

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](mailto: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.*

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## 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.*

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## 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