

Research & Development Project Manager

Endologix, Inc. is a developer and manufacturer of minimally invasive treatments for aortic disorders. The company is located in Irvine, CA, has over 300 employees and is listed on NASDAQ (ELGX). Since launching in the U.S. in 2005, Endologix has experienced average annual growth of 60%+ and was the top performing med tech stock in 2009. The Company's Powerlink® System is an endovascular stent graft for the treatment of abdominal aortic aneurysms (AAA). AAA is a weakening of the wall of the aorta, the largest artery in the body, resulting in a balloon-like enlargement. More than 2 million people in the U.S. have AAA, with 200,000 new cases diagnosed every year. If left untreated, AAAs become increasingly susceptible to rupture. The overall patient mortality rate for ruptured AAA is approximately 75%, making it the 13th leading cause of death in the U.S. More information is available on the Company's website at www.endologix.com.

The R&D Project Manager plans, directs and implements all aspects of the company's design and development of new medical device products. Oversees the investigation and evaluation of existing technologies. Guides the conceptualization of new methodologies, materials, machines, processes or products. Directs the development of new concepts from initial design to product release. Manages feasibility studies of the design to determine if capable of function as intended. Monitors documentation maintenance throughout all phases of research and development. Organizes the coordination of activities with outside suppliers and consultant to ensure timely delivery. Selects, develops and evaluates personnel to ensure the efficient operation of the function.

Essential Duties and Responsibilities:

- Manages the creation and development of new invasive and noninvasive medical device product(s).
- Manages the investigation and evaluation of existing technologies.
- Manages the design and implementation of new methodologies, materials, machines, processes or products.
- Manages the development new concepts from initial design to market release.
- Maintains documentation of project plans and GLP design control throughout all phases of research and development.
- Interfaces with other departments to coordinate cross-functional project activities. Communicates project status to management.
- Mentors and supervises technical personnel.
- Experience with Endovascular Products.
- Identifies hazards and mitigates risk associated with identified hazards in Design and Process FMEA.

Education, Training, Skills and Experience Requirements:

- Bachelor in engineering or related field
- Knowledge of FDA and ISO GMP design control regulation
- At least 5 years of working experience in medical device development or related fields