Marianela Perez-Torres, Ph.D.

Dr. Perez-Torres joined the FDA's Center for Devices and Radiological Health in the Office of In Vitro Diagnostics and Radiological Health in 2016. Currently, as the Acting Director of the Division of Chemistry and Toxicology devices (DCTD), she is involved in a wide variety of regulatory activities for in-vitro diagnostic devices including premarket clearance/approval, manufacturer assistance and post-market compliance actions. In this role, she serves as the technical and regulatory authority on scientific review evaluation of a wide variety of clinical chemistry in vitro diagnostic devices including tests for therapeutic drugs, diabetes management, cardiovascular diseases, women's health, and liver function. Her educational background and professional experience are in the fields of Medical Technology, Biochemistry and Cancer Biology.