## Congress of the United States

Washington, DC 20515

June 14, 2023

The Honorable Arati Prabhakar, Ph.D. Director Office of Science and Technology Policy Executive Office of the President Eisenhower Executive Office Building 1650 Pennsylvania Avenue Washington, D.C. 20504

Dear Director Prabhakar:

As you know, in December 2022, the President signed the PREVENT Pandemics Act ("PREVENT") into law as Title II of Division FF of the Consolidated Appropriations Act, 2023.<sup>1</sup> PREVENT intended to address lessons learned from the COVID-19 pandemic response, including improving and enhancing domestic biosafety and biosecurity. As part of this effort, the law directs the Office of Science and Technology Policy (OSTP) to undertake two specific activities: (1) develop a strategy for the management, maintenance, and oversight of federal high-containment laboratories (section 2312); and (2) review and update policies for the review and oversight of research involving enhanced pathogens of pandemic potential (section 2315). Your office is responsible for fulfilling both of these requirements by December 29, 2023.

Section 2312 was intended as an initial step toward addressing a longstanding Government Accountability Office (GAO) recommendation that a single federal entity be charged with leading the evaluation and coordination of high-containment laboratory capacity in the U.S.<sup>2</sup> Among other requirements, the law states that your strategy should:

- 1. Describe federal roles and responsibilities related to high-containment laboratories;
- 2. Assess federal capacity needs and summarize existing capacity that is either federally owned or was established through federal funding at non-federal facilities;
- 3. Summarize existing plans to maintain this capacity and describe how agencies determine whether it is necessary to maintain or expand capacity; and
- 4. Describe how federal departments and agencies will coordinate in overseeing this capacity to improve scientific collaboration and reduce any unnecessary duplication of effort.

Section 2315 was intended to capture OSTP's January 2017 *Recommended Policy Guidance for Departmental Development of Review Mechanisms for Potential Pandemic Pathogen Care and Oversight (P3CO)*. As written, the provision could also include updates to other policy frameworks, such as policies for the oversight of dual-use research of concern (DURC) at federal

<sup>&</sup>lt;sup>1</sup> Pub. L. No. 117-328, 136 Stat. 4459 (2022).

<sup>&</sup>lt;sup>2</sup> https://www.gao.gov/assets/gao-09-574.pdf.

and nonfederal institutions. Notably, both GAO and the National Science Advisory Board for Biosecurity, which Congress also codified through PREVENT, have recently issued recommendations to improve both policies with the goal of ensuring a more cohesive framework for oversight of this type of research.<sup>3</sup> Under the law, the updated policy should:

- 1. Include a clear scope to support the consistent identification of research proposals subject to such policy by relevant Federal departments and agencies;
- 2. Provide a framework for such reviews that accounts for safety, security, and ethical considerations related to the creation, transfer, or use of enhanced pathogens of pandemic potential;
- 3. Include measures to enhance the transparency and public availability of information related to such research actives in a manner that does not compromise national security, the safety and security of such research activities, or any identifiable, sensitive information of relevant individuals; and
- 4. Ensure consistent procedures across relevant Federal departments and agencies related to the identification and referral of research proposals.

Timely implementation of sections 2312 and 2315 is critical to enhancing our biosafety and biosecurity. Accordingly, we request a briefing for our staffs no later than Friday, June 30, 2023, from members of your staff. The goal of this briefing will be to learn about your progress to date in implementing these two provisions and your plan to fulfill these requirements by the statutory deadline of December 29, 2023. Thank you for your attention to this request.

Sincerely,

Bill Cassidy, M.D.

Bill Cassidy, M.D. Ranking Member Senate HELP Committee

Mitt Romney Member Senate HELP Committee

Cothe Mr L hodger

Cathy McMorris Rodgers Chair House Energy and Commerce Committee

Richard Hudson Member House Energy and Commerce Committee

<sup>&</sup>lt;sup>3</sup> https://www.gao.gov/assets/gao-23-105455.pdf; https://osp.od.nih.gov/wp-content/uploads/2023/03/NSABB-Final-Report-Proposed-Biosecurity-Oversight-Framework-for-the-Future-of-Science.pdf.