy of their passport and visa in addition to providing the following information no later than November 21, 2012: full name; gender; date/place of birth; citizenship; visa information (number, type, expiration date); passport information (number, country, expiration date); employer/affiliation information (name of institution, address, country, telephone); title/position of attendee; and home address to Charlene Williams at fax: (202) 358–4336. U.S. Citizens and Permanent Residents (green card holders) are requested to submit their name and affiliation 3 working days prior to the meeting to Charlene Williams.

Patricia D. Rausch,

Advisory Committee Management Officer, National Aeronautics and Space Administration.

[FR Doc. 2012–27487 Filed 11–9–12; 8:45 am]

BILLING CODE P

NATIONAL CREDIT UNION ADMINISTRATION

Sunshine Act; Notice of Agency Meeting

TIME AND DATE: 10:00 a.m., Thursday, November 15, 2012.

PLACE: Board Room, 7th Floor, Room 7047, 1775 Duke Street, Alexandria, VA 22314–3428.

STATUS: Open.

MATTERS TO BE CONSIDERED: 1. NCUA's 2013 Operating Budget.

2. NCUA/NCUSIF Overhead Transfer Rate.

3. Federal Credit Unions' Operating Fee Scale.

4. Board Briefing on the Estimated 2013 Premium Ranges for the NCUSIF and the Corporate Stabilization Fund.

FOR FURTHER INFORMATION CONTACT: Mary Rupp, Secretary of the Board, Telephone: 703–518–6304.

Mary Rupp,

Board Secretary.

[FR Doc. 2012–27648 Filed 11–8–12; 4:15 pm]

BILLING CODE 7535–01–P

NATIONAL SCIENCE FOUNDATION

Astronomy and Astrophysics Advisory Committee #13883; Notice of Meeting

In accordance with the Federal Advisory Committee Act (Pub. L. 92–463, as amended), the National Science Foundation announces the following Astronomy and Astrophysics Advisory Committee (#13883) meeting:

Date and Time: November 30, 2012, 8:30 a.m.–5:00 p.m.; December 1, 2012, 8:30 a.m.–1:00 p.m.

Place: National Science Foundation, Room 1235, Stafford I Building, 4201 Wilson Blvd., Arlington, VA, 22230.

Type of Meeting: Open.

Contact Person: Dr. Jim Ulvestad, Division Director, Division of Astronomical Sciences, Suite 1045, National Science Foundation, 4201 Wilson Blvd., Arlington, VA 22230. Telephone: 703–292–7165.

Purpose of Meeting: To provide advice and recommendations to the National Science Foundation (NSF), the National Aeronautics and Space Administration (NASA) and the U.S. Department of Energy (DOE) on issues