Current curriculum vitae, including complete contact information (telephone numbers, mailing address, and email address).

- Cover letter, including a description of the candidate qualifications and why the candidate would be a good fit for ACET.
- At least one letter of recommendation from person(s) not employed by the U.S. Department of Health and Human Services. (Candidates may submit letter(s) from current HHS employees if they wish, but at least one letter must be submitted by a person not employed by an HHS agency (e.g., CDC, NIH, FDA, etc.) (see https://www.hhs.gov/about/agencies/hhs-agencies-and-offices/index.html for a full list)).

Nominations may be submitted by the candidate him- or herself, or by a person/organization recommending the candidate.

The Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, 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.

Kalwant Smagh,
Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention.

SUPPLEMENTARY INFORMATION:

- **Request for Information**
  - State health agencies have a critical role in the identification and prevention of occupational illnesses and injuries. NIOSH has supported state agencies (primarily departments of public health and, in some cases, departments of labor) since the 1970s, through a combination of funding and technical assistance. Since that time, NIOSH has supported states to build capacity in occupational safety and health, ranging from the development of case-based surveillance to creating focused public health interventions addressing the occupational health needs of higher risk populations. The work of these state programs exemplifies the concept of “information for action” by ensuring that collection, analysis, interpretation, and dissemination of occupational health data are linked to prevention and control activities. Numerous examples of these successes can be found in the published literature, in state reports and on state websites, and NIOSH website topic pages.

In 2019, NIOSH funded 26 state occupational health surveillance programs of varying sizes and capacities. The portfolio of state based activities includes 49 projects addressing work related morbidity and mortality, exposure reduction, or special populations of interest. These states are funded through a research cooperative agreement mechanism. The most recent (2014) announcement can be found at the following web address: https://grants.nih.gov/grants/guide/pa-files/PAR-14-275.html.

For its state-based surveillance and C1 intervention cooperative agreements, NIOSH is considering switching from a research cooperative agreement approach to a non-research cooperative agreement approach. CDC generally defines public health research as an activity that develops or contributes to generalizable knowledge to improve public health practice; a non-research activity is one that is designed to identify and control a health problem or improve a public health program or service. A non-research mechanism could be a public health practice cooperative agreement or another cooperative agreement type, and may or may not be a better fit for the scope of activities ordinarily conducted by occupational health programs in a public health context.

Under the research mechanism currently used, submissions for funding are evaluated on the following criteria: Significance, investigators, innovation, approach, and environment. Under a non-research approach, proposals would likely be evaluated based upon how well the proposal identifies important occupational health burdens in the state; approach for tracking these concerns; relevance and potential impact of the public health actions proposed; and organizational capacity of the applicant to achieve the proposal.

This exploration of funding mechanism type presents an opportunity for NIOSH to receive stakeholder input and identify the best type from a programmatic, logistic, and administrative point of view. Exploring this and other approaches is recommended by the National Academies of Science, Engineering and Medicine in its report “A Smarter National Surveillance System for
Occupational Safety and Health in the 21st Century.”

To identify and assess different options, NIOSH plans to conduct the following activities: (1) Hold the public teleconference announced in this notice to receive comments regarding funding approaches for its state-based occupational health surveillance programs and (2) seek additional public comments through this docket.

NIOSH is interested in comments related to the funding mechanism as it relates to impact on the conduct of state agency activities, including comments on the following questions:

1. What are the advantages and disadvantages to the states if NIOSH continues using research cooperative agreements for funding of state occupational health surveillance programs?
2. What are the advantages and disadvantages to the states if NIOSH changed to using a non-research mechanism for funding state occupational health surveillance programs?
3. If the non-research mechanism would specifically prohibit the use of any funds for research, would this have a negative effect on state occupational health surveillance program development or direction? If so, please describe.
4. Only research cooperative agreements are covered by Certificates of Confidentiality that protect the confidentiality of sensitive information collected from research subjects by our grantees. Do states need or use these certificates?
5. Would a non-research cooperative agreement mechanism impact the ability of universities acting as bonafide agents of the states to apply and receive funding under this mechanism? If so, how?
6. Non-research proposals undergo “objective review,” which employs CDC reviewers in place of external peer reviewers. Scoring of applications would likely use the criteria described above (occupational health burdens in the state; approach for tracking these concerns; relevance and potential impact of the public health actions proposed; and organizational capacity of the state). Are there concerns related to these criteria or the use of objective review?
7. Is it possible that NIOSH will continue to employ an external peer review process for scoring of applications? Are there concerns related to the use of external peer review?

8. Using the principles of Burden, Need and Impact, the new Notice of Funding Opportunity will focus on surveillance activities that address the occupational safety and health burden of the applicant state. How will this directive impact the applying states?
9. The 2014 cooperative agreement (PAR–14–275) funded three programmatic levels (fundamental, fundamental plus, and expanded programs) to address the varying levels of surveillance capacity of applicant states. Should this 3-tier strategy be continued? If not, what other strategy might be considered?
10. How does the 3-tier funding strategy affect states’ ability to explore emerging occupational safety and health issues?
11. Occupational Health Indicators have been a central component of the NIOSH state-based surveillance program. What are the advantages and disadvantages to your state program of continuing to calculate and use the Occupational Health Indicators?

Public Meeting

NIOSH will hold a public teleconference meeting to solicit comments on the future funding mechanism of its state-based occupational health surveillance program. The meeting is open to the public, limited only by the capacity of 250 connections to the web-based conference.

Confirm your attendance to this meeting by sending an email to ksouza@cdc.gov by September 9, 2019. An email confirming registration will be sent from NIOSH and will include details needed to participate.

Requests to make a statement at the public meeting should be emailed to ksouza@cdc.gov by September 2, 2019. All requests to make statements should contain the name, address, telephone number, and relevant business affiliations of the presenter. Presenters will be assigned a 5-minute slot on the agenda. Oral statements only will be permitted—presentations of slides will not be permitted. NIOSH will confirm presentation requests by email, and will provide additional instructions regarding the presentation, including the approximate start time for the presentation.

If a presenter is not in attendance when his/her presentation is scheduled to begin, the remaining presenters will be heard in order. After the last scheduled presenter is heard, those who missed their opportunity may be allowed to present, limited by time available.

Attendees who wish to speak, but did not submit a request for the opportunity to make a presentation, may be given this opportunity after the scheduled presenters are heard, at the discretion of the presiding official and limited by time available. Those who do not have an opportunity to comment during the teleconference are encouraged to submit written comments to the NIOSH docket.

The public meeting will be recorded, transcribed, and posted without change to http://www.regulations.gov, including any personal information provided.

Frank J. Hearl,
Chief of Staff, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention, Department of Health and Human Services.

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BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–2567]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, HHS.

ACTION: Notice.

SUMMARY: The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS’ intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (the PRA), federal agencies are required to publish notices in the Federal Register concerning each proposed collection of information (including each proposed extension or reinstatement of an existing collection of information) and to allow 60 days for public comment on the proposed action. Interested persons are invited to send comments regarding our burden estimates or any other aspect of this collection of information, including the necessity and utility of the proposed information collection for the proper performance of the agency’s functions, the accuracy of the estimated burden, ways to enhance the quality, utility, and clarity of the information to be collected, and the use of automated collection techniques or other forms of

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See https://www.cdc.gov/niosh/programs/pps/bni.html for more information about Burden, Need and Impact.

See https://www.cste.org/page/OfHealthicators for more information about Occupational Health Indicators.

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