

119TH CONGRESS  
1ST SESSION

# S. 2276

To address patent thickets.

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IN THE SENATE OF THE UNITED STATES

JULY 15, 2025

Mr. WELCH (for himself, Mr. HAWLEY, and Ms. KLOBUCHAR) introduced the following bill; which was read twice and referred to the Committee on the Judiciary

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## A BILL

To address patent thickets.

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 “Eliminating Thickets  
5 to Increase Competition Act” or the “ETHIC Act”.

6 **SEC. 2. ADDRESSING PATENT THICKETS.**

7 (a) LIMIT ON NUMBER OF PATENTS PER PATENT  
8 GROUP THAT MAY BE ASSERTED IN ACTION FOR IN-  
9 FRINGEMENT.—Section 271(e) of title 35, United States  
10 Code, is amended by adding at the end the following:

1       “(7)(A) A person who brings an action for infringe-  
2       ment of a patent under this section against a party de-  
3       scribed in subparagraph (B) may assert in the action not  
4       more than one patent per Patent Group.

5       “(B) A party described in this subparagraph is—

6               “(i) a person who—

7                       “(I) submits an application for approval of  
8                       a drug under subsection (b)(2) or (j) of section  
9                       505 of the Federal Food, Drug, and Cosmetic  
10                      Act (21 U.S.C. 355), or is a holder of such an  
11                      approved application; or

12                     “(II) submits an application for licensure  
13                     of a biological product under section 351(k) of  
14                     the Public Health Service Act (42 U.S.C.  
15                     262(k)), or is a holder of such a licensure; or

16               “(ii) a person making, using, selling, offering  
17       for sale, introducing or delivering into interstate  
18       commerce, or importing—

19                     “(I) a drug approved pursuant to an appli-  
20                     cation under subsection (b)(2) or (j) of section  
21                     505 of the Federal Food, Drug, and Cosmetic  
22                     Act (21 U.S.C. 355); or

23                     “(II) a biological product licensed under  
24                     section 351(k) of the Public Health Service Act  
25                     (42 U.S.C. 262(k)).

1 “(C) A person who brings an action described in sub-  
 2 paragraph (A) asserting a patent against a party may not  
 3 bring any additional actions described in that subpara-  
 4 graph asserting a patent in the same Patent Group  
 5 against that party.

6 “(D)(i) For purposes of this paragraph, the term  
 7 ‘Patent Group’ means 2 or more commonly owned patents  
 8 or applications that—

9 “(I) are identified on 1 or more disclaimers  
 10 under section 253 to obviate obviousness-type double  
 11 patenting of another commonly owned patent; or

12 “(II) are subject to 1 or more disclaimers under  
 13 section 253 to obviate obviousness-type double pat-  
 14 enting of another commonly owned patent.

15 “(ii) For purposes of clause (i)(I)—

16 “(I) each patent or application that identifies  
 17 the same patent or application on a disclaimer under  
 18 section 253 is part of the same Patent Group; and

19 “(II) each patent or application that is identi-  
 20 fied on a disclaimer under section 253 is part of the  
 21 same Patent Group as the patent or application sub-  
 22 ject to the disclaimer.”.

23 (b) APPLICABILITY.—The amendment made by sub-  
 24 section (a) shall apply with respect to an application sub-  
 25 mitted under subsection (b)(2) or (j) of section 505 of the

1 Federal Food, Drug, and Cosmetic Act (21 U.S.C. 355)  
2 or section 351(k) of the Public Health Service Act (42  
3 U.S.C. 262(k)) on or after the date of enactment of this  
4 Act.

