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Andrew Grove is a Microbiologist in FDA's Office of In Vitro Diagnostic Device Evaluation and Safety. He is responsible for providing scientific review of preIDEs, 510(k)s and PMAs submitted to the Division of Microbiological Devices as well as acting as a hardware/software reviewer for the Division. Andrew received a B.A. in Biology from Eastern Mennonite University and a Ph.D. in Pharmacology and Toxicology from the Medical College of Virginia at Virginia Commonwealth University. After spending five years at a biotechnology company, he joined the Office of In Vitro Diagnostic Device Evaluation and Safety in 2007.