AMENDMENT TO RULES COMMITTEE PRINT 116–57

OFFERED BY MR. WENSTRUP OF OHIO

Page 485, after line 2, insert the following new subparagraphs (and revise the subsequent subparagraphs accordingly):

(D) an identification of any barriers that exist to manufacture finished drugs, biological products, vaccines, and critical medical supplies in the United States, including with respect to regulatory barriers by the Federal Government and whether the raw materials may be found in the United States;

(E) an identification of potential partners of the United States with whom the United States can work with to realign the manufacturing capabilities of the United States for such finished drugs, biological products, vaccines, and critical medical supplies;