

La Tronche, June 15th, 2010

EndoControl, start-up company developing innovative robotics solution for MIS is recruiting a:

QUALITY/REGULATORY AFFAIRS MANAGER

- MISSIONS:

1. Regulatory affairs: **EndoControl** manufactures and commercializes medical devices in European (CE), USA (FDA, UL), Canada (CAN/CSA), Korea (kFDA),... New products under development are also expected to reach these markets within 6 to 12 months. You will be in charge of regulatory clearance processes, post market follow-up and traceability.

2. QMS: **EndoControl** is ISO 9001:2008 and ISO 13485:2003 certified. You will be in charge of our Quality Management System, its implementation and continuous improvement.

- SKILLS AND PROFESSIONAL EXPERIENCE:

Bachelor Degree

3+ years professional experience in medical device regulatory affairs

93/42/CEE and 2007/47/CE directive – class IIa

FDA and GMP requirements

EN60601-1, EN60601-1-1, EN60601-1-2 and UL60601-1 for electromedical devices

Fluent in English

Project Management: Rigorous and organized, autonomous and proactive

- LOCATION:

La Tronche (Grenoble, France)

- CONTACT:

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www.endocontrol-medical.com