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JENNIFER DICKEY is a regulatory reviewer in the Office of *In Vitro* Diagnostics and Radiological Health (OIR) at the Center for Devices and Radiological Health (CDRH) in the Food and Drug Administration (FDA). Dr. Dickey reviews submissions in the Molecular Genetics Branch as part of the Division of Molecular Genetics and Pathology (DMGP). The Molecular Genetics Branch is responsible for reviewing a wide range of devices including next generation sequencing technologies and hybridization-based molecular techniques to detect genetic alterations associated with disease.

Dr. Dickey was the lead reviewer for the recent clearance of the Illumina Cystic Fibrosis Clinical Sequencing Assay. She additionally specializes in review of devices intended to aid in selection of therapy for patients with leukemia, and in the development of assays to determine absorbed radiation dose in crisis scenarios. Prior to her current position, Dr. Dickey was a reviewer in the Center for Drug Evaluation and Research (CDER) in the Office of Biotechnology Products. Dr. Dickey obtained her Ph.D. from Vanderbilt University and was a post-doctoral fellow at the National Cancer Institute.