

119TH CONGRESS  
1ST SESSION

# H. R. 3480

To amend the Patient Protection and Affordable Care Act to include fertility treatment and care as an essential health benefit.

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## IN THE HOUSE OF REPRESENTATIVES

MAY 19, 2025

Ms. UNDERWOOD introduced the following bill; which was referred to the Committee on Energy and Commerce

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## A BILL

To amend the Patient Protection and Affordable Care Act to include fertility treatment and care as an essential health benefit.

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 “Health Coverage for  
5 Inclusive and Valued Families Act of 2025” or the  
6 “Health Coverage for IVF Act of 2025”.

1   **SEC. 2. INCLUDING FERTILITY TREATMENT AND CARE AS**  
2                   **AN ESSENTIAL HEALTH BENEFIT.**

3       (a) IN GENERAL.—Section 1302(b) of the Patient  
4 Protection and Affordable Care Act (42 U.S.C. 18022(b))  
5 is amended—

6                   (1) in paragraph (1)—  
7                      (A) in the matter preceding subparagraph  
8                      (A), by striking “paragraph (2)” and inserting  
9                      “paragraphs (2) and (6)”; and

10                  (B) by adding at the end the following new  
11                  subparagraph:

12                  “(K) Fertility treatment and care.”; and

13                  (2) by adding at the end the following new  
14                  paragraph:

15                  “(6) FERTILITY TREATMENT AND CARE DE-  
16                  FINED.—For purposes of paragraph (1)(K), the  
17                  term ‘fertility treatment and care’ means the fol-  
18                  lowing medically appropriate items and services fur-  
19                  nished to an individual:

20                  “(A) Preservation of human oocytes,  
21                  sperm, or embryos for later reproductive use.

22                  “(B) Artificial insemination, including  
23                  intravaginal insemination, intracervical insemi-  
24                  nation, and intrauterine insemination.

25                  “(C) Assisted reproductive technology, in-  
26                  cluding in vitro fertilization and other treat-

1           ments or procedures in which reproductive ge-  
2           netic material, such as oocytes, sperm, fertilized  
3           eggs, and embryos, are handled, when clinically  
4           appropriate, and including at least 3 complete  
5           oocyte retrievals and an unlimited number of  
6           embryo transfers from such retrievals (regard-  
7           less of whether such retrieval was performed on,  
8           before, or after the date of the enactment of  
9           this paragraph) in accordance with the guide-  
10          lines of the American Society for Reproductive  
11          Medicine and using single embryo transfer  
12          when recommended and medically appropriate.

13                 “(D) Genetic testing of embryos.

14                 “(E) Medications prescribed, as indicated  
15                 for fertility.

16                 “(F) Gamete donation.

17                 “(G) Such other information, referrals,  
18                 treatments, procedures, medications, laboratory  
19                 testing, technologies, and services relating to  
20                 fertility as the Secretary determines appro-  
21                 priate.”.

22                 (b) ADDITIONAL REQUIREMENTS.—Subpart II of  
23                 part A of title XXVII of the Public Health Service Act  
24                 (42 U.S.C. 300gg–11 et seq.) is amended by adding at  
25                 the end the following new section:

1     **“SEC. 2730. REQUIREMENTS RELATING TO FERTILITY**

2                         **TREATMENT AND CARE.**

3         “(a) IN GENERAL.—In the case of health insurance  
4 coverage offered in the individual or small group market  
5 that provides both medical and surgical benefits and bene-  
6 fits for fertility treatment and care (as defined in section  
7 1302(b) of the Patient Protection and Affordable Care  
8 Act), such coverage shall ensure that—

9                 “(1) the financial requirements applicable to  
10 such fertility treatment and care benefits are no  
11 more restrictive than the predominant financial re-  
12 quirements applied to substantially all medical and  
13 surgical benefits covered by the coverage, and there  
14 are no separate cost sharing requirements that are  
15 applicable only with respect to fertility treatment  
16 and care benefits; and

17                 “(2) the treatment limitations applicable to  
18 such fertility treatment and care benefits are no  
19 more restrictive than the predominant treatment  
20 limitations applied to substantially all medical and  
21 surgical benefits covered by the coverage and there  
22 are no separate treatment limitations that are appli-  
23 cable only with respect to fertility treatment and  
24 care benefits.

25         “(b) PROHIBITION ON DENIAL OF CARE.—A health  
26 insurance issuer offering health insurance coverage in the

1 individual or small group market may not deny benefits  
2 for fertility treatment and care for individual on the basis  
3 that such individual lacks a diagnosis of infertility.

4       “(c) UTILIZATION MANAGEMENT TOOLS.—

5           “(1) IN GENERAL.—A health insurance issuer  
6 offering health insurance coverage in the individual  
7 or small group market that imposes any utilization  
8 management tool with respect to fertility treatment  
9 and care shall, for each of the first 5 plan years be-  
10 ginning on or after the date that is 1 year after the  
11 date of the enactment of this Act (and, upon request  
12 of the Secretary or the Comptroller General of the  
13 United States, for any subsequent plan year), con-  
14 duct an analysis of the application of any such tool  
15 to such treatment and care and submit such analysis  
16 to the Secretary and to the Comptroller General of  
17 the United States. Such analysis shall contain the  
18 following information:

19           “(A) The specific coverage terms or other  
20 relevant terms regarding the application of such  
21 tools to such benefits and a description of all  
22 such benefits.

23           “(B) The factors used to determine when  
24 utilization management tools apply to such ben-  
25 efits.

1                 “(C) The evidentiary standards used in de-  
2                 signing the application of such tools with re-  
3                 spect to such benefits and any other source or  
4                 evidence used to determine the application of  
5                 such tools to such benefits.

6                 “(D) Information demonstrating how ap-  
7                 plication of such tools to such benefits are con-  
8                 sistent with clinical guidelines for fertility treat-  
9                 ment and care.

10                 “(E) Any findings by the issuer that such  
11                 coverage is not in compliance with this section.

12                 “(2) REPORT.—For plan years beginning on or  
13                 after the date that is 1 year after the date of the  
14                 enactment of this section, the Comptroller General  
15                 of the United States shall submit to Congress and  
16                 make publicly available a report that contains the  
17                 following:

18                 “(A) A summary of the analyses submitted  
19                 under paragraph (1) with respect to such plan  
20                 year.

21                 “(B) An identification of each health in-  
22                 surance issuer that failed to submit an analysis  
23                 under paragraph (1).

24                 “(C) With respect to each health insurance  
25                 issuer that did submit such an analysis, a speci-

1 fication as to whether such issuer submitted in-  
2 formation sufficient to determine whether such  
3 issuer was in compliance with such require-  
4 ments.

5 “(D) For each health insurance issuer that  
6 did submit information sufficient to determine  
7 such compliance, a finding of whether such  
8 issuer was in compliance with such require-  
9 ments.

10 “(d) DEFINITIONS.—The terms ‘financial require-  
11 ment’, ‘predominant’, and ‘treatment limitation’ have the  
12 meaning given such terms in section 2726(a)(3).”.

13 (c) EFFECTIVE DATE.—The amendments made by  
14 this section shall apply to plan years beginning on or after  
15 the date that is 1 year after the date of the enactment  
16 of this Act.

