

115TH CONGRESS
2D SESSION

H. R. 6593

To direct the Comptroller General of the United States to conduct a study on the feasibility of establishing and maintaining an automated and searchable system that tracks prescription drug prices in the United States, and for other purposes.

IN THE HOUSE OF REPRESENTATIVES

JULY 26, 2018

Mr. DUNCAN of Tennessee introduced the following bill; which was referred to the Committee on Energy and Commerce

A BILL

To direct the Comptroller General of the United States to conduct a study on the feasibility of establishing and maintaining an automated and searchable system that tracks prescription drug prices in the United States, 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 “Medication Automated
5 Quotation System, or MEDAQ, Act of 2018”.

1 **SEC. 2. STUDY ON THE FEASIBILITY OF CREATING A SYS-**
2 **TEM THAT TRACKS PRESCRIPTION DRUG**
3 **COSTS.**

4 (a) STUDY.—The Comptroller General of the United
5 States, in consultation with the Commissioner of Food and
6 Drugs and the Administrator of CMS, shall conduct a
7 study on the feasibility of establishing a medication auto-
8 mated quotation system.

9 (b) ELEMENTS.—The feasibility study required under
10 subsection (a) shall include assessments of the following:

11 (1) The feasibility of—

12 (A) collecting or compiling data on the ac-
13 quisition price per unit that any purchaser (in-
14 cluding but not limited to a distributor, a re-
15 tailer, a third-party payor, or ultimate con-
16 sumer) pays for a prescription drug and the ef-
17 fect PBMs and other involved entities have on
18 such prices; and

19 (B) aggregating such data in a manner
20 that does not reveal personally identifiable in-
21 formation.

22 (2) The cost of procuring and maintaining a
23 medication automated quotation system, or
24 MEDAQ, based on the above data.

25 (3) The need for new or additional laws, regula-
26 tions, or policies to collect the data necessary for a

1 medication automated quotation system to track and
2 generate prescription drug pricing data in the
3 United States.

4 (4) The possibility of integrating or using the
5 FDA’s NDC system and CMS NADAC when cre-
6 ating the MEDAQ.

7 (5) Integration of CMS NADAC when creating
8 the MEDAQ and, notwithstanding any other provi-
9 sion of this Act, shall not consider pharmacy report-
10 ing, collecting, or compiling requirements beyond or
11 in addition to the requirements of CMS NADAC.

12 (6) Whether the system described under sub-
13 section (a) is likely to be effective in reducing overall
14 prescription drug prices by increasing competition
15 and transparency in the drug industry.

16 (c) REPORT.—Not later than 270 days after the date
17 of the enactment of this Act, the Comptroller General shall
18 submit to Congress a report on the results of the study
19 conducted under subsection (a).

20 (d) MEDICATION AUTOMATED QUOTATION SYSTEM
21 DEFINED.—The term “medication automated quotation
22 system” or “MEDAQ” means an automated and search-
23 able system that—

24 (1) with respect to the United States (and with
25 respect to each region thereof), tracks and generates

1 data on the acquisition price per unit of any pre-
2 scription drug that a purchaser pays for such unit
3 of such drug;

4 (2) produces reports in real time on average
5 prices per unit and other data; and

6 (3) is accessible to purchasers, who have paid
7 a user fee, with limitations to ensure appropriate
8 privacy but also transparency in price negotiations.

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