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Abstracts

REPORT INCLUDES:

Detailed description of the regulatory requirements for marketing and registration of medical devices in the U.S.

Insights into the current regulations and comprehensive procedures for the registration, renewal or notification of the medical devices, along with the information on timeline and fee required

Knowledge about labelling and advertising regulations for the medical device and details of the process for registration of the product with any specific variation

Information on Federal Food, Drug, and Cosmetic Act (the Act) and how it works towards the betterment of society

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