..... moves to amend H.F. No. 1138 as follows:

Page 4, delete lines 30 and 31 and insert:

"(b) Nothing in this section applies to manufacturers or distributors of a medical device as defined in the Federal Food, Drug, and Cosmetic Act, codified at United States Code, title 21, section 301 et seq., or a digital electronic product or software manufactured for use in a medical setting including diagnostic, monitoring, or control equipment or any product or service that they offer."