

Stuart Portnoy, MD, is Senior Director, Medical Device Consulting at PharmaNet where he advises medical device manufacturers regarding regulatory strategy, clinical trial design issues, and technical considerations to gain FDA market-approval for their products. He previously worked at the FDA for 8 years, most recently as Branch Chief of the Interventional Cardiology Devices Group and Acting Deputy Director of the Division of Cardiovascular Devices in the Center for Devices and Radiological Health.

At the FDA, Dr. Portnoy helped develop the FDA's approach to reviewing drug-eluting stents and other drug/device combination products. Prior to his experience as a manager, Dr. Portnoy specialized in the clinical review of cardiac electrophysiology devices including pacemakers, defibrillators, and EP catheters. Dr. Portnoy is a graduate of George Washington University School of Medicine and also has an MS in Bioengineering from the University of Pennsylvania.