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110TH CONGRESS
1ST SESSION

H. R. 2791

To amend the Federal Food, Drug, and Cosmetic Act to provide for establishment of a unique device identification system for medical devices.

IN THE HOUSE OF REPRESENTATIVES

JUNE 20, 2007

Ms. HOOLEY (for herself and Mr. DOYLE) 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 provide for establishment of a unique device identification system for medical devices.

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 ''Unique Device Identification System for Medical Devices Act of 2007''.

SEC. 2. UNIQUE DEVICE IDENTIFICATION SYSTEM.

Section 519 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360i) is amended—

(1) by redesignating subsection (f) as subsection (g); and

(2) by inserting after subsection (e) the following:

''Unique Device Identification System

''(f) The Secretary shall promulgate regulations establishing a unique device identification system for medical devices requiring the labeling of devices to bear a unique identifier.''.

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