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William Shackelford is a Biologist in FDA's Office of In Vitro Diagnostic Device Evaluation and Safety. He is responsible for providing manufacturing and Quality System GMP compliance review for PMA and PMA Supplements, Recalls, Establishment Inspections, and other compliance actions submitted to the Division of Microbiology Devices. William received a B.S. in Biology from the University of Maryland and an M.S. in Biotechnology from the Johns Hopkins University. After spending thirty years in the IVD industry at several biotechnology companies performing IVD device design and development, validation and scale-up, device manufacturing, Quality Assurance, and Regulatory Affairs for IVD devices, he joined the Office of In Vitro Diagnostic Device Evaluation and Safety in 2009.