

**Job title**

Supplier Development Engineer

Edit

**Company**

WishBone Medical, Inc

Edit

**Job location**

100 Capital Drive  
Warsaw, IN 46582

Edit

**Visible on job post:** No  
**Remote work allowed:** No

**Advertising location**

Warsaw, IN 46582

**Job Type**

Full-time

Edit

**Job description**

**JOB TITLE:** Supplier Development Engineer  
**BUSINESS DIVISION:** WISHBONE MEDICAL INC **Status:** FULL-TIME  
**DEPARTMENT NAME:** Supply Chain **Type:** EXEMPT (SALARY)  
**WORK LOCATION:** CORP - WARSAW **REPORTS TO:** VP OF SUPPLY CHAIN

Edit

**SUMMARY:** The position of Supplier Development Engineer is responsible for working with the suppliers and providing direction to establish a culture of continuous improvement with alignment to medical device regulatory controls. The goal is to create a trusting relationship with emphasis on removing waste and variability from the supply chain flow without product delay while reducing costs and maintaining quality.

This position is responsible for mentoring the supplier chain team in key project improvements in regulatory control, quality, lead time, on-time delivery and costs attributed to 'waste'. The position will engage external suppliers and internal partners to meet critical operational objectives utilizing supply chain development processes and tools.

**ESSENTIAL DUTIES AND RESPONSIBILITIES:** *(include the following. Other duties may be assigned.)*

- Direct, coach and mentor Supplier Development personnel.
- Develop, manage, audit and maintain Approved Supplier List.
- Create and monitor performance metrics for suppliers including manufacturing, sterile packaging, and distribution.

- Own Quality Management System elements related to suppliers (procedures, work instructions, agreements, evaluations, etc)
- Work with suppliers to resolve nonconformities, deviations, and Supplier Corrective Action Requests
- Develop and apply advanced technical principles, theories and concepts to improve quality, process control and product acceptance.
- Continually seek and evaluate potential new suppliers to ensure latest technologies are in use and backup resources are in place.
- Monitor and manage change control with suppliers and product development.

#### QUALIFICATION REQUIREMENTS:

- Degree - Preferred Bachelors in Engineering Technology
- Medical Manufacturing and Quality Engineering Expert
- 5+ years' experience in equipment/process validations, measurement system analysis, manufacturing process control, and production planning.

**OTHER SKILLS AND ABILITIES:** To perform this job successfully, an individual must demonstrate an exceptionally high proficiency in creative problem solving and communication. The individual must be able to perform each essential duty satisfactorily. The requirements listed below are representative of the knowledge, skill, and/or ability required. Reasonable accommodations may be made to enable individuals with disabilities to perform the essential functions.

- Knowledge and experience working with ISO 14971, 13485, 10993 and 21CFR Part 820 regulations.
- Analyze and interpret engineering drawing specifications per ANSI 14.5 and established drafting standards.
- Project Planning Experience
- Quality Control
- Validation
- Statistics

**PHYSICAL DEMANDS:** The physical demands described here are representative of those that must be met by an employee to successfully perform the essential functions of this job. Reasonable accommodations may be made to enable individuals with disabilities to perform the essential functions. While performing the duties of this job, the employee is regularly required to sit and talk or hear. The employee frequently is required to stand; walk; and use hands to finger, handle, or feel objects, tools, or controls. The employee is occasionally required to reach with hands and arms. The employee must occasionally lift and/or move up to 20 pounds.

**WORK ENVIRONMENT:** The work environment characteristics described here are representative of those an employee encounters while performing the essential functions of this job. Reasonable accommodations may be made to enable individuals with disabilities to perform the essential functions. Associate will normally work in an office environment but may also be subject to high noise levels from machines, and physical hazards from moving machine parts. WHEN OVERTIME IS SCHEDULED, IT IS MANDATORY.

---

#### Applicant Qualifications

[Edit](#)

You have requested that Indeed ask candidates the following questions:

- What is the highest level of education you have completed?
- Please list 2-3 dates and time ranges that you could do an interview.
- Are you authorized to work in the United States?
- How many years of Medical Device experience do you have?

---

#### Language

[Edit](#)

English

---

#### Expected Hiring Date

[Edit](#)

1 to 2 weeks

---

**Hires Needed**

Edit

1

---

**Schedule**

Edit

8 hour shift, Monday to Friday

---

**Application settings**

Edit

Apply method: **Email**  
Send updates to: **marywetzl@wishbonemedical.com**  
Employer Assist: **14 days**

Do you want applicants to submit a resume? **Yes**  
Do you want to let applicants start the conversation? **Yes**

---

**Selected Assessment Types**

Edit

Work style: Conscientiousness, Analyzing data

---

**Job Budget**

[Sponsor this job for more candidates](#)

---

By clicking "Confirm", you agree to candidates appearing in your dashboard based on the preferences you've selected above. You also agree to our [Indeed Terms of Service](#).

Cancel

View Preview

Confirm