

## AusMedtech2009

A ROADMAP FOR GLOBAL SUCCESS

Australia's Medical Technology National Conference

15 and 16 June

Sheraton on the Park, 161 Elizabeth Street on Hyde Park, Sydney

[www.ausmedtech2009.com.au](http://www.ausmedtech2009.com.au)



### **Dr Martin Devitt Medical Director Perficio Clinical & Regulatory Pty Ltd**



Martin graduated with a Bachelor of Medicine from the University of Newcastle (Australia) in 1997. He then went on to become a registrar in anaesthetics and intensive care.

In 2000, Martin joined the Therapeutic Goods Administration (TGA), a statutory body of the Australian Government that regulates the supply of therapeutic goods in Australia. He initially undertook a role as a medical advisor in the pharmaceutical area of the TGA (working with anti-infective pharmaceuticals, radiopharmaceuticals, and vaccines), then moved to the medical devices area of the TGA and eventually went on to become the Head of the medical devices Clinical Section, underpinned by a sound knowledge of the regulatory requirements for medical devices in Australia and internationally. His role included the provision of high level advice on clinical considerations in the regulation of medical devices; participation on the TGA's Medical Devices Expert Committee (MDEC) and Medical Device

Incident Review Committee (MDIRC); the development of legislation and policy initiatives both nationally and internationally, including input into the workings of the Global Harmonisation Task Force (GHTF); participation as a member on a committee of Standards Australia; and the auditing of quality management systems to ISO 13485.

In 2005, he joined Cook Australia (the Australian subsidiary of Cook Medical, a global medical devices company) to undertake the position of Medical Director in a global capacity, providing high-level clinical advice on pre- and post-market safety, quality, performance, and marketing issues pertaining to the manufacture and supply of medical devices globally.

Currently, Martin is Perficio Clinical & Regulatory's Medical Director. He possesses medical devices experience from both the regulator and the industry. His expertise encompasses clinical and regulatory input into all aspects of medical device research and development, manufacturing, and commercialisation, including clinical trials, clinical evidence, and post-marketing issues; health technology assessment of medical devices for reimbursement; and quality management system auditing to ISO 13485 and clinical trial auditing to ICH GCP. In addition to these roles, he also still practices clinically part-time in emergency medicine.