

Medical Device Law Committee Mission

The purpose of the Medical Device Law Committee is, in part, to keep the medical device industry and legal practitioners up to date and provide practical advice concerning issues that are of interest to the industry. Such issues involve the U.S. Food and Drug Administration's regulation of medical devices, including the approval and post market surveillance of novel technologies such as mobile applications and combination products.

The Committee will seek to engage members in programming and activities to contribute to the development of sound laws, policies, and regulations concerning the medical device industry. These activities will provide members with opportunities to network with colleagues on the committee and with relevant external organizations and individuals.