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 should—

(1) review the medical device reporting requirements under part 803 of title 21, Code of Federal Regulations; and

(2) in cooperation with appropriate law enforcement agencies, thoroughly investigate any instance in which such requirements were violated.