

116TH CONGRESS  
2D SESSION

**S.** \_\_\_\_\_

To prohibit distribution abroad of COVID–19 vaccines developed with the support of Federal funding until the domestic need for the vaccine has been met.

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IN THE SENATE OF THE UNITED STATES

Mr. TILLIS introduced the following bill; which was read twice and referred to the Committee on \_\_\_\_\_

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**A BILL**

To prohibit distribution abroad of COVID–19 vaccines developed with the support of Federal funding until the domestic need for the vaccine has been met.

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 “America First Vaccine  
5 Act”.

6 **SEC. 2. COVID–19 VACCINE DISTRIBUTION.**

7 (a) CERTIFICATION REQUIREMENT.—

8 (1) IN GENERAL.—Notwithstanding any other  
9 provision of law, subject to paragraph (2), a manu-

1       facturer of a vaccine described in subsection (b) that  
2       is headquartered in the United States may not dis-  
3       tribute such vaccine outside of the United States un-  
4       less the Secretary of Health and Human Services  
5       (referred to in this section as the “Secretary”), in  
6       consultation with the Commissioner of Food and  
7       Drugs, certifies that domestic need for the vaccine  
8       already has been met.

9           (2) WAIVER.—The Secretary may waive the  
10       certification requirement under paragraph (1) with  
11       respect to a vaccine described in subsection (b) and  
12       permit the distribution of such vaccine outside of the  
13       United States, if the Secretary determines that such  
14       waiver and distribution is in the interest of the  
15       American public health.

16       (b) VACCINES DESCRIBED.—A vaccine described in  
17       this subsection is a COVID–19 vaccine that—

18           (1) is authorized under section 564 of the Fed-  
19       eral Food, Drug, and Cosmetic Act (21 U.S.C.  
20       360bbb–3) or licensed under section 351 of the Pub-  
21       lic Health Service Act (42 U.S.C. 262); and

22           (2) was developed with support of Federal fund-  
23       ing made available under the Coronavirus Prepared-  
24       ness and Response Supplemental Appropriations Act  
25       (Public Law 116–123), the Families First

1        Coronavirus Response Act (Public Law 116–127),  
2        the Coronavirus Aid, Relief, and Economic Security  
3        Act (Public Law 116–136), or the Paycheck Protec-  
4        tion Program and Health Care Enhancement Act  
5        (Public Law 116–139).

6        (c) VIOLATIONS.—In the case of a manufacturer who  
7        violates subsection (a), such manufacturer shall be re-  
8        quired to reimburse the Federal Government for all  
9        amounts received by the manufacturer in support of devel-  
10       opment of the vaccine, plus interest in an amount deter-  
11       mined by the Secretary of the Treasury.

12       (d) RULE OF CONSTRUCTION.—Nothing in this Act  
13       shall be construed to limit or otherwise restrict the Sec-  
14       retary’s authority to enter into bilateral or multilateral  
15       agreements with international manufacturers to procure  
16       vaccines described in subsection (b) for the people of the  
17       United States.