AMENDMENT TO
RULES COMMITTEE PRINT 114-22
OFFERED BY MR. FITZPATRICK OF
PENNSYLVANIA

Page 235, after line 2, insert the following:

Subtitle R—Other Provisions

SEC. 2321. SENSE OF CONGRESS.

It is the sense of the Congress that the Food and Drug Administration, in coordination with the medical device industry, should prioritize and expedite the development of a robust, postmarket medical device surveillance system that can quickly identify and remove dangerous medical devices from market.