

**Suspend the Rules and Pass the Bill, H.R. 4764, with an Amendment**

**(The amendment strikes all after the enacting clause and inserts a new text)**

116<sup>TH</sup> CONGRESS  
1<sup>ST</sup> SESSION

# H. R. 4764

To reauthorize the Stem Cell Therapeutic and Research Act of 2005, and for other purposes.

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IN THE HOUSE OF REPRESENTATIVES

OCTOBER 18, 2019

Ms. MATSUI (for herself, Mr. BILIRAKIS, and Ms. PINGREE) introduced the following bill; which was referred to the Committee on Energy and Commerce

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## A BILL

To reauthorize the Stem Cell Therapeutic and Research Act of 2005, and for other purposes.

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 “Timely ReAuthoriza-  
5 tion of Necessary Stem-cell Programs Lends Access to  
6 Needed Therapies Act of 2020” or the “TRANSPLANT  
7 Act of 2020”.

1 **SEC. 2. REAUTHORIZATION OF THE C.W. BILL YOUNG CELL**  
2 **TRANSPLANTATION PROGRAM.**

3 (a) **ADVISORY COUNCIL MEETINGS.**—Subsection (a)  
4 of section 379 of the Public Health Service Act (42 U.S.C.  
5 274k) is amended by adding at the end the following new  
6 paragraph:

7 “(7) The Secretary shall convene the Advisory  
8 Council at least two times each calendar year.”.

9 (b) **INCREASING COLLECTION.**—

10 (1) **TECHNICAL CLARIFICATION.**—Effective as  
11 if included in the enactment of Public Law 114–104  
12 (the Stem Cell Therapeutic and Research Reauthor-  
13 ization Act of 2015), the amendment to section  
14 379(d)(2)(B) of the Public Health Service Act (42  
15 U.S.C. 274k(d)(2)(B)) in section 2(a)(2) of Public  
16 Law 114–104 is amended by inserting “goal of in-  
17 creasing collections of high quality” before “cord  
18 blood units,”.

19 (2) **ELIMINATING DEADWOOD.**—Subparagraph  
20 (B) of section 379(d)(2) of the Public Health Serv-  
21 ice Act (42 U.S.C. 274k(d)(2)) is amended by strik-  
22 ing the second and third sentences in such subpara-  
23 graph.

24 (c) **PERIODIC REVIEW OF STATE OF SCIENCE.**—Sec-  
25 tion 379 of the Public Health Service Act (42 U.S.C.

1 274k) is amended by adding at the end the following new  
2 subsection:

3 “(o) PERIODIC REVIEW OF STATE OF SCIENCE.—

4 “(1) REVIEW.—Not less than every two years,  
5 the Secretary, in consultation with the Director of  
6 the National Institutes of Health, the Commissioner  
7 of Food and Drugs, the Administrator of the Health  
8 Resources and Services Administration, the Advisory  
9 Council, and other stakeholders, where appropriate  
10 given relevant expertise, shall conduct a review of  
11 the state of the science of using adult stem cells and  
12 birthing tissues to develop new types of therapies for  
13 patients, for the purpose of considering the potential  
14 inclusion of such new types of therapies in the Pro-  
15 gram.

16 “(2) RECOMMENDATIONS.—Not later than  
17 June 30, 2024, the Secretary shall—

18 “(A) complete the second review required  
19 by paragraph (1); and

20 “(B) informed by such review, submit to  
21 the Committee on Health, Education, Labor,  
22 and Pensions of the Senate and the Committee  
23 on Energy and Commerce of the House of Rep-  
24 resentatives recommendations on the appro-

1           priateness of the inclusion of new types of  
2           therapies in the Program.”.

3           (d) **AUTHORIZATION OF APPROPRIATIONS.**—Section  
4 379B of the Public Health Service Act (42 U.S.C. 274m)  
5 is amended by striking “\$33,000,000 for fiscal year 2015  
6 and \$30,000,000 for each of fiscal years 2016 through  
7 2020” and inserting “\$30,000,000 for each of fiscal years  
8 2021 through 2025”.

9 **SEC. 3. CORD BLOOD INVENTORY.**

10          Subsection (g) of section 2 of the Stem Cell Thera-  
11 peutic and Research Act of 2005 (42 U.S.C. 274k note)  
12 is amended to read as follows:

13          “(g) **AUTHORIZATION OF APPROPRIATIONS.**—To  
14 carry out this section, there is authorized to be appro-  
15 priated \$23,000,000 for each of fiscal years 2021 through  
16 2025.”.

17 **SEC. 4. ADVANCING THE FIELD OF REGENERATIVE MEDI-**  
18 **CINE.**

19          Section 402 of the Public Health Service Act (42  
20 U.S.C. 282) is amended by adding at the end the fol-  
21 lowing:

22          “(o) **REGENERATIVE MEDICINE.**—The Director of  
23 NIH shall, as appropriate, continue to consult with the  
24 directors of relevant institutes and centers of the National  
25 Institutes of Health, other relevant experts from such in-

1 stitutes and centers, and relevant experts within the Food  
2 and Drug Administration, to further the field of regenera-  
3 tive medicine using adult stem cells, including autologous  
4 stem cells, therapeutic tissue engineering products, human  
5 cell and tissue products, human gene therapies, and ge-  
6 netically modified cells.”.

7 **SEC. 5. GAO REPORT ON REGENERATIVE MEDICINE WORK-**  
8 **FORCE.**

9 Not later than 2 years after the date of enactment  
10 of this Act, the Comptroller General of the United States  
11 shall submit to the Committee on Health, Education,  
12 Labor, and Pensions of the Senate and the Committee on  
13 Energy and Commerce of the House of Representatives  
14 a report that assesses the national blood stem cell work-  
15 force, including those related to the C.W. Bill Young Cell  
16 Transplantation Program established under section 379 of  
17 the Public Health Service Act (42 U.S.C. 274k). The re-  
18 port shall include—

- 19 (1) an overview of the current employment lev-  
20 els, in both commercial and academic settings, for—  
21 (A) positions necessary for the collection  
22 and transplantation of stem cell therapeutics,  
23 including bone marrow and cord blood;

1 (B) positions in the field of regenerative  
2 medicine using adult stem cells and related to  
3 product development; and

4 (C) Federal funding for extramural stem  
5 cell programs;

6 (2) an overview of the current employment lev-  
7 els in Federal stem cell programs, including the  
8 scope of, staffing models of, and vacancies within  
9 such programs;

10 (3) the identification of gaps, if any, in the pro-  
11 jected workforce capacity for—

12 (A) positions described in paragraph  
13 (1)(A); and

14 (B) the field of regenerative medicine using  
15 adult stem cells, including workforce gaps re-  
16 lated to the development of new cellular thera-  
17 pies using adult stem cells;

18 (4) an overview of the availability of training  
19 programs related to the development, refinement,  
20 and utilization of adult stem cells, including training  
21 on good manufacturing practices for such activities,  
22 and the performance of such programs; and

23 (5) recommendations, if any, for improving the  
24 workforce capacity related to—

1 (A) the positions described in paragraph  
2 (1)(A); or  
3 (B) the field of regenerative medicine using  
4 adult stem cells.