

Job Title: Manufacturing Associate III

FLSA Status: Non-Exempt EEO Category: Technician

Management Level Position: ☐ Yes ☒ No Reports to: Manufacturing Supervisor Department or Division: Manufacturing Job Description Effective Date: 11/01/2021

POSITION DESCRIPTION:

A Manufacturing Associate III produces standard and/or custom medical devices to meet customer and Quality requirements by following standard work instructions and company policies while manufacturing Argen products in a team environment.

ESSENTIAL DUTIES AND RESPONSIBILITIES:

- Follows department SOPs and work instructions and adheres to customer and Argen standards for manufacturing products.
- Adheres to work instructions to properly operate and maintain equipment as needed.
- Visually inspects product quality & escalates issues.
- Identifies and recommends disposition of defective items for rework or scrap.
- Understands the product manufacturing process from initial steps through final inspection.
- Completes all documentation and training as required.
- Maintains a safe and clean work area, wears appropriate PPE, and adheres to safety standards.
- Cross trains in manufacturing procedures.
- Other duties as assigned

EXPERIENCE & QUALIFICATIONS:

- High school diploma or equivalent required.
- **Two** plus years of experience working in a manufacturing environment required. **Four** plus years of experience preferred.
- Proficiency in **ten or more** areas of Argen manufacturing.
- Leads a manufacturing team in achieving results in a high-volume manufacturing environment
- Possesses leadership qualities such as self-awareness, team focus, ability to translate learning into action, and train others to do so, as well as drive success.
- Ability to work in a team environment.
- Ability to follow instructions and readily accept additional responsibilities.
- Attention to detail and quality focused.
- Passionate about industry and desire to contribute where needed.
- Schedule adherence and dependability.
- Ability to meet tight deadlines and meet production goals.

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- Ability to learn technical concepts by reading work instructions and standard operating procedures, and completing on-the-job training.
- Ability to follow detailed directions in a Good Manufacturing Practices (GMP) environment required.
- Knowledge of cGMP regulations ISO 13485, 21CFR Part 820, CMDR SOR/98-282, 93/42/EEC, RDC 16 2013, MHLW MO 169 and TG(MD)R Sch3 preferred.

Personal Protective Equipment Required ☑ Yes ☐ No	
If Yes Describe: Eye Protection	

PHYSICAL REQUIREMENTS, ENVIRONMENT & WORKING CONDITIONS			
Description	Regularly	Frequently	Occasionally
Sitting	Х		
Standing		Х	
Walking		Х	
Climbing/Balancing		Х	
Reaching-with arms & hands	Х		
Stooping/Kneeling/Crouching/Crawling			Х
Talking	Х		
Hearing	Х		
Feeling/Touching	Х		
Vision-Close, Peripheral, Depth, Ability to Adjust Focus	Х		
Light to moderate lifting (50lbs or less)			Х
Moderate to Heavy Lifting (more than 50lbs)			
Travel Required			

Environment & Working Conditions	Applicable
Loud noise level	X
Overtime	X
Working in a factory environment	X
Ability to work in a confined area	X
Ability to sit at a computer for an extended period of time	X
Exposure to airborne powder (non-toxic)	X
Work near moving mechanical parts	X
Ability to sit and work on one machine for an extended period of time	X
Ability to stand for an extended Period of Time	X

The intent of this job description is to provide a representative and level of the types of duties and responsibilities that will be required of positions given this title and shall not be construed as a declaration of the total of the specific duties and responsibilities of any particular position. An employee may be directed to perform job-related tasks other than those specifically present in this description.

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Change History

Version	Change Description
1	Initial Release
2	Header and Footer, removed signature line, JD effective date CO# 201

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