FOR IMMEDIATE RELEASE

Spaulding Clinical Receives FDA 510(k) Clearance for 12-Lead ECG Device

Wireless, hand-held electrocardiograph designed to streamline the collection and transmission of ECG data for doctors and patients.

May 24, 2016 - West Bend, Wisconsin, USA - Spaulding Medical, LLC, a wholly owned subsidiary of Spaulding Clinical Research, LLC and provider of cardiac safety solutions, today announced the FDA 510(k) Clearance for their newest 12-Lead ECG device: the Spaulding Electrocardiograph 2100iQ™, and the device is now commercially available at www.SpauldingMedical.com.

"The Spaulding Electrocardiograph 2100iQ is the tangible culmination of our mission to deliver high-quality data in less time and at a reduced cost," said Randol Spaulding, CEO of Spaulding Clinical.

The Spaulding 12-Lead ECG product line was originally launched in 2011 and is now in use in over 35 countries. The Spaulding Electrocardiograph 2100iQ is optimized to visually collect real-time digital ECG data via Bluetooth® using iOS™ 7+, Android™ 4.2+ and Windows™ 7+ devices. Data is wirelessly uploaded to the Spaulding webECG™ management cloud for Cardiologist over-reads and is designed to integrate with electronic medical records and clinical information management systems.

"With our FDA 510(k) Clearance in place, we've reinforced our commitment to being an innovation leader in cardiac safety by providing patients, partners and physicians with a mobile ECG solution our competitors cannot," said Amanda Baltz, President of Spaulding Medical.

About Spaulding Medical

Spaulding Medical, LLC is a wholly owned subsidiary of Spaulding Clinical Research, LLC. Spaulding Medical is where innovative engineering meets purposeful design, ushering in a new paradigm in cardiac safety. The Mission: to make hospital quality cardiac care affordable and accessible by leveraging 150 collective years of cardiac safety expertise into viable health solutions.

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