

Medical Devices and Medical Equipment

The process of bringing a new medical device to market has a number of important milestones. In addition to product design, it is important to anticipate many of the obstacles that will need to be overcome to achieve success. The medical device industry is a rapidly growing, competitive and regulatory landscape, with shorter product lifecycles, increased regulatory scrutiny, and the need to manage complex partnerships across the value chain.



Our Approach

Thomas Group is a business process-improvement consulting firm; thus, our general approach is to look at processes and sub processes in-depth and then evaluate the effectiveness of each one. We examine how the processes interrelate with one another and find improvements and linkages that will result in enhanced business results.

For a company developing a new medical device, the ultimate goal is to ensure the safety, reliability, and quality of the product. Medical device companies work with Thomas Group to manage R&D, device design and control processes, and vendor selection and retention, as well as the intellectual property generated by innovation. We provide specific industry solutions that reduce the complexity of regulated business processes while reducing costs associated with production, design, and customer service.

How Thomas Group Can Help

Thomas Group can help ease the regulatory, clinical trial, and quality assurance burdens, allowing you to focus your time and resources on achieving your business goals. Thomas Group can also help overcome challenges associated with:

- Inefficient systems
- Contract manufacturing
- The high cost of validation

At Thomas Group, we are focused on assisting medical device, pharmaceutical, and equipment manufacturers across the globe with services that help them achieve efficiency in process improvement. We provide our clients with professionals who are armed with a proven and flexible consulting methodology. This approach enables us to provide clients a customized service that meets your specific needs.

Among the benefits realized by Thomas Group clients:

- Faster time-to-market
- Reduced costs
- Lower cost of regulatory compliance
- Lower costs due to poor quality
- Improved product quality
- Reduced noncompliance risk

Whether you need to define your regulatory pathway, support your product with a well designed clinical trial, prepare your team for an FDA or ISO audit, or ensure ongoing compliance, Thomas Group has the experience and knowledge that will help you achieve success. 