

(such as 18–35), or patients enrolled in limited scale quality improvement projects.

The estimated burden for completing the nomination form is 30 minutes.

Amount of the Prize:

Up to a total of 20 of highest scoring individual clinicians, clinical practices or health systems will be recognized as Million Hearts® Hypertension Control Champions and will receive a cash award of \$2,000. A maximum of \$40,000 will be awarded in this challenge.

Payment of the Prize:

Prizes awarded under this challenge will be paid by electronic funds transfer and may be subject to Federal income taxes. HHS will comply with the Internal Revenue Service withholding and reporting requirements, where applicable.

Basis Upon Which Winner Will Be Selected:

The nomination will be scored based on hypertension control rate (95% of score); and sustainable systems in the practice that support hypertension control (5% of score).

Nominees with the highest score will be required to participate in a two-phase process to verify their data. Nominees who are non-compliant or non-responsive with the data requests or timelines will be removed from further consideration. Phase 1 includes verification of the hypertension prevalence and blood pressure control rate data submitted and a background check. For nominees whose Phase 1 data is verified as accurate, phase 2 consists of a medical chart review.

National Quality Forum Measure #0018 for controlling high blood pressure will be used to evaluate the submitted data. Details about NQF 0018 can be found at <http://www.qualityforum.org/QPS/0018>.

A CDC-sponsored panel of three to five experts consisting of HHS/CDC staff will review the nominations that pass the verification to select Champions. Final selection will take into account all the information from the nomination form, the background check, and data verification. Geographic location and population treated may be used to break ties in the event of tie scores at any point in the selection process.

Some Champions will participate in a post-challenge telephone interview. The interview will include questions about the strategies employed by the individual or organization to achieve high rates of hypertension control, including barriers and facilitators for those strategies. The interview will focus on systems and processes and should not require preparation time by the Champion. The estimated time for the interview is two hours, which

includes time to review the interview protocol with the interviewer, respond to the interview questions, and review a summary data about the Champion's practices. The summary will be written as a success story and will be posted on the Million Hearts® Web site.

Additional Information:

Information received from nominees will be stored in a password protected file on a secure server. The challenge Web site may post the number of nominations received but will not include information about individual nominees. The database of information submitted by nominees will not be posted on the Web site. Information collected from nominees will include general details, such as the business name, address, and contact information of the nominee. This type of information is generally publically available. The nomination will collect and store only aggregate clinical data through the nomination process; no individual identifiable patient data will be collected or stored. Confidential or propriety data, clearly marked as such, will be secured to the full extent allowable by law.

Information for selected Champions, such as the provider, practice, or health system's name, location, hypertension control rate, and clinic practices that support hypertension control will be shared through press releases, the challenge Web site, and Million Hearts® and HHS/CDC resources.

Summary data on the types of systems and processes that all nominees use to control hypertension may be shared in documents or other communication products that describe generally used practices for successful hypertension control. HHS/CDC will use the summary data only as described.

Compliance With Rules and Contacting Contest Winners:

Finalists and Champions must comply with all terms and conditions of these official rules, and winning is contingent upon fulfilling all requirements herein. The initial finalists will be notified by email, telephone, or mail after the date of the judging.

Privacy:

Personal information provided by entrants on the nomination form through the challenge Web site will be used to contact selected finalists. Information is not collected for commercial marketing. Winners are permitted to cite that they won this challenge.

The names, cities, and states of selected Champions will be made available in promotional materials and at recognition events.

General Conditions:

The HHS/CDC reserves the right to cancel, suspend, and/or modify the challenge, or any part of it, for any reason, at HHS/CDC's sole discretion.

Award Approving Official: Thomas R. Frieden, MD, MPH, Director, Centers for Disease Control and Prevention, and Administrator, Agency for Toxic Substances and Disease Registry

Authority: 15 U.S.C. 3719.

Dated: August 15, 2014.

Ron A. Otten,

Acting Deputy Associate Director for Science, Centers for Disease Control and Prevention.

[FR Doc. 2014–19754 Filed 8–19–14; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[CMS–7033–N]

Health Insurance Marketplace, Medicare, Medicaid, and Children's Health Insurance Programs; Meeting of the Advisory Panel on Outreach and Education (APOE), September 12, 2014

AGENCY: Centers for Medicare & Medicaid Services (CMS), HHS.

ACTION: Notice of meeting.

SUMMARY: This notice announces a meeting of the Advisory Panel on Outreach and Education (APOE) (the Panel) in accordance with the Federal Advisory Committee Act. The Panel advises and makes recommendations to the Secretary of the U.S. Department of Health and Human Services and the Administrator of the Centers for Medicare & Medicaid Services on opportunities to enhance the effectiveness of consumer education strategies concerning the Health Insurance Marketplace, Medicare, Medicaid, and the Children's Health Insurance Program (CHIP). This meeting is open to the public.

DATES:

Meeting Date: Friday, September 12, 2014, 8:30 a.m. to 4:00 p.m. eastern daylight time (e.d.t.).

Deadline for Meeting Registration, Presentations and Comments: Friday, August 29, 2014, 5:00 p.m., e.d.t.

Deadline for Requesting Special Accommodations: Friday, August 29, 2014, 5:00 p.m., e.d.t.

ADDRESSES:

Meeting Location: U.S. Department of Health & Human Services, Hubert H. Humphrey Building, 200 Independence Avenue SW., Room 738 G, Conference Room, Washington, DC 20201.

Presentations and Written Comments: Kirsten Knutson, Acting Designated Federal Official (DFO), Division of Forum and Conference Development, Office of Communications, Centers for Medicare & Medicaid Services, 7500 Security Boulevard, Mailstop S1-13-05, Baltimore, MD 21244-1850 or contact Ms. Knutson via email at Kirsten.Knutson@cms.hhs.gov.

Registration: The meeting is open to the public, but attendance is limited to the space available. Persons wishing to attend this meeting must register at the Web site <http://events.SignUp4.com/SEP2014APOEMTG> or by contacting the DFO at the address listed in the **ADDRESSES** section of this notice or by telephone at number listed in the **FOR FURTHER INFORMATION CONTACT** section of this notice, by the date listed in the **DATES** section of this notice. Individuals requiring sign language interpretation or other special accommodations should contact the DFO at the address listed in the **ADDRESSES** section of this notice by the date listed in the **DATES** section of this notice. In accordance with the U.S. Department of Health & Human Services standards, and an effort for the public to engage virtually in the open meetings, this APOE meeting will be available to view via live web streaming by visiting the link www.cms.gov/live during the designated time of the meeting.

FOR FURTHER INFORMATION CONTACT: Kirsten Knutson, (410) 786-5886. Additional information about the APOE is available on the Internet at: <http://www.cms.gov/Regulations-and-Guidance/Guidance/FACA/APOE.html>.

Press inquiries are handled through the CMS Press Office at (202) 690-6145.

SUPPLEMENTARY INFORMATION: In accordance with section 10(a) of the Federal Advisory Committee Act (FACA), this notice announces a meeting of the Advisory Panel on Outreach and Education (APOE) (the Panel). Section 9(a)(2) of the Federal Advisory Committee Act authorizes the Secretary of the U.S. Department of Health and Human Services (the Secretary) to establish an advisory panel if the Secretary determines that the panel is “in the public interest in connection with the performance of duties imposed . . . by law.” Such duties are imposed by section 1804 of the Social Security Act (the Act), requiring the Secretary to provide informational materials to Medicare beneficiaries about the Medicare program, and section 1851(d) of the Act, requiring the Secretary to provide for “activities . . . to broadly disseminate information to [M]edicare beneficiaries . . . on the coverage options provided

under [Medicare Advantage] in order to promote an active, informed selection among such options.”

The Panel is also authorized by section 1114(f) of the Act (42 U.S.C. 1314(f)) and section 222 of the Public Health Service Act (42 U.S.C. 217a). The Secretary signed the charter establishing this Panel on January 21, 1999 (64 FR 7899, February 17, 1999) and approved the renewal of the charter on December 18, 2012 (78 FR 32661, May, 31, 2013).

Pursuant to the amended charter, the Panel advises and makes recommendations to the Secretary of Health and Human Services and the Administrator of the Centers for Medicare & Medicaid Services (CMS) concerning optimal strategies for the following:

- Developing and implementing education and outreach programs for individuals enrolled in, or eligible for, the Health Insurance Marketplace, Medicare, Medicaid and the Children’s Health Insurance Program (CHIP).
- Enhancing the federal government’s effectiveness in informing Health Insurance Marketplace, Medicare, Medicaid, and CHIP consumers, providers, and stakeholders pursuant to education and outreach programs of issues regarding these and other health coverage programs; including the appropriate use of public-private partnerships to leverage the resources of the private sector in educating beneficiaries, providers, and stakeholders.
- Expanding outreach to vulnerable and underserved communities, including racial and ethnic minorities, in the context of the Health Insurance Marketplace, Medicare, Medicaid, and CHIP education programs.
- Assembling and sharing an information base of “best practices” for helping consumers evaluate health plan options.
- Building and leveraging existing community infrastructures for information, counseling, and assistance.
- Drawing the program link between outreach and education, promoting consumer understanding of health care coverage choices and facilitating consumer selection/enrollment, which in turn support the overarching goal of improved access to quality care, including prevention services, envisioned under health care reform.

The current members of the Panel are: Samantha Artiga, Principal Policy Analyst, Kaiser Family Foundation; Joseph Baker, President, Medicare Rights Center; Kellan Baker, Associate Director, Center for American Progress; Philip Bergquist, Manager, Health Center Operations, CHIPRA Outreach &

Enrollment Project and Director, Michigan Primary Care Association; Marjorie Cadogan, Executive Deputy Commissioner, Department of Social Services; Jonathan Dauphine, Senior Vice President, AARP; Barbara Ferrer, Executive Director, Boston Public Health Commission; Shelby Gonzales, Senior Health Outreach Associate, Center on Budget & Policy Priorities; Jan Henning, Benefits Counseling & Special Projects Coordinator, North Central Texas Council of Governments’ Area Agency on Aging; Louise Knight, Director, The Sidney Kimmel Comprehensive Cancer Center at Johns Hopkins; Miriam Mobley-Smith, Dean, Chicago State University, College of Pharmacy; Ana Natale-Pereira, Associate Professor of Medicine, Rutgers-New Jersey Medical School; Roanne Osborne-Gaskin, M.D., Associate Medical Director, Neighborhood Health Plan of Rhode Island; Megan Padden, Vice President, Sentara Health Plans; Jeanne Ryer, Director, New Hampshire Citizens Health Initiative, University of New Hampshire; Carla Smith, Executive Vice President, Healthcare Information and Management Systems Society (HIMSS); Winston Wong, Medical Director, Community Benefit Director, Kaiser Permanente and Darlene Yee-Melichar, Professor & Coordinator, San Francisco State University.

The agenda for the September 12, 2014 meeting will include the following:

- Welcome and listening session with CMS leadership
- Recap of the previous (May 22, 2014) meeting
- Affordable Care Act initiatives
- An opportunity for public comment
- Meeting summary, review of recommendations, and next steps

Individuals or organizations that wish to make a 5-minute oral presentation on an agenda topic should submit a written copy of the oral presentation to the DFO at the address listed in the **ADDRESSES** section of this notice by the date listed in the **DATES** section of this notice. The number of oral presentations may be limited by the time available. Individuals not wishing to make an oral presentation may submit written comments to the DFO at the address listed in the **ADDRESSES** section of this notice by the date listed in the **DATES** section of this notice.

Authority: Sec. 222 of the Public Health Service Act (42 U.S.C. 217a) and sec. 10(a) of Pub. L. 92-463 (5 U.S.C. App. 2, sec. 10(a) and 41 CFR 102-3).

(Catalog of Federal Domestic Assistance Program No. 93.733, Medicare—Hospital

Insurance Program; and Program No. 93.774, Medicare—Supplementary Medical Insurance Program)

Dated: August 12, 2014.

Marilyn Tavenner,

Administrator, Centers for Medicare & Medicaid Services.

[FR Doc. 2014–19698 Filed 8–19–14; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Title: Maternal and Infant Home Visiting Program Evaluation (MIHOPE)

Check-In project—Update contact information, consent forms, child and family outcomes survey.

OMB No.: 0970–0402.

Description: The Administration for Children and Families (ACF), in partnership with the Health Resources and Services Administration (HRSA), both of the U.S. Department of Health and Human Services (HHS), is proposing a data collection activity as part of the Maternal and Infant Home Visiting Program Evaluation (MIHOPE) Check-In project. The purpose of the MIHOPE Check-In project is to maintain up-to-date contact information for families that participated in MIHOPE (the national evaluation of the Maternal, Infant, and Early Childhood Home Visiting program), so it is possible to conduct future follow-up studies and

assess the potential long-term impact of the program. In addition to contact information, the MIHOPE Check-In project will also maintain up-to-date consent forms for the collection of administrative data and administer a brief survey on child and family outcomes.

Respondents: Adult participants in MIHOPE and adult primary caregivers of children who participated in MIHOPE.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Annual number of respondents	Number of responses per respondent	Average burden hours per response	Total annual burden hours
Child and Family Outcome Assessment and Updating Contact Information	4,300	1,433	3	.50	2,150
Updating Administrative Data Consent Forms	4,300	1,433	3	.25	1,075

Estimated total annual burden hours: 3225.

In compliance with the requirements of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded 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. Email address: OPREinfocollection@acf.hhs.gov. All requests should be identified by the title of the information collection.

The Department specifically requests comments on (a) whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or

other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Karl Koerper,

Reports Clearance Officer.

[FR Doc. 2014–19717 Filed 8–19–14; 8:45 am]

BILLING CODE 4184–22–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

Agency Information Collection Activities; Proposed Collection; Comment Request; Extension of the Help America Vote Act (HAVA), (HAVA Narrative Annual Report)

AGENCY: Administration on Intellectual & Developmental Disabilities, Administration for Community Living, HHS.

ACTION: Notice.

SUMMARY: The Administration for Community Living (ACL) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of

information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the information collection requirements relating to the Help America Vote Act (HAVA), Public Law 107–252, Title II, Subtitle D, Part 2, Sections 261 to 265 (HAVA Narrative Annual Report).

DATES: Submit written or electronic comments on the collection of information by September 19, 2014.

ADDRESSES: Submit written comments on the collection of information by fax 202.395.5806 or by email to OIRA_submission@omb.eop.gov, Attn: OMB Desk Officer for ACL, Office of Information and Regulatory Affairs, OMB.

FOR FURTHER INFORMATION CONTACT: Clare Barnett, Program Specialist, Administration for Community Living, Washington, DC 20201.

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. “Collection of information” is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency request or requirements that members of the public submit reports, keep records, or