

AMENDMENT TO
RULES COMMITTEE PRINT 116-51
OFFERED BY MR. KRISHNAMOORTHY OF ILLINOIS

At the end of title I, insert the following new section:

1 **SEC. 111. REPORTING OF CONSUMER COMPLAINTS BY**
2 **ENDS MANUFACTURERS AND IMPORTERS.**

3 Section 904 of the Federal Food, Drug, and Cosmetic
4 Act (21 U.S.C. 287d) is amended by adding at the end
5 the following new subsection:

6 “(f) **REPORTING OF CONSUMER COMPLAINTS BY**
7 **ENDS MANUFACTURERS AND IMPORTERS.—**

8 “(1) **IN GENERAL.—**Not later than the date
9 that is 1 year after the date of enactment of the
10 Protecting American Lungs and Reversing the
11 Youth Tobacco Epidemic Act of 2020, and annually
12 thereafter, each manufacturer or importer of an
13 electronic nicotine delivery system shall submit a re-
14 port to the Secretary describing each consumer com-
15 plaint received by the manufacturer or importer dur-
16 ing the reporting period with respect to such system.

1 “(2) COMPLAINTS INCLUDED.—The consumer
2 complaints to be reported under paragraph (1) shall
3 include complaints regarding—

4 “(A) adverse health effects associated with
5 use of the electronic nicotine delivery system;

6 “(B) problematic marketing techniques as-
7 sociated with the electronic nicotine delivery
8 system; and

9 “(C) the illegal presence or offering for
10 sale of the electronic nicotine delivery system in
11 a retail facility or online.

12 “(3) CONSUMER PRIVACY.—The Secretary—

13 “(A) shall maintain the confidentiality of
14 any individually identifiable consumer informa-
15 tion reported pursuant to this subsection; and

16 “(B) shall not require the inclusion of any
17 such information in a report under this sub-
18 section.”.

