

## Galen Robotics Quality Assurance Department

<b>Position Title:</b>	Quality Engineer	<b>Date:</b>	
<b>Department:</b>	Quality Assurance	<b>Job Status:</b>	Full Time Employee
<b>Location:</b>	Baltimore MD	<b>Travel Required:</b>	Approximately 10%
		<b>FLSA:</b>	Exempt
<b>Reports To:</b>	Director, QA and RA	<b>Direct Reports:</b>	N/A
<b>Objective of Position:</b>	The Quality Engineer will focus on developing, implementing, and performing Quality Assurance and Quality Control related activities to support the products designed and manufactured by Galen Robotics.		
<b>Other position notes:</b>			
<i>This job description is not designed to cover or contain a comprehensive listing of activities, duties, or responsibilities that are required of the employee.</i>			
<b>Job Description</b>			
<p><b>Key Activities:</b></p> <ul style="list-style-type: none"> <li>• Assist in the continuous evaluation and improvement of the Galen Quality System.</li> <li>• Serve as an independent reviewer for design reviews.</li> <li>• Perform supplier evaluation related activities.</li> <li>• Develop metrics to evaluate supplier performance and perform supplier performance evaluations.</li> <li>• Maintain and update Supplier Performance Files.</li> <li>• Develop and execute incoming raw material inspection activities.</li> <li>• Develop and execute in-process manufacturing quality inspection activities.</li> <li>• Participate in Risk Management, Design Verification, Transfer and Change activities.</li> </ul> <p><b>Minimum Education and Experience Requirements:</b></p> <p><b>Education:</b></p> <ul style="list-style-type: none"> <li>• Bachelor’s Degree in engineering or scientific discipline.</li> </ul> <p><b>Experience:</b></p> <ul style="list-style-type: none"> <li>• 3 years of medical device, pharmaceutical or other regulated industry (e.g., automotive, nuclear, telecom, defense, etc.) experience</li> <li>• 2 years of quality assurance and/or quality control experience in a regulated industry (e.g., medical device, pharmaceutical development, automotive, etc.)</li> </ul> <p><b>Additional Desired Skills/Qualifications/Competencies:</b></p> <ul style="list-style-type: none"> <li>• Medical Device product design experience</li> <li>• ASQ CQE/CSQE/CQA Certification</li> <li>• ISO 13485:2016 Auditor or Lead Auditor Certification</li> <li>• Lean and/or 6-Sigma training or certification</li> </ul>			

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**Physical Requirements:**

- Must be able to lift 30 lbs.
- Ability to sit at a desk and/or stand at a manufacturing location for extended periods of time.  
**NOTE:** Reasonable accommodations may be made to enable individuals with disabilities to perform this task

Reviewed By (HR):		Date:	
Approved By (MGR):		Date:	
Last Updated By:		Date/Time:	