

Agency in full compliance with section 510 of the FD&C Act before January 9, 2014. As previously stated, drug products covered by this document that are currently marketed but not listed with the Agency on the date of this document must, as of the effective date of this document, have approved applications before their shipment in interstate commerce. Moreover, any person or firm that has submitted or submits an application but has yet to receive approval for such products is still responsible for full compliance with this document.

V. Discontinued Products

Some firms may have previously discontinued manufacturing or distributing products covered by this document without removing them from the listing of their products under section 510(j) of the FD&C Act. Other firms may discontinue manufacturing or distributing listed products in response to this document. Firms are required to electronically update the listing of their products under section 510(j) of the FD&C Act to reflect discontinuation of unapproved products covered by this document (21 CFR 207.21(b)). Questions on electronic drug listing updates should be sent to: eDRLS@fda.hhs.gov. In addition to the required update, firms can also notify the Agency of product discontinuation by sending a letter, signed by the firm's chief executive officer and fully identifying the discontinued product(s), including the product NDC number(s), and stating that the manufacturing and/or distribution of the product(s) has (have) been discontinued. The letter should be sent electronically to Astrid Lopez-Goldberg (see **ADDRESSES**). FDA plans to rely on its existing records, including its drug listing records, the results of any subsequent inspections, or other available information when it targets violations for enforcement action.

VI. Reformulated Products

FDA cautions firms against reformulating their products into unapproved new drugs without codeine sulfate, codeine phosphate, or dihydrocodeine bitartrate, and marketing them under the same name or substantially the same name (including a new name that contains the old name) in anticipation of an enforcement action based on this document. As stated in the Marketed Unapproved Drugs CPG, FDA intends to give higher priority to enforcement actions involving unapproved drugs that are reformulated to evade an anticipated FDA enforcement action. In addition, reformulated products marketed under a

name previously identified with a different active ingredient have the potential to confuse healthcare practitioners and harm patients.

Dated: January 6, 2014.

Leslie Kux,

Assistant Commissioner for Policy.

[FR Doc. 2014-00257 Filed 1-9-14; 8:45 am]

BILLING CODE 4160-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Statement of Organization, Functions and Delegations of Authority

This notice amends Part R of the Statement of Organization, Functions and Delegations of Authority of the Department of Health and Human Services (HHS), Health Resources and Services Administration (HRSA) (60 FR 56605, as amended November 6, 1995; as last amended at 78 FR 42089-42090 dated July 15, 2013).

This notice reflects organizational changes in the Health Resources and Services Administration. This notice corrects the administrative codes for the Bureau of Clinician Recruitment and Service (RU); the Division of Regional Operations (RU2) and the Office of Business Operations (RU3).

Chapter RU—Bureau of Clinician Recruitment and Service

Section RU-10, Organization

Delete and replace in its entirety.

The Office of the Associate Administrator (RU) is headed by the Associate Administrator, Bureau of Clinician Recruitment and Service (BCRS), who reports directly to the Administrator, Health Resources and Services Administration. BCRS includes the following components:

- (1) Office of the Associate Administrator (RU);
- (2) Office of Legal and Compliance (RU1);
- (3) Division of Regional Operations (RU2);
- (4) Office of Business Operations (RU3);
- (5) Division of National Health Service Corps (RU5);
- (6) Division of Nursing and Public Health (RU6);
- (7) Division of External Affairs (RU7);
- (8) Division of Policy and Shortage Designation (RU8); and
- (9) Division of Program Operations (RU9).

Section RU-30, Delegations of Authority

All delegations of authority and re-delegations of authority made to HRSA officials that were in effect immediately prior to this reorganization, and that are consistent with this reorganization, shall continue in effect pending further re-delegation.

This reorganization is effective upon date of signature.

Dated: December 26, 2013.

Mary K. Wakefield,

Administrator.

[FR Doc. 2014-00221 Filed 1-9-14; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Indian Health Service

Organization, Function, and Delegations of Authority; Part G; Proposed Functional Statement

AGENCY: Indian Health Service, HHS.

ACTION: Notice of change in name of an organizational component.

SUMMARY: The Indian Health Service is announcing the name change of the Aberdeen Area Indian Health Service to the Great Plains Area Indian Health Service at the request of Tribes served by the Aberdeen Area Indian Health Service.

FOR FURTHER INFORMATION CONTACT: Ms. Mona Galpin, Office of Management Services, Management Policy and Internal Control Staff, 801 Thompson Avenue, TMP Suite 625A, Rockville, MD 20852, Telephone 301-443-2650.

Section GF-10, Indian Health Service Area Offices—Organization

An Area Office is a second echelon organization under the direction of an Area Director, who reports to the IHS Director.

Indian Health Service Area Offices of the Indian Health Service in alphabetical order:

- Alaska Area Office (GFB)
- Albuquerque Area Office (GFC)
- Bemidji Area Office (GFE)
- Billings Area Office (GFF)
- California Area Office (GFG)
- Great Plains Area Office (GFA)
- Nashville Area Office (GFH)
- Navajo Area Office (GFJ)
- Oklahoma Area Office (GFK)
- Phoenix Area Office (GFL)
- Portland Area Office (GFM)
- Tucson Area Office (GFN)

Dated: December 6, 2013.

Yvette Roubideaux,

Acting Director, Indian Health Service.

[FR Doc. 2014-00264 Filed 1-9-14; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center for Scientific Review; Notice of Closed Meetings

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of the following meetings.

The meetings will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: Center for Scientific Review Special Emphasis Panel; RFAs: PAR12-138 and PAR13-026—NHLBI Systems Biology.

Date: February 4–5, 2014.

Time: 10:00 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (Virtual Meeting).

Contact Person: Ai-Ping Zou, MD, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4118, MSC 7814, Bethesda, MD 20892, 301-408-9497, zouai@csr.nih.gov.

Name of Committee: Brain Disorders and Clinical Neuroscience Integrated Review Group; Clinical Neuroplasticity and Neurotransmitters Study Section.

Date: February 6–7, 2014.

Time: 8:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Melrose Hotel, 2430 Pennsylvania Ave. NW., Washington, DC 20037.

Contact Person: Suzan Nadi, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5217B, MSC 7846, Bethesda, MD 20892, 301-435-1259, nadis@csr.nih.gov.

Name of Committee: Biobehavioral and Behavioral Processes Integrated Review Group; Cognition and Perception Study Section.

Date: February 6–7, 2014.

Time: 8:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: The Westin St. Francis, 335 Powell Street, San Francisco, CA 94102.

Contact Person: Dana Jeffrey Plude, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3176, MSC 7848, Bethesda, MD 20892, (301) 435-2309, pluded@csr.nih.gov.

Name of Committee: Risk, Prevention and Health Behavior Integrated Review Group; Psychosocial Development, Risk and Prevention Study Section.

Date: February 6–7, 2014.

Time: 8:00 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Renaissance Mayflower Hotel, 1127 Connecticut Avenue NW., Washington, DC 20036.

Contact Person: Anna L Riley, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3114, MSC 7759, Bethesda, MD 20892, 301-435-2889, rileyann@csr.nih.gov.

Name of Committee: Endocrinology, Metabolism, Nutrition and Reproductive Sciences Integrated Review Group Clinical and Integrative Diabetes and Obesity Study Section.

Date: February 6, 2014.

Time: 8:00 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Residence Inn Bethesda, 7335 Wisconsin Avenue, Bethesda, MD 20814.

Contact Person: Nancy Sheard, SCD, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 6046-E, MSC 7892, Bethesda, MD 20892, 301-408-9901, sheardn@csr.nih.gov.

(Catalogue of Federal Domestic Assistance Program Nos. 93.306, Comparative Medicine; 93.333, Clinical Research, 93.306, 93.333, 93.337, 93.393–93.396, 93.837–93.844, 93.846–93.878, 93.892, 93.893, National Institutes of Health, HHS).

Dated: January 6, 2014.

Carolyn Baum,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2014-00162 Filed 1-9-14; 8:45 am]

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DEPARTMENT OF HOMELAND SECURITY

U.S. Customs and Border Protection

Agency Information Collection Activities: Vessel Entrance or Clearance Statement

AGENCY: U.S. Customs and Border Protection, Department of Homeland Security.

ACTION: 30-Day Notice and request for comments; Extension of an existing collection of information: 1651-0019.

SUMMARY: U.S. Customs and Border Protection (CBP) of the Department of

Homeland Security will be submitting the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act: Vessel Entrance or Clearance Statement (CBP Form 1300). This is a proposed extension of an information collection that was previously approved. CBP is proposing that this information collection be extended with no change to the burden hours. This document is published to obtain comments from the public and affected agencies. This proposed information collection was previously published in the **Federal Register** (76 FR 39114) on October 29, 2013, allowing for a 60-day comment period. This notice allows for an additional 30 days for public comments. This process is conducted in accordance with 5 CFR 1320.10.

DATES: Written comments should be received on or before February 10, 2014 to be assured of consideration.

ADDRESSES: Interested persons are invited to submit written comments on this proposed information collection to the Office of Information and Regulatory Affairs, Office of Management and Budget. Comments should be addressed to the OMB Desk Officer for Customs and Border Protection, Department of Homeland Security, and sent via electronic mail to oir_submission@omb.eop.gov or faxed to (202) 395-5806.

FOR FURTHER INFORMATION CONTACT: Requests for additional information should be directed to Tracey Denning, U.S. Customs and Border Protection, Regulations and Rulings, Office of International Trade, 90 K Street NE., 10th Floor, Washington, DC 20229-1177, at 202-325-0265.

SUPPLEMENTARY INFORMATION: CBP invites the general public and other Federal agencies to comment on proposed and/or continuing information collections pursuant to the Paperwork Reduction Act of 1995 (Pub. L. 104-13; 44 U.S.C. 3507). The comments should address: (a) Whether the collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimates of the burden of the collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; (d) ways to minimize the burden including the use of automated collection techniques or the use of other forms of information technology; and (e) the annual costs burden to respondents or record keepers from the collection of information (a