meeting participants not only to provide input, but also to respond to the input provided by others. A more specific agenda will be provided to meeting registrants before the meeting.

AHRQ requests that interested persons register with the PSO PPC to participate in the meeting. The contact at the PSO PPC is Mark Baliff who can be reached by telephone at (866) 571–7712 and by email at *SUPPORT@.PSOPPC.ORG.* Additional logistical information for the meeting is also available from the PSO PPC. The meeting space will accommodate approximately 150 participants. Interested persons are encouraged to register as soon as possible for the meeting. Non-registered individuals will be able to attend the meeting in person if space is available.

Prior to the meeting, AHRQ invites review of the technical specifications for Common Formats. The formats can be accessed through AHRQ's PSO Web site at *http://www.pso.AHRQ.GOV/formats/commonfmt.htm.* AHRQ is committed to continuing refinement of the Common Formats, and welcomes questions from prospective meeting participants and interested individuals on the technical specifications. These questions should be emailed to *SUPPORT@PSOPPC.ORG* no later than April 10th, 2013. AHRQ will use the input received at this meeting to further update and refine the Common Formats.

Dated: February 7, 2013.

Carolyn M. Clancy,

Director.

[FR Doc. 2013–03911 Filed 2–20–13; 8:45 am]

BILLING CODE 4160–90–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Patient Safety Organizations: Delisting for Cause for Independent Data Safety Monitoring, Inc.

AGENCY: Agency for Healthcare Research and Quality (AHRQ), HHS.

ACTION: Notice of Delisting.

SUMMARY: AHRQ has delisted Independent Data Safety Monitoring, Inc. due to its failure to correct a deficiency. The Patient Safety and Quality Improvement Act of 2005 (Patient Safety Act) authorizes the listing of PSOs, which are entities or component organizations whose mission and primary activity is to conduct activities to improve patient safety and the quality of health care delivery. HHS issued the Patient Safety and Quality Improvement Final Rule (Patient Safety Rule) to implement the Patient Safety Act. AHRQ administers the provisions of the Patient Safety Act and Patient Safety Rule relating to the listing and operation of PSOs.

DATES: The directories for both listed and delisted PSOs are ongoing and reviewed weekly by AHRQ. The delisting was effective at 12:00 Midnight ET (2400) on January 15, 2013.

ADDRESSES: Both directories can be accessed electronically at the following HHS Web site: *http://www.pso.AHRQ.gov/index.html.*

FOR FURTHER INFORMATION CONTACT: Eileen Hogan, Center for Quality Improvement and Patient Safety, AHRQ, 540 Gaither Road, Rockville, MD 20850; Telephone (toll free): (866) 403–3697; Telephone (local): (301) 427–1111; TTY (toll free): (866) 438–7231; TTY (local): (301) 427–1130; *Email: psoAHRQ.hhs.gov.*

SUPPLEMENTARY INFORMATION:

Background

The Patient Safety Act, Public Law 109–41,42 U.S.C. 299b–21 b–26, provides for the formation of PSOs, which collect, aggregate, and analyze confidential information regarding the quality and safety of health care delivery. The Patient Safety Rule, 42 CFR part 3, authorizes AHRQ, on behalf of the Secretary of HHS, to list as a PSO an entity that attests that it meets the statutory and regulatory requirements for listing. A PSO can be ''delisted'' by the Secretary if it is found no longer to meet the requirements of the Patient Safety Act and Patient Safety Rule. Section 3.108(d) of the Patient Safety Rule requires AHRQ to provide public notice when it removes an organization from the list of federally approved PSOs.

Independent Data Safety Monitoring, Inc. failed to respond to a Notice of Preliminary Finding of Deficiency sent by AHRQ pursuant to 42 CFR 3.108(a)(2) and a Notice of Proposed Revocation and Delisting sent by AHRQ pursuant to 42 CFR 3.108(a)(3)(iii)(C) which found that Independent Data Safety Monitoring, Inc. had not complied with its attestation to notify the Secretary if there has been a change in the accuracy of the information it submitted for initial listing, which includes contact information for the PSO (42 CFR 3.102(a)(1)(vi)). Independent Data Safety Monitoring, Inc. did not exercise its opportunity to be heard in writing to respond to the deficiency specified in the notices, and has not provided any evidence of a good faith effort to correct the deficiency. As such, pursuant to 42 CFR 3.108(a)(4)(iii), the notice of proposed revocation became final as a matter of law and the basis for revocation.

Accordingly, AHRQ has revoked the listing of Independent Data Safety Monitoring, Inc., PSO number P0114, effective at 12:00 Midnight ET (2400) on January 15, 2013.

More information on PSOs can be obtained through AHRQ's PSO Web site at *http://www.pso.AHRQ.gov/index.html.*

Dated: February 11, 2013.

Carolyn M. Clancy,

Director.

[FR Doc. 2013–03909 Filed 2–20–13; 8:45 am]

BILLING CODE 4160–90–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

National Institute for Occupational Safety and Health Personal Protective Technology for Pesticide Handlers: Stakeholder Meeting

AGENCY: The National Institute for Occupational Safety and Health (NIOSH) of the Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice of public meeting.

SUMMARY: The National Institute for Occupational Safety and Health (NIOSH) of the Centers for Disease Control and Prevention (CDC) announces the following public meeting: ''Pesticide Handler Personal Protective Technology Stakeholder Meeting''.

Stakeholder Meeting Time and Date:

4 p.m.–6 p.m. EDT, March 25, 2013 [Optional]

8 a.m.–6 p.m. EDT, March 26, 2013.

Place: NIOSH, Patriots Plaza 1, 395 E. Street, SW., Room 9000, Washington, DC 20201. This meeting will also be accessible remotely through Live Meeting with advanced registration.

Purpose of the Meeting: This meeting is being held to motivate and educate pesticide handlers and pesticide workers to use best pesticide personal protective equipment practices. This stakeholder meeting allows NIOSH to facilitate focused communication and exchange ideas and solutions between key stakeholder groups. Stakeholder feedback is sought to provide input to future updates of the NIOSH Personal

Protective Technology program research agenda.

Day 1, March 25th, 2013 (4 a.m.–6 p.m.) is optional. This first day includes informal introductions and discussions of partnering opportunities and on-going collaborations.

Day 2, March 26, 2013 (8 a.m.–6 p.m.) will include formal sessions on potential health effects of pesticide exposure, work safety culture, and the use and limitations of storytelling to motivate safer and healthier work practices. Pesticide handlers and pesticide workers will share their personal stories. The afternoon session will include an update from the U.S. Environmental Protection Agency (EPA).

This meeting will also include interactive sessions involving an expert panel in which stakeholders will brainstorm ways to collaboratively promote wide-spread adoption of best practices.

Status: The meeting is open to the public, limited only by the space available. The meeting room accommodates 33 people. This meeting will also be available to participants at 100 locations via Live Meeting. Live Meeting participants can simultaneously listen, speak, and view presentations via a telephone call and Internet connection.

Registration will be accepted on a first come first served basis. Registration to participate in person is available on the NIOSH NPPTL Web site: *www.cdc.gov/niosh/npptl.*

Registration to participate via Live Meeting is available at: *https://www.livemeeting.com/lrs/1100003614/Registration.aspx?pageName=lx5wbfv03vl92kzj.* Preregistration is required for both remote and in-person attendees.

An email confirming registration will be sent from NIOSH and will include details needed to participate. A government-issued photo ID will be required to obtain entrance to Patriots Plaza for those who will attend in-person. Non-US citizens must be cleared in advance. This clearance takes a minimum of 30 days.

FOR FURTHER INFORMATION CONTACT: Dr. Kimberly Faulkner, Epidemiologist, NIOSH NPPTL at *NPPTLEventsPesticide@cdc.gov,* telephone (412) 386–6111, fax (412) 386–6617.

Dated: February 14, 2013.

John Howard,

Director, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention.

[FR Doc. 2013–03925 Filed 2–20–13; 8:45 am]

BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Tracking of the Early Head Start Research and Evaluation Project (EHSRET) Sample.

OMB No.: 0970–0388.

Description: The EHSREP is a longitudinal study originally designed to meet the 1994 requirement for a national evaluation of the Early Head Start program. Child and family assessments were conducted when children were 14 months old, 24 months old, 36 months old, in the spring prior to kindergarten entry, and again in the spring of the sixth year of formal schooling (5th grade for most children). Today, children of the EHSREP are approximately 14–17 years of age (depending on their age at the time of enrollment in the study).

The Administration for Children and Families (ACF) within the Department of Health and Human Services (HHS) is proposing to track the children/families who participated in the Early Head Start Research and Evaluation Project (EHSREP) until the children reach 18 years of age. The purpose of tracking these participants is to maintain up-to-date contact information for the children/families in the event that the ACF determines that a future follow-up to the EHSREP will take place.

Respondents: Participants in the Early Head Start Research and Evaluation Project.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours	Annual burden hours
Tracking Survey	2533	3	.25	1,899	633
Administrative Records Consent Form	1700	1	.08	136	45

Estimated Total Annual Burden Hours: 678.

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 370 L'Enfant Promenade SW., Washington, DC 20447, Attn: OPRE Reports Clearance Officer. All requests should be identified by the title of the information collection. Email address: *OPREinfocollection@acf.hhs.gov.*

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, Email: *OIRA SUBMISSION@OMB.E0P.GOV,* Attn: Desk Officer for the Administration for Children and Families.

Steven M. Hanmer,

OPRE Reports Clearance Officer.

[FR Doc. 2013–03868 Filed 2–20–13; 8:45 am]

BILLING CODE 4184–22–P