

118TH CONGRESS  
1ST SESSION

# S. 36

To review domestic biopharmaceutical manufacturing capabilities in order to improve public health and medical preparedness and response capabilities and domestic biopharmaceutical manufacturing capabilities.

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## IN THE SENATE OF THE UNITED STATES

JANUARY 24 (legislative day, JANUARY 3), 2023

Mr. RUBIO (for himself and Mr. COONS) introduced the following bill; which was read twice and referred to the Committee on Health, Education, Labor, and Pensions

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## A BILL

To review domestic biopharmaceutical manufacturing capabilities in order to improve public health and medical preparedness and response capabilities and domestic biopharmaceutical manufacturing capabilities.

1 *Be it enacted by the Senate and House of Representa-*  
2 *tives of the United States of America in Congress assembled,*

3 **SECTION 1. SHORT TITLE.**

4 This Act may be cited as the “Agility in Manufac-  
5 turing Preparedness Act of 2023”.

1 **SEC. 2. REVIEW OF DOMESTIC BIOPHARMACEUTICAL MAN-**  
2 **UFACTURING CAPABILITIES.**

3 (a) IN GENERAL.—The Secretary of Health and  
4 Human Services (referred to in this section as the “Sec-  
5 retary”), in cooperation with the Director of the Bio-  
6 medical Advanced Research and Development Authority,  
7 shall seek to enter into an agreement with the National  
8 Institute for Innovation in Manufacturing Biopharma-  
9 ceuticals to perform the services described in subsection  
10 (b).

11 (b) REVIEW AND RECOMMENDATIONS.—Under an  
12 agreement described in subsection (a) between the Sec-  
13 retary, the Director of the Biomedical Advanced Research  
14 and Development Authority, and the National Institute for  
15 Innovation in Manufacturing Biopharmaceuticals, the Na-  
16 tional Institute for Innovation in Manufacturing Bio-  
17 pharmaceuticals shall—

18 (1) review current domestic biopharmaceutical  
19 manufacturing capacity at the Department of  
20 Health and Human Services and such department’s  
21 adaptability to various threats;

22 (2) draft recommendations for developing, dem-  
23 onstrating, deploying, and advancing new domestic  
24 biopharmaceutical manufacturing technologies that  
25 address gaps identified under paragraph (1) and  
26 align Federal technologies with technologies avail-

1       able to the private sector, including through the new  
2       BioMAP initiative of the Biomedical Advanced Re-  
3       search and Development Authority; and

4               (3) identify other opportunities and priorities to  
5       improve the United States public health and medical  
6       preparedness and response capabilities and domestic  
7       biopharmaceutical manufacturing capabilities.

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