which they promote as a resource for consumers in search of financial products such as loans and insurance. In numerous instances, Respondents have described the content on the website, including their rate tables, star ratings, and rankings of the companies offering these financial products, as ''objective,'' ''honest,'' ''accurate,'' and ''unbiased.'' Contrary to their claims, Respondents have provided financial services companies with higher numerical rankings or star ratings and higher positions on rate tables based on compensation. Respondents also have added or removed companies from their content based on compensation.

In addition, Respondents have touted positive consumer reviews about their company and website that, in fact, were written by LendEDU employees or their friends, family members, and associates. Of 126 reviews of LendEDU on the third-party review platform Trustpilot, 90% were written or made up by LendEDU employees or their family, friends, or other associates. Respondents also have reposted and touted the Trustpilot reviews on LendEDU's website, as well as fake reviews written by LendEDU employees who purport to be, but are not, actual users.

The proposed order will prevent Proposed Respondents from engaging in similar acts or practices. Part I would prohibit Proposed Respondents from making the challenged and related misrepresentations. Part II would require Proposed Respondents to disclose the influence of compensation on representations made on its website and to disclose material connections among the Proposed Respondents and the various parties represented on the website. Part III would require Proposed Respondents, jointly and severally, to pay to the Commission $350,000 within 8 days of the effective date of the Order.

Part IV sets out additional requirements related to the monetary relief. Part V requires Proposed Respondents to provide sufficient customer information to enable the Commission to efficiently administer consumer redress. Part VI is an order distribution provision that requires Proposed Respondents to provide the order to current and future principals, officers, directors, and LLC managers and members, as well as current and future managers, agents and representatives who participate in certain duties related to the subject matter of the proposed complaint and order, and to secure statements acknowledging receipt of the order. Part VII requires Proposed Respondents to submit a compliance report one year after the order is entered. It also requires Proposed Respondents to notify the Commission of corporate changes that may affect compliance obligations within 14 days of such a change.

Part VIII requires Proposed Respondents to maintain and upon request make available certain compliance-related records, including accounting records and unique websites. Part IX requires Proposed Respondents to submit additional compliance reports within 10 business days of a written request by the Commission. Part X is a provision ''sunsetting'' the order after twenty (20) years, with certain exceptions.

The purpose of this analysis is to aid public comment on the proposed order. It is not intended to constitute an official interpretation of the complaint or proposed order, or to modify in any way the proposed order's terms.

By direction of the Commission.

April J. Tabor,

Acting Secretary.

[FR Doc. 2020–02798 Filed 2–11–20; 8:45 am]

BILLING CODE 6750–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2020–N–0255]

Patient-Focused Drug Development for Vitiligo; Public Meeting; Request for Comments

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public meeting; request for comments.

SUMMARY: The Food and Drug Administration (FDA, the Agency, or we) is announcing the following public meeting entitled ''Patient-Focused Drug Development for Vitiligo.'' The purpose of the public meeting is to allow FDA to obtain patient perspectives on the impact of vitiligo on daily life, patient views on treatment approaches, and decision factors considered when selecting a treatment.

DATES: The public meeting will be held on March 30, 2020, from 1 p.m. to 5 p.m. Submit either electronic or written comments on this public meeting by June 1, 2020. See the **SUPPLEMENTARY INFORMATION** section for registration date and information.

ADDRESSES: The public meeting will be held at FDA's White Oak Campus, 10903 New Hampshire Ave., Bldg. 31 Conference Center, the Great Room (Rm. 1503), Silver Spring, MD 20993. Entrance for the public meeting participants (non-FDA employees) is through Building 1 where routine security check procedures will be performed. For parking and security information, please refer to *https://www.fda.gov/AboutFDA/WorkingatFDA/BuildingsandFacilities/WhiteOakCampusInformation/ucm241740.htm.*

You may submit comments as follows. Please note that late, untimely filed comments will not be considered. Electronic comments must be submitted on or before June 1, 2020. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of June 1, 2020. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are postmarked or the delivery service acceptance receipt is on or before that date.

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal: https://www.regulations.gov.* Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to *https://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 *https://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):* Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Dockets Management Staff, 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–2020–N–0255 for ''Patient-Focused Drug Development on Vitiligo; Public Meeting; Request for Comments.'' Received comments, those filed in a timely manner (see **ADDRESSES**), will be placed in the docket and, except for those submitted as ''Confidential Submissions,'' publicly viewable at *https://www.regulations.gov* or at the Dockets Management Staff between 9 a.m. and 4 p.m., Monday through Friday.

• Confidential Submissions—To submit a comment with confidential information that you do not wish to be made publicly available, submit your comments only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states ''THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION.'' The Agency will review this copy, including the claimed confidential information, in its consideration of comments. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on *https://www.regulations.gov.* Submit both copies to the Dockets Management Staff. If you do not wish your name and contact information to be made publicly available, you can provide this information on the cover sheet and not in the body of your comments and you must identify this information as ''confidential.'' Any information marked as ''confidential'' will not be disclosed except in accordance with 21 CFR 10.20 and other applicable disclosure law. For more information about FDA's posting of comments to public dockets, see 80 FR 56469, September 18, 2015, or access the information at: *https://www.govinfo.gov/content/pkg/FR-2015-09-18/pdf/2015-23389.pdf.*

Docket: For access to the docket to read background documents or the electronic and written/paper comments received, go to *https://www.regulations.gov* and insert the docket number, found in brackets in the heading of this document, into the ''Search'' box and follow the prompts and/or go to the Dockets Management Staff, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Shannon Cole, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 51, Rm. 6306, Silver Spring, MD 20993–0002, 301–796–9208, *PatientFocused@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION:

I. Background

This meeting will provide FDA the opportunity to obtain patient and patient representative input on the aspects of vitiligo that matter most to patients, including how it affects daily life, and on current approaches to treating vitiligo. Vitiligo is an autoimmune disease that causes the loss of skin color. The loss of color can affect skin, hair, and other areas of the body. The area affected by color loss can range in individual patients from small discrete areas to near total involvement. Although there is no cure or FDA-approved treatment for repigmentation, there are available therapies, such as prescription medications or non-drug therapies, which may often be used to manage aspects of vitiligo. FDA is interested in patients' (including adult and pediatric patients) perspectives on: (1) The impact of their vitiligo; (2) treatment approaches; and (3) decision factors considered when selecting a treatment.

The questions that will be asked of patients and patient representatives at the meeting are listed in the following section and organized by topic. For each topic, a brief initial patient panel discussion will begin the dialogue. This discussion will be followed by a facilitated discussion inviting comments from other patients and patient representatives. In addition to input generated through this public meeting, FDA is interested in receiving patient and patient representative input addressing these questions through written comments, which can be submitted to the public docket (see **ADDRESSES**). When submitting comments, if you are commenting on behalf of a patient, please indicate that you are doing so and answer the following questions as much as possible from the patient's perspective.

FDA will post the agenda and other meeting materials approximately 5 days before the meeting at: *https://www.fda.gov/drugs/news-events-human-drugs/public-meeting-patient-focused-drug-development-vitiligo-03302020-03302020.*

II. Topics for Discussion at the Public Meeting

Topic 1: Health Effects and Daily Impacts That Matter Most to Patients

1. Which aspects of vitiligo have the most significant impact on your life? (Examples may include depigmentation, itching, sensitivity to sunlight, etc.)

2. Are there specific activities that are important to you but that you cannot do at all or as fully as you would like because of your vitiligo? (Examples of activities may include participating in social events, playing sports, being outside in the sunlight, etc.)

How does your vitiligo and its impacts affect your daily life on the best days? On the worst days?

3. How has your vitiligo changed over time?

a. How has your vitiligo changed from childhood to adulthood (such as vitiligo severity, disease acceptance)?

b. Would you define your vitiligo today as being well-managed?

4. What worries you most about your vitiligo?

Is there a particular body area affected by vitiligo (such as face, hands, limbs) that is of most concern to you?

Topic 2: Patients' Perspectives on Current Approaches to Treatment

1. What are you currently doing to help treat your vitiligo? (Examples may include prescription medicines, over-the-counter products, and other therapies, including non-drug therapies such as diet modification.)

How has your treatment regimen changed over time, and why?

2. How well does your current treatment regimen treat the most significant aspects of your vitiligo? For example, how well do your treatments improve your ability to do specific activities?

3. What are the most significant downsides to your current treatments, and how do they affect your daily life? (Examples of downsides may include bothersome side effects, depigmentation of affected area is more noticeable, hospital treatments, etc.)

4. Assuming there is no complete cure for your vitiligo, what specific things would you look for in an ideal treatment for your vitiligo?

Is there a particular body area affected by vitiligo (such as face, hands, limbs) that you would prioritize for treatment?

5. What factors do you consider when making decisions about selecting a course of treatment?

III. Participating in the Public Meeting

Registration: To register for the public meeting, visit *https://vitiligo-pfdd.eventbrite.com.* Please register by March 23, 2020. Persons without access to the internet can call 301–796–9208 to register. If you are unable to attend the meeting in person, you can register to view a live webcast of the meeting. You will be asked to indicate in your registration if you plan to attend in person or via the webcast. Please

provide complete contact information for each attendee, including name, title, affiliation, address, email, and telephone.

Registration is free and based on space availability, with priority given to early registrants. Persons interested in attending this public meeting must register by March 23, 2020, 11:59 p.m. Eastern Time. Early registration is recommended because seating is limited; therefore, FDA may limit the number of participants from each organization. If time and space permit, onsite registration on the day of the public meeting will be provided beginning at 12 p.m.

If you need special accommodations due to a disability, please contact Shannon Cole (see **FOR FURTHER INFORMATION CONTACT**) no later than March 23, 2020.

Panelist Selection: Patients or patient representatives who are interested in presenting comments as part of the initial panel discussions will be asked to indicate in their registration which topic(s) they wish to address. These patients or patient representatives also will be asked to send *PatientFocused@fda.hhs.gov* a brief summary of responses to the topic questions by March 9, 2020. Panelists will be notified of their selection approximately 7 days before the public meeting. We will try to accommodate all patients and patient stakeholders who wish to speak, either through the panel discussion or audience participation; however, the duration of comments may be limited by time constraints.

Open Public Comment: There will be time allotted during the meeting for open public comment. Signup for this session will be on a first-come, first-served basis on the day of the meeting. Individuals and organizations with common interests are urged to consolidate or coordinate and request time for a joint presentation. No commercial or promotional material will be permitted to be presented or distributed at the public meeting.

Persons attending FDA's meetings are advised that FDA is not responsible for providing access to electrical outlets.

Streaming Webcast of the Public Meeting: This public meeting will also be webcast. Please register for the webcast by visiting *https://vitiligo_pfdd.eventbrite.com.*

If you have never attended a Connect Pro event before, test your connection at *https://collaboration.fda.gov/common/help/en/support/meeting_test.htm.* To get a quick overview of the Connect Pro program, visit *https://www.adobe.com/go/connectpro_overview.* FDA has verified the website addresses in this document, as of the date this document publishes in the **Federal Register**, but websites are subject to change over time.

Transcripts: Please be advised that as soon as a transcript of the public meeting is available, it will be accessible at *https://www.regulations.gov.* It may be viewed at the Dockets Management Staff (see **ADDRESSES**). A link to the transcript will also be available on the internet at *https://www.fda.gov/drugs/news-events-human-drugs/public-meeting-patient-focused-drug-development-vitiligo-03302020-03302020.*

Dated: February 6, 2020.

Lowell J. Schiller,

Principal Associate Commissioner for Policy.

[FR Doc. 2020–02767 Filed 2–11–20; 8:45 am]

BILLING CODE 4164–01–P

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, 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; PAR Panel: Alzheimer's Clinical Trials.

Date: March 10, 2020.

Time: 12:00 p.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

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

Contact Person: Unja Hayes, Ph.D., Scientific Review Officer, National Institutes of Health, Center for Scientific Review, 6701 Rockledge Drive, Bethesda, MD 20892, 301–827–6830, *unja.hayes@nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel; SBIR Small Business: Computational, Modeling, and Biodata Management.

Date: March 11, 2020.

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

Agenda: To review and evaluate grant applications.

Place: Embassy Suites at the Chevy Chase Pavilion, 4300 Military Road NW, Washington, DC 20015.

Contact Person: Allen Richon, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 6184, MSC 7892, Bethesda, MD 20892, 301–379–9351, *allen.richon@nih.hhs.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel; Small Business: Psycho/Neuropathology Lifespan Development, STEM Education.

Date: March 12–13, 2020.

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

Agenda: To review and evaluate grant applications.

Place: Residence Inn Pentagon City, 550 Army Navy Drive, Arlington, VA 22202.

Contact Person: Elia K. Ortenberg, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3108, Bethesda, MD 20892, 301–827–7189, *femiaee@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel; Small Business: Medical Imaging.

Date: March 12–13, 2020.

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

Agenda: To review and evaluate grant applications.

Place: Holiday Inn Bayside, 4875 North Harbor Drive, San Diego, CA 92106.

Contact Person: Leonid V. Tsap, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5128, MSC 7854, Bethesda, MD 20892, (301) 435–2507, *tsapl@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel; Small Business: Neuroscience Assay, Diagnostics and Animal Model Development.

Date: March 12–13, 2020.

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

Agenda: To review and evaluate grant applications.

Place: Residence Inn Washington, DC Downtown, 1199 Vermont Ave. NW, Washington, DC 20005.

Contact Person: Joseph G. Rudolph, Ph.D., Chief and Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5186, MSC 7844, Bethesda, MD 20892, 301–408–9098, *josephru@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel; PAR–17–190: Maximizing Investigators' Research Award for Early Stage Investigators (R35).

Date: March 12, 2020.

Time: 8:00 a.m. to 4:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Hyatt Regency Bethesda, Metro Center, 1, Bethesda, MD 20814.

Contact Person: Baishali Maskeri, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 2022, Bethesda, MD 20892, 301–827–2864, *maskerib@mail.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel; Fellowships: Oncology.