

MRI Medical appoints Ira Duesler as Director of Regulatory Affairs and Quality Assurance

Tucson, Arizona September 14, 2010

For Immediate Release

MRI Medical, a leader in the design, development and manufacturing of complex medical devices, strengthened its executive team with the appointment of Ira Duesler, as Director of Regulatory Affairs and Quality Assurance (RA/QA).

Mr. Duesler will lead the company's efforts to streamline, expand, and strengthen its regulatory affairs and quality systems to support rapid growth.

Joseph Lee, MRI's President said, "Ira's appointment is a significant step in expanding our commitment to surpassing the highest quality standards like the ISO, CE, and FDA requirements. We welcome him on board and eagerly anticipate his contribution. His guidance will accelerate our ongoing device developments in gastroenterology, minimally invasive surgery, cardiology and urology. Ira understands the challenges of getting a device to market and of creating a quality system that supports our customers' goals and does not overburden manufacturing operations."

Mr. Duesler has over 18 years experience in the medical device industry, having held positions with increasing responsibility from Quality Engineer to Quality Manager to Director of RA/QA to Director of Corporate Regulatory Affairs with medical device manufacturers. His experience ranges from quality assurance and engineering to domestic and international regulatory affairs and quality management system design and oversight. He has managed the compliance of various devices and technologies including ECG monitoring, pulse oximetry, electrosurgery, endoscopic/laparoscopic surgical instrumentation, TENS/NMES, defibrillation, and IV Administration

MR. Duesler is a member of the
Regulatory Affairs Professional Society (RAPS),
Association for the Advancement of Medical Instrumentation (AAMI),
American Society for Quality (ASQ), and a
former member of AAMI Technical Committee TC/210, Working Groups WG01 and
WG02 (Quality Systems)

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