



## QUALITY MANAGER

We are looking for a hands-on, self-starter Quality focused leader that can help support our ISO 13485 Quality Management System as well as help with the transition to the 2016 standard. This person would be responsible for the two facilities (Vaughn and Burlington), so ideally candidates would be best from the Mississauga or Brampton area.

### Responsibilities and requirements:

- Maintain the current Quality Management System (QMS) as well as guide the updates to meet the ISO 13485:2016 standard
- Work with the Site General Manager to prioritize the improvements and implementation plan
- Identify and implement Areas for Improvement within the Quality and Business Systems
- Perform Internal Audits and Supplier Audits
- Lead audits from ISO Registrars, Regulators and Customers
- Provide Quality-related training
- Travel between sites
- Travel to suppliers and customers ~ 20%
- Clearance to travel internationally

### Experience:

- 3-5 years in the Medical Device Industry or heavily regulated industry
- Manufacturing and / or repair service experience a must
- 2+ years leading and implementing Quality Improvements within the organization
- ASQ CQA or CQE Certified or Equivalent (and current)
- Experience working with small to medium size companies

To apply, please email a copy of your resume to **Bill White** at [bill@apex-careers.com](mailto:bill@apex-careers.com). Please ensure the subject line of your email reads: **QUALITY MANAGER**

**Job Posting Expiration: August 31, 2017**