that the offense warranted the imposition of a 5-year period of debarment.

The proposal informed Ms. Godding of the proposed debarment and offered her an opportunity to request a hearing, providing her 30 days from the date of receipt of the letter in which to file the request, and advised her that failure to request a hearing constituted a waiver of the opportunity for a hearing and of any contentions concerning this action. Ms. Godding received the proposal and notice of opportunity for a hearing on August 30, 2022. Ms. Godding failed to request a hearing within the timeframe prescribed by regulation and has, therefore, waived her opportunity for a hearing and waived any contentions concerning her debarment (21 CFR part 12).

II. Findings and Order

Therefore, the Assistant Commissioner, Office of Human and Animal Food Operations, under section 306(b)(3)(C) of the FD&C Act, under authority delegated to the Assistant Commissioner, finds that Ms. Linda Godding has been convicted of a felony under Federal law for conduct relating to the importation into the United States of any drug or controlled substance. FDA finds that the offense should be accorded a debarment period of 5 years as provided by section 306(c)(2)(A)(iii) of the FD&C Act.

As a result of the foregoing finding, Ms. Godding is debarred for a period of 5 years from importing or offering for import any drug into the United States, effective (see **DATES**). Pursuant to section 301(cc) of the FD&C Act (21 U.S.C. 331(cc)), the importing or offering for import into the United States of any drug by, with the assistance of, or at the direction of Ms. Godding is a prohibited act.

Any application by Ms. Godding for termination of debarment under section 306(d)(1) of the FD&C Act should be identified with Docket No. FDA–2022–N–1398 and sent to the Dockets Management Staff (see **ADDRESSES**). The public availability of information in these submissions is governed by 21 CFR 10.20(j).

Publicly available submissions will be placed in the docket and will be viewable at *https://www.regulations.gov* or at the Dockets Management Staff (see **ADDRESSES**) between 9 a.m. and 4 p.m., Monday through Friday, 240–402–7500.

Dated: January 12, 2023.

Lauren K. Roth,

Associate Commissioner for Policy.

[FR Doc. 2023–00997 Filed 1–19–23; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Mental Health; Amended Notice of Meeting

Notice is hereby given of a change in the meeting of the National Institute of Mental Health Special Emphasis Panel, January 24, 2023, 12:00 p.m. to January 24, 2023, 4:00 p.m., National Institutes of Health, Neuroscience Center, 6001 Executive Boulevard, Rockville, MD, 20852, which was published in the **Federal Register** on December 30, 2022, FR Doc 2022–28446, 87 FR 80554.

This notice is being amended to change the meeting date from January 24, 2023, to February 2, 2023. Meeting location and time remain the same. The meeting is closed to the public.

Dated: January 13, 2023.

Melanie J. Pantoja,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2023–01018 Filed 1–19–23; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Prospective Grant of an Exclusive Patent License: Small Cell Lung Cancer Subtyping Using Plasma Cell-Free Nucleosomes

AGENCY: National Institutes of Health, HHS.

ACTION: Notice.

SUMMARY: The National Cancer Institute (NCI), an institute of the National Institutes of Health, Department of Health and Human Services, is contemplating the grant of an exclusive, sublicensable patent license to Yissum Research and Development (''Yissum''), the technology transfer company of the Hebrew University of Jerusalem, a non-profit research institution located in Jerusalem, Israel for NCI's rights to 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 National Cancer Institute's Technology Transfer Center on or before February 6, 2023 will be considered.

ADDRESSES: Requests for copies of the patent application, inquiries, and comments relating to the contemplated exclusive patent license should be directed to: Michaela McCrary, Ph.D., Licensing and Patenting Manager, NCI Technology Transfer Center, at: Email: *michaela.mccrary@nih.gov.*

SUPPLEMENTARY INFORMATION:

Intellectual Property

The following and all continuing U.S. and foreign patents/patent applications thereof are the intellectual properties to be licensed under the prospective agreement to Yissum: United States Provisional Patent Application No. 63/342,763, filed May 17, 2022 and entitled ''SMALL CELL LUNG CANCER SUBTYPING USING PLASMA CELL–FREE NUCELOSOMES'' [HHS Reference No. E–172–2022–0–US–01].

The patent rights in these inventions have been assigned to the Government of the United States of America and Yissum. The prospective license will be for the purpose of consolidating the patent rights to Yissum, the co-owners of said rights, for commercial development and marketing. Consolidation of these co-owned rights is intended to expedite development of the invention, consistent with the goals of the Bayh-Dole Act codified as 35 U.S.C. 200–212.

The prospective patent license territory will be worldwide, exclusive, and may be limited to those fields of use commensurate in scope with the patent rights. It will be sublicensable, and any sublicenses granted by Yissum will be subject to the provisions of 37 CFR part 401 and 404.

This technology discloses a non-invasive method to molecularly subtype SCLC from plasma samples using chromatin immunoprecipitation of cell-free nucleosomes carrying active chromatin modification followed by sequencing (cfChIP–seq).

This notice is made in accordance with 35 U.S.C. 209 and 37 CFR part 404. The prospective exclusive license will include terms for the sharing of royalty income with NCI from commercial sublicenses of the patent rights. The prospective exclusive license may be granted unless within fifteen (15) days from the date of this published notice, the National Cancer Institute receives written evidence and argument that establishes that the grant of the license would not be consistent with the requirements of 35 U.S.C. 209 and 37 CFR part 404.

Complete applications for a license that are timely filed in response to this notice will be treated as objections to the grant of the contemplated exclusive patent license. In response to this Notice, the public may file comments or objections. Comments and objections, other than those in the form of a license