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CTTC
Center for Technology Transfer & Commercialization
VANDERBILT

Understanding new FDA Guidelines: Medical Software, Apps & Medical Devices

Join Us for a Special Seminar!

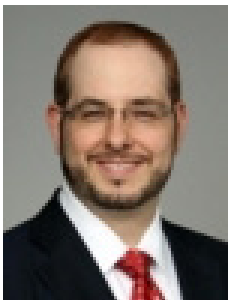
Seth Mailhot, J.D., Michael Best and Friedrich, LLP

Seminar: "When Does a Medical Software/APP Become a Medical Device: Practical Considerations and FDA Guidance"

Tuesday, Nov. 19
1:30 - 2:30 pm

Jacobs Believed in Me Auditorium, Featheringill Hall (FGH 134)

This seminar will address the regulatory framework for medical software, specifically for the rapidly expanding field of mobile apps. With Vanderbilt's significant involvement in software driven medicine and medical apps, this talk should be of interest to entrepreneurs, students, researchers, marketers and clinicians in this area.



Seth Mailhot, J.D., currently leads the FDA Regulatory practice at Michael Best and Friedrich, and is a member of the firm's Transactional Practice Group in the Washington, D.C. office. Prior to entering private practice, Mailhot worked for the U.S. Food and Drug Administration from 1994 to 2006. Mailhot has worked on FDA-related matters for more than 20 years, starting in various technical and enforcement positions at the U.S. Food and Drug Administration. He counsels clients on all aspects of regulation by FDA, as well as related matters regulated by USDA and Department of Health and Human Services.

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In partnership with Michael Best and Friedrich, LLP. Light refreshments will be provided.