assistance and consultation to the activities within ESRO; and (6) coordinates, develops, and monitors implementation of program initiatives.

Senior Executive Compensation and Performance Activity (CAJQG2). (1) Provides advisory services, and technical assistance on pay and compensation guidelines in accordance with OPM rules and regulations, HHS and CDC/ATSDR established pay and compensation recommendation policies, and procedures; (2) provides expert human resources advisory services and technical assistance support to the CDC performance review boards and compensation committees; (3) reviews actions for statutory and regulatory compliance; (4) manages strategic recruitment, relocation, and retention incentives to facilitate attraction of a quality, diverse workforce to ensure accomplishment of the CDC/ATSDR missions; (5) provides performance management training for all SES and Title 42 executives with emphasis on performance systems, timelines, supervisory and employee responsibilities; (6) provides guidance on establishing performance plans, conducting mid-year reviews, and conducting final performance rating discussions and closing performance plans; (7) develops and maintains a standard Department-wide performance management system and forms for executives; (8) conducts reviews of SES performance plans and appraisals and provides feedback; (9) prepares and submits SES performance system certification request to OPM and OMB; (10) processes performance awards and performance-based pay adjustments; (11) provides advice, assistance, templates and training workshops on performance award and Presidential Rank Award requirements; (12) manages the HHS Executive Development Program, including developmental activities, rotational assignments, and the Candidate Development Program; and (13) advises on development of executive succession planning activities.

Title 42 Staffing and Recruitment Activity (CAJQG3). (1) Provides leadership, technical assistance, guidance, and consultation in the administration of policies and procedures for appointment of individuals through the distinguished consultants, experts, consultants, and fellows under Title 42 appointment authorities; and (2) administers and manages the Guest Researcher and Oak Ridge Institute for Science and Education Program.

Immigration Activity (CAJQG4). (1) provides technical guidance and visa-assistance for employment based, CDC-sponsored visas; (2) administers and Manages the Exchange Visitor Program; (3) works closely with the US Office of Exchange and Cultural Affairs, US Citizenship and Immigration Services, US Department of Homeland Security, US Department of State, Office of the Secretary/DHHS, and US Department of Labor) to facilitate immigration procedures; (4) reviews, processes and files H–1B, O–1, and Green Card (I–140) Petitions with the U.S. Citizenship and Immigration Services; (5) provides advisory services and guidance on employment based green card petitions in the Alien of Extraordinary Ability category; (6) issues Certificate of Eligibility for J–1 Exchange Visitor Status through the Student and Exchange Visitor Information System to non US citizens seeking CDC J–1 visa sponsorship; (7) coordinates and provides consultations and guidance on Interested Government Agency Waivers; (8) provides Immigration Training Workshops to CDC/ATSDR Administrative Staff; and (9) determines the appointment mechanism, legal status, and work authorizations for non U.S. citizens through the Visitors Management System.

Sherri A. Berger,

Chief Operating Officer, Centers for Disease Control and Prevention.

[FR Doc. 2016–28225 Filed 11–22–16; 8:45 am]

BILLING CODE 4160–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Request for Nominations of Potential Reviewers To Serve on the Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP)

The Centers for Disease Control and Prevention (CDC) is soliciting nominations for possible membership on the Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP) in the National Center for Injury Prevention and Control (NCIPC).

The Disease, Disability, and Injury Prevention and Control Special Emphasis Panel provides advice and guidance to the Secretary, Department of Health and Human Services (HHS); the Director, Centers for Disease Control and Prevention (CDC), and the Administrator, Agency for Toxic Substances and Disease Registry (ATSDR) regarding the concept review, scientific and technical merit of grant and cooperative agreement assistance applications, and contract proposals relating to the causes, prevention, and control of diseases, disabilities, injuries, and impairments of public health significance; exposure to hazardous substances in the environment; health promotion and education; and other related activities that promote health and well-being.

Nominations are being sought for individuals who have expertise and qualifications necessary to contribute to the accomplishment of NCIPC SEP objectives. Reviewers with expertise in the following research fields for injury and violence prevention are sought to serve on the NCIPC SEPs, for research and evaluation related, but not limited to: Child abuse and neglect, prescription drug overdose, intimate partner violence, motor vehicle injury, older adult falls, self-directed violence, sexual violence, traumatic brain injury, youth sports concussion, and youth violence. Reviewers with expertise in the following methodological fields for injury and violence prevention are also sought to serve on the CDC SEP for NCIPC programs: economic evaluation, etiology, implementation and translation, intervention research, policy evaluation, program evaluation, qualitative research design, quantitative research design, statistics, and surveillance.

Members and Chairs shall be selected by the Secretary, HHS, or other official to whom the authority has been delegated, on an ''as needed'' basis in response to specific applications being reviewed with expertise to provide advice. Members will be selected from authorities in the various fields of prevention and control of diseases, disabilities, and injuries. Members of other chartered HHS advisory committees may serve on the panel if their expertise is required. Consideration is given to professional training and background, points of view represented, and upcoming applications to be reviewed by the committee.

Information about nominated potential reviewers will be maintained in the NCIPC Reviewer and Advisor Database. The work of reviewers' appointed to NCIPC SEPS includes the initial review, discussion, and written critique and evaluation of applications. This work will enable the CDC/NCIPC to fulfill its mission of funding meritorious research that provides vital knowledge about underlying risk and protective factors and strategies for violence and injury prevention *http://www.cdc.gov/injury/index.html.*

The U.S. Department of Health and Human Services policy stipulates that

committee membership be balanced in terms of points of view represented and the committee's function. Appointments shall be made without discrimination on the basis of age, race, ethnicity, gender, sexual orientation, gender identity, HIV status, disability, and cultural, religious, or socioeconomic status. Nominees must be U.S. citizens appointed to serve on a CDC SEP and can be full-time employees of the U.S. Government. Current participation on CDC federal workgroups or prior experience serving on another federal advisory committee does not disqualify a reviewer, except for service on the Board of Scientific Counselors, NCIPC. However, HHS policy is to avoid excessive individual service on advisory committees and multiple committee memberships. Reviewers appointed to the SEP, CDC are not considered Special Government Employees, and will not be required to file financial disclosure reports.

Nominees interested in serving as a potential reviewer on a SEP, CDC for NCIPC programs should submit the following items:

• Current *curriculum vitae,* including complete contact information (name, affiliation, mailing address, telephone number, and email address).

Nomination materials must be postmarked by March 31, 2017 and sent by U.S. mail to: NCIPC Extramural Research Program Office (ERPO): Centers for Disease Control and Prevention, 1600 Clifton Road NE., Mailstop F–63, Atlanta, Georgia 30329 or to the ERPO Mailbox *NCIPC_ERPO@cdc.gov.*

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Elaine L. Baker,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 2016–28207 Filed 11–22–16; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Advisory Committee on Breast Cancer in Young Women, Centers for Disease Control and Prevention: Notice of Charter Renewal

This gives notice under the Federal Advisory Committee Act (Pub. L. 92–463) of October 6, 1972, that the Advisory Committee on Breast Cancer in Young Women, Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS), has been renewed for a 2-year period through June 17, 2018.

For information, contact Temeika L. Fairley, Ph.D., Designated Federal Officer, Advisory Committee on Breast Cancer in Young Women, HHS, CDC, 4770 Buford Highway NE., Mailstop K52, Atlanta, Georgia 30341, telephone 770/488–4518, fax 770/488–4760.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Elaine L. Baker,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 2016–28206 Filed 11–22–16; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2015–D–4033]

Nonprescription Sunscreen Drug Products—Format and Content of Data Submissions; Guidance for Industry; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of availability.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing the availability of a guidance for industry entitled ''Nonprescription Sunscreen Drug Products—Format and Content of Data Submissions.'' This guidance addresses FDA's current thinking on the format and content of information provided to support a request for a determination whether a nonprescription sunscreen active ingredient is generally recognized as safe and effective (GRASE), as provided under the Federal Food, Drug, and Cosmetic Act (the FD&C Act), as amended by the Sunscreen Innovation Act (SIA).

DATES: Submit either electronic or written comments on Agency guidances at any time.

ADDRESSES: You may submit comments as follows:

Electronic Submissions

Submit electronic comments in the following way:

• *Federal eRulemaking Portal: http://www.regulations.gov.* Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to *http://www.regulations.gov* will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on *http://www.regulations.gov.*

• If you want to submit a comment with confidential information that you do not wish to be made available to the public submit the comment as a written/paper submission and in the manner detailed (see ''Written/Paper Submissions'' and ''Instructions'').

Written/Paper Submissions

Submit written/paper submissions as follows:

• *Mail/Hand delivery/Courier (for written/paper submissions):* Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

• For written/paper comments submitted to the Division of Dockets Management, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in ''Instructions.''

Instructions: All submissions received must include the Docket No. FDA–2015–D–4033 for ''Nonprescription Sunscreen Drug Products—Format and Content of Data Submissions; Guidance for Industry; Availability.'' Received comments will be placed in the docket