

go/connectpro overview. FDA has verified the Web site addresses in this document, as of the date this document publishes in the **Federal Register**, but Web sites are subject to change over time.

Transcripts: Please be advised that as soon as a transcript of the public workshop 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/MedicalDevices/NewsEvents/WorkshopsConferences/default.htm>.

IV. References

The following references are on display in the Dockets Management Staff (see **ADDRESSES**) and are available for viewing by interested persons between 9 a.m. and 4 p.m., Monday through Friday; they are also available electronically at <https://www.regulations.gov>. FDA has verified the Web site addresses, as of the date this document publishes in the **Federal Register**, but Web sites are subject to change over time.

1. National Cancer Institute, "Cancer Stat Facts: Cervix Uteri Cancer," (<http://seer.cancer.gov/statfacts/html/cervix.html>).
2. Horner, M.J., S.F. Altekruse, Z. Zou, L. Wideroff, et al. "U.S. Geographic Distribution of Pre-Vaccine Era Cervical Cancer Screening, Incidence, Stage, and Mortality." *Cancer Epidemiology, Biomarkers & Prevention*. 2011 Jan.; 20(4):591–9. doi: 10.1158/1055-9965.EPI-10-1183.
3. Freeman, H.W.B. "Excess Cervical Cancer Mortality: A Marker for Low Access to Health Care in Poor Communities." Rockville (MD): National Cancer Institute, Center to Reduce Cancer Health Disparities; 2005.

Dated: September 1, 2017.

Anna K. Abram,

Deputy Commissioner for Policy, Planning, Legislation, and Analysis.

[FR Doc. 2017–19029 Filed 9–7–17; 8:45 am]

BILLING CODE 4164-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Heart, Lung, and Blood Institute; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of a meeting of the NHLBI Special Emphasis Panel.

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 contract proposals and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the contract proposals, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Heart, Lung, and Blood Institute Special Emphasis Panel; Resources for Clinical Trials.

Date: September 20, 2017.

Time: 11:00 a.m. to 1:00 p.m.

Agenda: To review and evaluate contract proposals.

Place: National Institutes of Health, 6701 Rockledge Drive, Room 7200, Bethesda, MD 20892 (Telephone Conference Call).

Contact Person: Michael P. Reilly, Ph.D., Scientific Review Officer, Office of Scientific Review/DERA, National Heart, Lung, and Blood Institute, 6701 Rockledge Drive, Room 7200, Bethesda, MD 20892, 301–827–7975, reillymp@nhlbi.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. 93.233, National Center for Sleep Disorders Research; 93.837, Heart and Vascular Diseases Research; 93.838, Lung Diseases Research; 93.839, Blood Diseases and Resources Research, National Institutes of Health, HHS)

Dated: September 1, 2017.

Michelle Trout,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2017–19027 Filed 9–7–17; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Cancer Institute; 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: National Cancer Institute Special Emphasis Panel; Pancreatic Cancer Detection Consortium (U01).

Date: November 3, 2017.

Time: 11:00 a.m. to 2:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Cancer Institute, Shady Grove, 9609 Medical Center Drive, Room 7W240, Rockville, MD 20850, (Telephone Conference Call).

Contact Person: Hasan Siddiqui, Ph.D., Scientific Review Officer, Special Review Branch, Division of Extramural Activities, National Cancer Institute, NIH, 9609 Medical Center Drive, Room 7W240, Bethesda, MD 20892–9750, 240–276–5122, hasan.siddiqui@nih.gov.

Name of Committee: National Cancer Institute Special Emphasis Panel; Cooperative Agreement To Develop Targeted Agents Used With Systemic Agents Plus Radiotherapy.

Date: November 17, 2017.

Time: 10:00 a.m. to 2:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Cancer Institute, Shady Grove, 9609 Medical Center Drive, Room 7W640, Rockville, MD 20850, (Telephone Conference Call).

Contact Person: Saejeong J. Kim, Ph.D., Scientific Review Officer, Resources and Training Review Branch, Division of Extramural Activities, National Cancer Institute, NIH, 9609 Medical Center Drive, Room 7W640, Bethesda, MD 20892–9750, 240–276–7684, saejeong.kim@nih.gov.

(Catalogue of Federal Domestic Assistance Program Nos. 93.392, Cancer Construction; 93.393, Cancer Cause and Prevention Research; 93.394, Cancer Detection and Diagnosis Research; 93.395, Cancer Treatment Research; 93.396, Cancer Biology Research; 93.397, Cancer Centers Support; 93.398, Cancer Research Manpower; 93.399, Cancer Control, National Institutes of Health, HHS)

Dated: September 1, 2017.

Melanie J. Pantoja,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2017–19026 Filed 9–7–17; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; 30-Day Comment Request; Application To Participate in the National Institutes of Health Technical Assistance Programs: Commercialization Accelerator Program (CAP)

AGENCY: National Institutes of Health, HHS.

ACTION: Notice.

SUMMARY: In compliance with the Paperwork Reduction Act of 1995, the National Institutes of Health (NIH) has submitted to the Office of Management and Budget (OMB) a request for review and approval of the information collection listed below. This proposed information collection was previously published in the **Federal Register** on June 15, 2017, page 27516 (82 FR 27516) and allowed 60 days for public comment. No public comments were received. The purpose of this notice is to allow an additional 30 days for public comment.

DATES: Comments regarding this information collection are best assured of having their full effect if received within 30-days of the date of this publication.

ADDRESSES: Written comments and/or suggestions regarding the item(s) contained in this notice, especially regarding the estimated public burden and associated response time, should be directed to the: Office of Management and Budget, Office of Regulatory Affairs, *OIRA_submission@omb.eop.gov* or by fax to 202-395-6974, Attention: Desk Officer for NIH.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of

the data collection plans and instruments, contact: J. P. Kim, NIH SBIR/STTR Program Manager & NIH Extramural Data Sharing Policy Officer, Small Business Innovation Research (SBIR) and Small Business Technology Transfer (STTR) Program Office, Office of Extramural Programs (OEP)/Office of Extramural Research (OER), Office of the Director (OD)/National Institutes of Health (NIH), 6705 Rockledge Drive, Suite 350; Bethesda, Maryland 20892-7963 or call non-toll-free number (301) 435-0189 or Email your request, including your address to: *jpkim@nih.gov*. Formal requests for additional plans and instruments must be requested in writing.

SUPPLEMENTARY INFORMATION: The Commercialization Accelerator Program (CAP), National Institutes of Health, may not conduct or sponsor, and the respondent is not required to respond to, an information collection that has been extended, revised, or implemented on or after October 1, 1995, unless it displays a currently valid OMB control number.

In compliance with Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Institutes of Health (NIH) has submitted to the Office of Management and Budget (OMB) a request for review and

approval of the information collection listed below.

Proposed Collection: Application to Participate in the National Institutes of Health Technical Assistance Programs: Commercialization Accelerator Program (CAP)—0925—Existing Without OMB Approval, Office of Extramural Programs (OEP)/Office of Extramural Research (OER), Office of the Director (OD)/National Institutes of Health.

Need and Use of Information Collection: The purpose of this application is to collect information to be used internally by the NIH SBIR/STTR staff to identify and select small businesses that would most benefit if selected as participants in the NIH Commercialization Accelerator Program (CAP). The data will not be used to formulate or change policies. Rather, it will be used to enable NIH SBIR/STTR staff to be responsive to its constituents by offering commercialization training to meet the goals of the Phase II small business NIH awardees. The form will be online for any potential CAP applicant companies and completed electronically.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 150.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Number of respondents	Number of responses per respondent	Average time per response	Total annual burden hour
SBIR Phase II Awardees	100	1	90	150
Total	100	100		150

Dated: September 1, 2017.

Lawrence A. Tabak,

Principal Deputy Director, National Institutes of Health.

[FR Doc. 2017-19078 Filed 9-7-17; 8:45 am]

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DEPARTMENT OF HOMELAND SECURITY

Coast Guard

[Docket No. USCG-2016-1084]

Navigation and Vessel Inspection Circular (NVIC) 05-17; Guidelines for Addressing Cyber Risks at Maritime Transportation Security Act (MTSA) Regulated Facilities

AGENCY: Coast Guard, DHS.

ACTION: Notice of availability and request for comments; extension of comment period.

SUMMARY: The Coast Guard is extending the comment period for its notice of availability and request for comments, published on July 12, 2017. The notice announced the availability of the draft Navigation and Vessel Inspection Circular (NVIC) 05-17 entitled Guidelines for Addressing Cyber Risks at Maritime Transportation Security Act (MTSA) Regulated Facilities, and requested public comments on the draft. The comment period was set to close on September 11, 2017. The Coast Guard has received requests to extend the comment period by 30 days due to the conditions caused by hurricane Harvey, which prevent some members of the public from submitting comments by the original deadline.

DATES: Comments and related material must reach the Coast Guard on or before October 11, 2017.

ADDRESSES: You may submit comments identified by docket number USCG-2016-1084 using the Federal eRulemaking Portal at <http://www.regulations.gov>. See the "Public Participation and Request for Comments" portion of the **SUPPLEMENTARY INFORMATION** section of the July 12, 2017 notice for further instructions on submitting comments.

FOR FURTHER INFORMATION CONTACT: If you have questions on this notice, call or email, Jason Warren, Coast Guard; telephone 202-372-1106, email Jason.S.Warren@uscg.mil or LCDR Josephine Long, Coast Guard; telephone 202-372-1109, email Josephine.A.Long@uscg.mil.