Technology Transfer Center, 9609 Medical Center Drive, RM 3W–204, MSC 9702, Bethesda, MD 20892–9702 (for business mail), Rockville, MD 20850–9702; Telephone: (240)–276–5530; Email: *suna.gulay@nih.gov.*

SUPPLEMENTARY INFORMATION:

Intellectual Property

1. United States Provisional Patent Application No. 62/327,529, filed April 26, 2016 and entitled ''Anti-KK–LC–1 T Cell Receptors'' [HHS Reference No. E–153–2016–0–US–01];
2. PCT Patent Application No. PCT/US2017/027865, filed April 17, 2017 and entitled ''Anti-KK–LC–1 T Cell Receptors'' [HHS Reference No. E–153–2016–0–PCT–02];
3. Australian Patent No. 2017258745, issued July 14, 2022 and entitled ''Anti-KK–LC–1 T Cell Receptors'' [HHS Reference No. E–153–2016–0–AU–03];
4. Canadian Patent Application No. 3021898, filed April 17, 2017 and entitled ''Anti-KK–LC–1 T Cell Receptors'' [HHS Reference No. E–153–2016–0–CA–04];
5. European Patent No. 3448882, issued November 24, 2021 and entitled ''Anti-KK–LC–1 T Cell Receptors'' [HHS Reference No. E–153–2016–0–EP–05];
a. Validated in the following jurisdictions: CH, DE, BE, DK, ES, FI, FR, GB, IE, IT, NL, NO and SE.
6. U.S. Patent No. 11,352,410, issued June 7, 2022 and entitled ''Anti-KK–LC–1 T Cell Receptors'' [HHS Reference No. E–153–2016–0–US–06].

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

The prospective exclusive license territory may be worldwide and the field of use may be limited to the following:

''1. Development, manufacture, and commercialization of autologous T cell therapy products, including T cells with stem-like properties, engineered via retrovirus-mediated gene transfer to express T cell receptors reactive to Kita-Kyushu Lung Cancer Antigen 1 (KK–LC–1), as claimed in the Licensed Patent Rights; such products to be developed for treatment of patients carrying HLA–A*01:01 histocompatibility haplotype, and diagnosed with a cancer expressing KK–LC–1 protein (''KK–LC–1 Targeting TCR–T Products'').

2. Development, manufacture, and commercialization of a combination therapy for the treatment of KK–LC–1 expressing human cancers, independent of their HLA phenotype, wherein the treatment comprises:

a. Modification of the patient's tumor using Licensee's proprietary technology to express the HLA–A*01:01 restriction element, and

b. Treatment with the KK–LC–1 Targeting TCR–T Products.

For the avoidance of doubt, specifically excluded from these Fields of Use are Natural Killer cell therapy products engineered via viral vectors (including lentivirus or retrovirus) to express the TCR(s) claimed in the Licensed Patent Rights.''

This technology discloses isolated T cell receptors (TCR) reactive to the Kita-Kyushu lung cancer antigen 1 (KK–LC–1) within the context of human leukocyte antigen (HLA) A*01:01. KK–LC–1 is expressed by various epithelial cancers including carcinomas of the bladder, cervix, stomach, breast, lung, and pancreas. Due to its minimal expression in normal tissues, this antigen may be targeted on KK–LC–1-expressing tumors with minimal normal tissue toxicity.

This notice is made in accordance with 35 U.S.C. 209 and 37 CFR part 404. The prospective exclusive license will be royalty bearing, and 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.

In response to this Notice, the public may file comments or objections. Comments and objections, other than those in the form of a license application, will not be treated confidentially, and may be made publicly available.

License applications submitted in response to this Notice will be presumed to contain business confidential information and any release of information in these license applications will be made only as required and upon a request under the Freedom of Information Act, 5 U.S.C. 552.

Dated: October 1, 2024.

Richard U. Rodriguez,

Associate Director, Technology Transfer Center, National Cancer Institute.

[FR Doc. 2024–23029 Filed 10–4–24; 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 1009 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: Biological Chemistry and Macromolecular Biophysics Integrated Review Group; Drug Discovery and Molecular Pharmacology B Study Section.

Date: October 31-November 1, 2024.

Time: 8:00 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: Razvan Cornea, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 904L, Bethesda, MD 20892, (301) 480–1955, *cornearl@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel; Imaging and Bioengineering Technology for Visual Systems (IBV).

Date: November 4–5, 2024.

Time: 8:00 a.m. to 7:00 p.m.

Agenda: To review and evaluate grant applications.

Place: The Watergate, 2650 Virginia Avenue NW, Washington, DC 20037.

Contact Person: Susan Gillmor, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (240) 762–3076, *susan.gillmor@nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel; Fellowships: Learning, Memory, Language, Communication and Related Neuroscience.

Date: November 4–5, 2024.

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

Agenda: To review and evaluate grant applications.

Place: Canopy by Hilton, 940 Rose Avenue, North Bethesda, MD 20852.

Contact Person: Alexei Kondratyev, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5200, MSC 7846, Bethesda, MD 20892, 301–435–1785 *kondratyevad@csr.nih.gov.*

Name of Committee: Applied Therapeutics for Cancer Integrated Review Group; Drug Discovery and Molecular Pharmacology C Study Section.

Date: November 4–5, 2024.

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

Agenda: To review and evaluate grant applications.

Place: The Westin Georgetown, 2350 M Street NW, Washington, DC 20037.

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: Center for Scientific Review Special Emphasis Panel; Member Conflict: Radiation Therapeutics and Biology.

Date: November 4, 2024.

Time: 9:00 a.m. to 8:00 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: Lambratu Rahman Sesay, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 6214, MSC 7804, Bethesda, MD 20892, 301–905–8294, *rahman-sesay@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel; Member conflict: Topics on Biobehavioral Processes

Date: November 4–5, 2024.

Time: 10:00 a.m. to 8:00 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: Natalie S Dailey, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (301) 827–4451, *daileyns@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel; PAR–24–129: Specific Pathogen Free Macaque Colonies.

Date: November 4, 2024.

Time: 1:00 p.m. to 3:00 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: Latha Malaiyandi, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 812Q, Bethesda, MD 20892, (301) 435–1999, *malaiyandilm@csr.nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.306, Comparative Medicine; 93.333, Clinical Research, 93.306, 93.333, 93.337, 93.393–93.396, 93.837–93.844, 93.846–93.878, 93.892, 93.893, National Institutes of Health, HHS)

Dated: October 1, 2024.

Victoria E. Townsend,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2024–23037 Filed 10–4–24; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HOMELAND SECURITY

Federal Emergency Management Agency

[Docket ID FEMA–2024–0002; Internal Agency Docket No. FEMA–B–2463]

Proposed Flood Hazard Determinations

AGENCY: Federal Emergency Management Agency, Department of Homeland Security.

ACTION: Notice.

SUMMARY: Comments are requested on proposed flood hazard determinations, which may include additions or modifications of any Base Flood Elevation (BFE), base flood depth, Special Flood Hazard Area (SFHA) boundary or zone designation, or regulatory floodway on the Flood Insurance Rate Maps (FIRMs), and where applicable, in the supporting Flood Insurance Study (FIS) reports for the communities listed in the table below. The purpose of this notice is to seek general information and comment regarding the preliminary FIRM, and where applicable, the FIS report that the Federal Emergency Management Agency (FEMA) has provided to the affected communities. The FIRM and FIS report are the basis of the floodplain management measures that the community is required either to adopt or to show evidence of having in effect in order to qualify or remain qualified for participation in the National Flood Insurance Program (NFIP).

DATES: Comments are to be submitted on or before January 6, 2025.

ADDRESSES: The Preliminary FIRM, and where applicable, the FIS report for each community are available for inspection at both the online location *https://hazards.fema.gov/femaportal/prelimdownload* and the respective Community Map Repository address listed in the tables below. Additionally, the current effective FIRM and FIS report for each community are accessible online through the FEMA Map Service Center at *https://msc.fema.gov* for comparison.

You may submit comments, identified by Docket No. FEMA–B–2463, to Rick Sacbibit, Chief, Engineering Services Branch, Federal Insurance and Mitigation Administration, FEMA, 400 C Street SW, Washington, DC 20472, (202) 646–7659, or (email) *patrick.sacbibit@fema.dhs.gov.*

FOR FURTHER INFORMATION CONTACT: Rick Sacbibit, Chief, Engineering Services Branch, Federal Insurance and Mitigation Administration, FEMA, 400 C Street SW, Washington, DC 20472, (202) 646–7659, or (email) *patrick.sacbibit@fema.dhs.gov;* or visit the FEMA Mapping and Insurance eXchange (FMIX) online at *https://www.floodmaps.fema.gov/fhm/fmx_main.html.*

SUPPLEMENTARY INFORMATION: FEMA proposes to make flood hazard determinations for each community listed below, in accordance with section 110 of the Flood Disaster Protection Act of 1973, 42 U.S.C. 4104, and 44 CFR 67.4(a).

These proposed flood hazard determinations, together with the floodplain management criteria required by 44 CFR 60.3, are the minimum that are required. They should not be construed to mean that the community must change any existing ordinances that are more stringent in their floodplain management requirements. The community may at any time enact stricter requirements of its own or pursuant to policies established by other Federal, State, or regional entities. These flood hazard determinations are used to meet the floodplain management requirements of the NFIP.

The communities affected by the flood hazard determinations are provided in the tables below. Any request for reconsideration of the revised flood hazard information shown on the Preliminary FIRM and FIS report that satisfies the data requirements outlined in 44 CFR 67.6(b) is considered an appeal. Comments unrelated to the flood hazard determinations also will be considered before the FIRM and FIS report become effective.

Use of a Scientific Resolution Panel (SRP) is available to communities in support of the appeal resolution process. SRPs are independent panels of experts in hydrology, hydraulics, and other pertinent sciences established to review conflicting scientific and technical data and provide recommendations for resolution. Use of the SRP only may be exercised after FEMA and local communities have been engaged in a collaborative consultation process for at least 60 days without a mutually acceptable resolution of an appeal. Additional information regarding the SRP process can be found