

clearing/approving forms, processing forms, and acknowledging data entered.

The total estimated annualized burden hours for all respondents are 1,280. There are no costs to respondents

other than their time. CDC will seek a three-year approval from OMB.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per CIO	Average burden per response (in hours)	Total burden (in hours)
Initiator/C//O	CDC 0.4601	64	5	1	320
Initiator/C//O	CDC 0.410A	64	5	1	320
Initiator/C//O	CDC 0.410B	64	5	1	320
Initiator/C//O	Section C of the CDC 0.1475	64	5	1	320
Totals	1,280

Leroy A. Richardson,

Chief, Information Collection Review Office, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2016-19461 Filed 8-15-16; 8:45 am]

BILLING CODE 4163-18-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 (5 U.S.C. App.), 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 Clinical Trial Implementation Cooperative Agreement (U01).

Date: September 22, 2016.

Time: 10:00 a.m. to 12:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Room 3F100, 5601 Fishers Lane, Rockville, MD 20892 (Telephone Conference Call).

Contact Person: Lynn Rust, Ph.D., Scientific Review Officer, Scientific Review Program, Division of Extramural Activities, Room 3G42A, National Institutes of Health/NIAID, 5601 Fishers Lane, MSC 9823,

Bethesda, MD 20892-9823, (240) 669-5069, lrust@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 10, 2016.

Natasha M. Copeland,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2016-19417 Filed 8-15-16; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Prospective Grant of Start-Up Exclusive Evaluation Option License Agreement: Small Molecule Therapeutic Compounds Encompassed Within the Licensed Patent Rights for the Treatment of Thioesterase Deficiency Disorder

AGENCY: National Institutes of Health, HHS.

ACTION: Notice.

SUMMARY: This notice, in accordance with 35 U.S.C. 209 and 37 CFR part 404, that the National Institutes of Health, Department of Health and Human Services, is contemplating the grant of a Start-Up Exclusive Evaluation Option License Agreement to practice the inventions embodied in the following Patent Applications to Circumvent Pharmaceuticals Inc. (“Circumvent”) located in Pasadena, California, USA:

Intellectual Property

United States Provisional Patent Application No. 61/473,692, filed April 8, 2011, titled “Small molecule therapeutic compounds targeting thioesterase deficiency disorders and methods of using the same” [HHS

Reference No. E-157-2011/0-US-01], status: Expired;

International Patent Application No. PCT/US2012/32772 filed April 9, 2012 titled “Small molecule therapeutic compounds targeting thioesterase deficiency disorders and methods of using the same” [HHS Reference No. E-157-2011/0-PCT-02], status: Converted;

European Patent Application No. 12716889.6, filed November 7, 2013, titled “Small molecule therapeutic compounds targeting thioesterase deficiency disorders and methods of using the same” [HHS Reference No. E-157-2011/0-EP-03], status: Pending; and

United States Patent Application No. 14/110,393, filed October 7, 2013, titled “Small molecule therapeutic compounds targeting thioesterase deficiency disorders and methods of using the same” [HHS Reference No. E-157-2011/0-US-04], status: Pending.

The patent rights in these inventions have been assigned and/or exclusively licensed to the government of the United States of America.

The territory of the prospective Start-Up Exclusive Evaluation Option License Agreement may be worldwide and the field of use may be limited to: “Small molecule therapeutic compounds encompassed within the Licensed Patent Rights for the treatment of thioesterase deficiency disorders”

Upon the expiration or termination of the Start-up Exclusive Evaluation Option License Agreement, Circumvent will have the exclusive right to execute a Start-Up Exclusive Patent License Agreement which will supersede and replace the Start-up Exclusive Evaluation Option License Agreement, with no greater field of use and territory than granted in the Start-up Exclusive Evaluation Option License Agreement.

DATES: Only written comments and/or applications for a license which are received by the NIH Office of