

Job Title: Manufacturing Supervisor I FLSA Status: Exempt EEO Category: First Level Supervisor/Manager Management Level Position: ⊠ Yes □ No Reports to: Production Manager or Operations Director Department or Division: Manufacturing Job Description Effective Date: 02/28/23

## **POSITION DESCRIPTION:**

The Manufacturing Supervisor I supervises operations and production of medical devices to ensure the safe, timely, and quality completion of all production work by effectively utilizing people, equipment and materials to meet company standards.

### **ESSENTIAL DUTIES AND RESPONSIBILITIES:**

- Supervises the department's day to day operations.
- Ensures that employees follow department work instructions and adhere to customer and industry standards for dental product.
- Supervises employee activities including evaluation and overall performance management, training and development, initiating or suggesting plans to motivate workers to achieve goals, and effective communication with associates.
- Understands and drives manufacturing philosophy of Safety (number 1), Quality and Delivery.
- Understands the product manufacturing process from initial steps through final inspection.
- Assures company assets (equipment and material) are used and maintained appropriately and employees promptly report Non-Conforming material (NCR) and/or equipment issues.
- Establishes or adjusts work procedures to meet production schedules and goals.
- Makes recommendations to management to improve production methods, equipment performance, overall work quality, working conditions, and use of equipment.
- Coordinates time and production records and ensures that job assignments are clearly communicated to employees in the department.
- Coordinates maintenance activities with department requirements and schedules.
- Assists employees with diagnosing malfunctions in machinery and equipment and alerts maintenance and management of problems with equipment.
- Assures the accurate and timely reporting of various records which may including time sheets, production reports, rework reports, check sheets, employee job transfers, etc.
- Coordinates and collaborates with other departments in establishing and carrying out responsibilities.
- Maintains safe working conditions in assigned area by continually working on safety awareness with associates and enforcing safety regulations.
- Develops high performing work teams and appropriate performance monitoring systems.
- Motivates and coaches employees by assessing employee performance and providing helpful feedback and training opportunities.
- Delegates tasks and manages progress to ensure successful completion of department objectives.

- Creates and reinforces a culture of teamwork and actively resolves conflicts within the team.
- Manages proactively by holding regular 1on1s and team meetings to ensure open lines of communication and professional development for direct reports.
- Other duties as assigned.

### **EXPERIENCE & QUALIFICATIONS:**

ARGEN

- High School (HS) diploma required or equivalent.
- One plus years of supervisory experience in a high-volume, low automation assembly or light manufacturing environment.
- One plus years of people management experience preferably supervising teams of 10+ employees.
- Experience leading people through coaching, utilizing the employee performance tool, hiring and onboarding.
- Motivates and coaches employees by assessing employee performance and providing helpful feedback and training opportunities.
- Delegates tasks and manages progress to ensure successful completion of department objectives.
- Manufacturing certification or related training preferred.
- Proficient in MS Office (Word, Excel, Outlook, PowerPoint).
- Experience with an ERP computer program is a plus.
- Excellent communication skills both verbal and written.
- Problem solving capabilities.
- Passionate about industry and desire to contribute where needed.
- Schedule adherence and dependability.
- Ability to meet tight deadlines and meet production goals.
- Ability to follow detailed directions in a manufacturing Good Manufacturing Practices (GMP) environment is required.
- Ability to learn technical concepts by reading work instructions and standard operating procedures, and completing on-the-job training.
- 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: Depending on department: Eye Protection, Gloves

PHYSICAL REQUIREMENTS, ENVIRONMENT & WORKING CONDITIONS			
Description	Regularly	Frequently	Occasionally
Sitting		X	
Standing	Х		
Walking		X	
Climbing/Balancing			Х
Reaching-with arms & hands		X	
Stooping/Kneeling/Crouching/Crawling			Х

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Talking	X		
Hearing	x		
Feeling/Touching		х	
Vision-Close, Peripheral, Depth, Ability to Adjust Focus	x		
Light to moderate lifting (50lbs or less)	x		
Moderate to Heavy Lifting (more than 50lbs)			Х
Travel Required			Х

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

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.

#### **Change History**

Version	Date of Change	Change Description	
1	10/18/2018	Initial Release	
2	10/1/2019	Header and Footer, removed signature line, JD effective date	