552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended for the review and discussion of grant applications. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the contact person listed below in advance of the meeting.

Name of Committee: National Advisory Child Health and Human Development Council.

Date: September 14, 2017.

Open: September 14, 2017.

Time: 8:00 a.m. to 12:00 p.m.

Agenda: The agenda will include opening remarks, administrative matters, Director's Report, Division of Extramural Research Report and, other business of the Council.

Place: National Institutes of Health, Building 31, C-Wing, Conference Room 6, 9000 Rockville Pike, Bethesda, MD 20892.

Closed: September 14, 2017.

Time: 1:00 p.m. to Adjournment.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Building 31, C-Wing, Conference Room 6, 9000 Rockville Pike, Bethesda, MD 20892.

Contact Person: Della Hann, Ph.D., Director, Division of Extramural Research, Eunice Kenney Shriver, National Institute of Child Health and Human Development, NIH, 6710 Rockledge Blvd., MSC 7002, Bethesda, MD 20892, 301–496–8535.

Any interested person may file written comments with the committee by forwarding the statement to the contact person listed on this notice. The statement should include the name, address, telephone number, and when applicable, the business or professional affiliation of the interested person.

In the interest of security, NIH has instituted stringent procedures for entrance onto the NIH campus. All visitor vehicles, including taxis, hotel, and airport shuttles, will be inspected before being allowed on campus. Visitors will be asked to show one form of identification (for example, a government-issued photo ID, driver's license, or passport) and to state the purpose of their visit.

In order to facilitate public attendance at the open session of Council in the main meeting room, Conference Room 6, please contact Ms. Lisa Kaeser, Program and Public Liaison Office, NICHD, at 301–496–0536 to make your reservation, additional seating will be available in the meeting overflow rooms, Conference Rooms 7 and 8. Individuals will also be able to view the meeting via NIH Videocast. Please go to the following link for Videocast access instructions at: *http://www.nichd.nih.gov/about/advisory/nachhd/Pages/virtual-meeting.aspx.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.864, Population Research; 93.865, Research for Mothers and Children; 93.929, Center for Medical Rehabilitation Research; 93.209, Contraception and Infertility Loan Repayment program, National Institutes of Health, HHS).

Dated: August 1, 2017.

Michelle Trout,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2017–16520 Filed 8–4–17; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Biomedical Imaging and Bioengineering; 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 Biomedical Imaging and Bioengineering Special Emphasis Panel; JHU Translational Immuno-Engineering BTRC (2018/01).

Date: September 26, 2017.

Time: 09:00 a.m. to 8:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Two Democracy Plaza, Suite 920, 6707 Democracy Boulevard, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: John K. Hayes, Ph.D., Scientific Review Officer, 6707 Democracy Blvd., Suite 959, Democracy Two, Bethesda, MD 20892, (301) 451–3398, *hayesj@mail.nih.gov.*

Dated: August 1, 2017.

David Clary,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2017–16533 Filed 8–4–17; 8:45 am]

BILLING CODE 4140–01–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; Pilot Clinical Trials Targeting HIV–1 Reservoirs in Children (U01).

Date: August 23, 2017.

Time: 11:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 5601 Fishers Lane, Rockville, MD 20892 (Telephone Conference Call).

Contact Person: J. Bruce Sundstrom, Ph.D., Scientific Review Officer, Scientific Review Program, Division of Extramural Activities, Rm. 3G11A, National Institutes of Health/NIAID, 5601 Fishers Lane, MSC 9823, Bethesda, MD 20892–9823, 240–669–5045, *sundstromj@niaid.nih.gov.*

(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: August 1, 2017.

Sylvia L. Neal,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2017–16523 Filed 8–4–17; 8:45 am]

BILLING CODE 4140–01–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; Asthma and Allergic Diseases Cooperative Research Centers.

Date: September 7–14, 2017.

Time: 12:00 p.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: The William F. Bolger Center, 9600 Newbridge Drive, Potomac, MD 20854.

Contact Person: Paul A. Amstad, Ph.D., Scientific Review Officer, Scientific Review Program, NIAID/NIH/DHHS, Division of Extramural Activities, Room 3G41, 5601 Fishers Lane, Bethesda, MD 20892–7616, 240–669–5067, *pamstad@niaid.nih.gov.*

(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: August 1, 2017.

Sylvia L. Neal,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2017–16522 Filed 8–4–17; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Prospective Grant of Exclusive Patent License: The Development of a Bispecific, Biparatopic Antibody-Drug Conjugate to GPC3 for the Treatment of Human Liver Cancers

AGENCY: National Institutes of Health, HHS.

ACTION: Notice.

SUMMARY: The National Cancer Institute, National Institutes of Health, Department of Health and Human Services, is contemplating the grant of an Exclusive Patent License to Salubris Biotherapeutics, Inc. (Salubris), located in Gaithersburg, Maryland, to practice the inventions embodied in the patent applications listed in the **SUPPLEMENTARY INFORMATION** section of this notice.

DATES: Only written comments and/or applications for a license which are received by the NCI Technology Transfer Center on or before August 22, 2017 will be considered.

ADDRESSES: Requests for copies of the patent applications, inquiries, and comments relating to the contemplated Exclusive Patent License should be directed to: David A. Lambertson, Ph.D., Senior Licensing and Patenting Manager, NCI Technology Transfer Center, 9609 Medical Center Drive, RM 1E530 MSC 9702, Bethesda, MD 20892–9702 (for business mail), Rockville, MD 20850–9702; Telephone: (240) 276–6467; Email: *david.lambertson@nih.gov.*

SUPPLEMENTARY INFORMATION: The following represents the intellectual property to be licensed under the prospective agreement: (A) U.S. Provisional Patent Application 61/654,232 entitled ''High-affinity Monoclonal Antibodies To Glypican-3 And Use Thereof'' [HHS Ref. E–136–2012/0–US–01], PCT Patent Application PCT/US2013/043633 entitled ''High-affinity Monoclonal Antibodies To Glypican-3 And Use Thereof'' [HHS Ref. E–136–2012/0–PCT–02], Chinese Patent Application 201380039993.7 entitled ''High-affinity Monoclonal Antibodies To Glypican-3 And Use Thereof'' [HHS Ref. E–136–2012/0–CN–03], Japanese Patent Application 2015–515243 entitled ''High-affinity Monoclonal Antibodies To Glypican-3 And Use Thereof'' [HHS Ref. E–136–2012/0–JP–04], South Korean Patent Application 10–2014–7037046 entitled ''High-affinity Monoclonal Antibodies To Glypican-3 And Use Thereof'' [HHS Ref. E–136–2012/0–KR–05], Singapore Patent Application 11201407972R entitled ''High-affinity Monoclonal Antibodies To Glypican-3 And Use Thereof'' [HHS Ref. E–136–2012/0–SG–06], and United States Patent 9,409,994 entitled ''High-affinity Monoclonal Antibodies To Glypican-3 And Use Thereof'' [HHS Ref. E–136–2012/0–US–07], and all continuing U.S. and foreign patents/patent applications for the technology family; and (B) U.S. Provisional Patent Application 61/477,020 entitled ''Human Monoclonal Antibody Specific for Glypican-3 And Use Thereof'' [HHS Ref. E–130–2011/0–US–01], PCT Patent Application PCT/US2012/034186 entitled ''Human Monoclonal Antibodies Specific for Glypican-3 And Use Thereof'' [HHS Ref. E–130–2011/0–PCT–02], Chinese Patent 201280029201.3 entitled ''Human Monoclonal Antibodies Specific for Glypican-3 And Use Thereof'' [HHS Ref. E–130–2011/0–CN–03], European Patent 2699603 entitled ''Human Monoclonal Antibodies Specific for Glypican-3 And Use Thereof'' [HHS Ref. E–130–2011/0–EP–04], and validated in France [HHS Ref. E–130–2011/0–FR–09], Germany [HHS Ref. E–130–2011/0–DE–08] and the United Kingdom [HHS Ref. E–130–2011/0–GB–10] and lodged in Hong Kong [HHS Ref. E–130–2011/0–HK–11], United States Patent 9,206,257 entitled ''Human Monoclonal Antibodies Specific for Glypican-3 And Use Thereof'' [HHS Ref. E–130–2011/0–US–05], United States Patent 9,394,364, entitled ''Human Monoclonal Antibodies Specific for Glypican-3 And Use Thereof'' [HHS Ref. E–130–2011/0–US–06], European Patent Application 15188264.4 entitled ''Human Monoclonal Antibodies Specific for Glypican-3 And Use Thereof'' [HHS Ref. E–130–2011/0–EP–07], United States Patent Application 15/090,873 entitled ''Human Monoclonal Antibodies Specific for Glypican-3 And Use Thereof'' [HHS Ref. E–130–2011/0–US–12], Chinese Patent Application 201610290837.3 entitled ''Human Monoclonal Antibodies Specific for Glypican-3 And Use Thereof'' [HHS Ref. E–130–2011/0–CN–13], European Patent Application 16166924.7 entitled ''Human Monoclonal Antibodies Specific for Glypican-3 And Use Thereof'' [HHS Ref. E–130–2011/0–EP–14], and all continuing U.S. and foreign patents/patent applications for the technology family, to Salubris. The patent rights in these inventions have been assigned to and/or exclusively licensed to the Government of the United States of America.

With respect to persons who have an obligation to assign their right, title and interest to the Government of the United States of America, the patent rights in these inventions have been assigned to the Government of the United States of America.

The prospective Exclusive Patent License territory may be worldwide for the following field of use:

The development and commercialization of a bispecific, biparatopic antibody-drug conjugate (ADC) having:

(1) The CDR sequences of both the hYP7 and HN3 anti-GPC3 monoclonal antibodies; and

(2) a microtubule inhibitor payload including, but not limited to, auristatin and mertansine;

for the treatment of human liver cancer. The licensed field of use excludes any (a) non-specified immunoconjugates, including, but not limited to, chimeric antigen receptors (CARs) and variants thereof, immunotoxins, ADCs with payloads that are not microtubule inhibitors, and monospecific versions of the aforementioned immunoconjugates, and (b) unconjugated antibodies.

The present inventions to be licensed concern monoclonal antibodies that are specific for the cell surface domain of GPC3: HN3 and hYP7. These antibodies can potentially be used for the treatment of GPC3-expressing cancers such as HCC. In the subject situation, the antibodies can be used in conjunction to target a toxic payload specifically to GPC3-expressing cells, leading to the selective destruction of the cancerous cells.

This notice is made in accordance with 35 U.S.C. 209 and 37 CFR part 404.