(d) A requirement that payments are made to the recipient of the grant, contract, or cooperative agreement only when agreed upon outcomes are achieved, excluding payments made to a third party conducting the evaluation.

See 42 U.S.C. 711(k)(4).

The forthcoming SIR will provide further instructions to recipients in proposing a PFO initiative and submitting the required information to HRSA. Recipients are not required to propose or implement a PFO initiative, but if they wish to do so, they must submit a PFO SIR Response describing how their PFO initiative will meet all of the applicable statutory requirements. HRSA will use the information collected through the PFO SIR Response to ensure that MIECHV recipients' proposals to use grant funds for PFO initiatives meet statutory requirements and to provide technical assistance to recipients. The implementation of a PFO initiative is not intended to disrupt current services or negatively impact communities that have benefited from home visiting programs and must not result in a reduction of funding for home visiting services.

Likely Respondents: MIECHV Program recipients that are states, territories, and, where applicable, nonprofit organizations providing home visiting services within states.

Burden Statement: Burden in this context means the time expended by persons to generate, maintain, retain, disclose, or provide the information requested. This includes the time needed to review instructions and supporting materials; to collect and analyze data and information to develop the PFO SIR Response; engage with stakeholders and coordinate with state level partners; and to draft and submit the PFO SIR Response. The table below summarizes the total annual burden hours estimated for this SIR.

Total Estimated Annualized Burden Hours:

Instrument	Number of respondents	Number of responses per respondent	Total responses	Average burden hours per response	Total burden hours
MIECHV PAY FOR OUTCOMES SIR	15	1	15	92	1,380
Total	15		15		1,380

HRSA specifically requests comments on (1) the necessity and utility of the proposed information collection for the proper performance of the agency's functions, (2) the accuracy of the estimated burden, (3) ways to enhance the quality, utility, and clarity of the information to be collected, and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

Maria G. Button,

Director, Executive Secretariat.

[FR Doc. 2021–09910 Filed 5–10–21; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Allergy and Infectious Diseases; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of the following meeting.

The meeting 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: National Institute of Allergy and Infectious Diseases Special Emphasis Panel; NIAID Investigator Initiated Program Project Applications (P01 Clinical Trial Not Allowed).

Date: May 17, 2021.

Time: 10:00 a.m. to 1:30 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institute of Allergy and Infectious Diseases, National Institutes of Health, 5601 Fisher Lane, Room 3G45, Rockville, MD 20892 (Virtual Meeting).

Contact Person: Vanitha Sundaresa Raman, Ph.D., Scientific Review Officer, Scientific Review Program, Division of Extramural Activities, National Institute of Allergy and Infectious Diseases, National Institutes of Health, 5601 Fishers Lane, Room 3G45, Rockville, MD 20852, 301–761–7949, *vanitha.raman@nih.gov.*

This notice is being published less than 15 days prior to the meeting due to the timing limitations imposed by the review and funding cycle.

(Catalogue of Federal Domestic Assistance Program Nos. 93.855, Allergy, Immunology, and Transplantation Research; 93.856, Microbiology and Infectious Diseases Research, National Institutes of Health, HHS)

Dated: May 5, 2021.

Tyeshia M. Roberson,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2021–09913 Filed 5–10–21; 8:45 am]

BILLING CODE 4140–01–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–20–131: Mammalian Models for Translational Research.

Date: June 7, 2021.

Time: 9:00 a.m. to 7:00 p.m.

Agenda: To review and evaluate grant applications.

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

Contact Person: Jeffrey Smiley, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 6194, MSC 7804, Bethesda, MD 20892, (301) 272–4596, *smileyja@csr.nih.gov.*

Name of Committee: Vascular and Hematology Integrated Review Group; Basic