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118TH CONGRESS
1ST SESSION

H. R. 3723

To amend the Federal Food, Drug, and Cosmetic Act to expand the types of devices for which required labeling may be made available solely by electronic means.

IN THE HOUSE OF REPRESENTATIVES

MAY 25, 2023

Mr. OBERNOLTE (for himself, Ms. KUSTER, Mr. CRENSHAW, and Ms. CRAIG) introduced the following bill; which was referred to the Committee on Energy and Commerce

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A BILL

To amend the Federal Food, Drug, and Cosmetic Act to expand the types of devices for which required labeling may be made available solely by electronic means.

Be it enacted by the Senate and House of Representatives of the United States of America in Congress assembled,

SECTION 1. SHORT TITLE.

This Act may be cited as the ''Medical Device Electronic Labeling Act''.

SEC. 2. ALLOWING REQUIRED LABELING OF DEVICES TO BE MADE AVAILABLE SOLELY BY ELECTRONIC MEANS.

Section 502(f) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 352(f)) is amended by striking ''prescription devices intended for use in health care facilities or by a health care professional and required labeling for in vitro diagnostic devices intended for use by health care professionals or in blood establishments'' and inserting ''devices (including in vitro diagnostic devices)''.

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