

**Ready, Set, Grow!**



**VESTA NUTRA** represents a customer-centric contract manufacturer of dietary supplements and a world-class ingredient distributor under the combined endeavor to ensure high-quality nutraceutical supplements at affordable prices, across the globe. We take responsibility for delivering outstanding products and results to our partners and the consuming public while working with our partners to generate innovation within the supplement industry.

**VESTA NUTRA** is ready to meet the challenges of the dynamic global nutritional supplement market, from research, development, supply chain, production, packaging, and regulatory conditions.

**VESTA INGREDIENTS, INC.** understands there are purer, more effective, and more potent ingredients just waiting to be discovered. Skilled research and development staff are constantly investigating emerging scientific and botanical data in order to formulate the next breakthrough dietary supplement your customers will demand.

**VESTA PHARMACEUTICALS, INC.** is a customer-centric contract manufacturer of dietary supplements has grown into a premier cGMP and FDA registered (21CFR111) full-service provider of dietary supplements and nutritional ingredients. Thriving in the growth of the private label nutraceutical space has provided the ability to meet our customer's needs in a timely and price-efficient manner.



### **Quality specialist (FI4010)**

**On-site 5767 Thunderbird Road, Indianapolis, IN 46236**

QA person helps to formulate what the actual batch "drugs" get blended into making the pills/tablets. They formulate the raw ingredients list.

#### **GENERAL DESCRIPTION:**

- Assist Quality Manager with FDA cGMP Compliance
- Assist Quality Manager to maintain cGMP Certification
- Raw material and Finished Product sampling and testing
- Material Disposition

- Document Control
- Assist with generation of controlled documents – Specification Sheets, SOPs, Work Instructions, Forms, etc
- Manage Retention Samples
- Conduct Line Clearance and ATP testing
- Assist with monitoring production to maintain product quality and cGMP compliance
- Conduct Internal Audits as Assigned by Quality Manger
- Complete cGMP and SOP Training – refer to Training Matrix for complete list of SOP training required for Quality Assurance Assistant
- Assist with conducting internal audits
- Complete Product Formula calculation prior to preparing the batch record
- Additional duties as assigned

**WORK EXPERIENCE REQUIREMENTS:**

- At least 3 year(s) of relevant experience

**EDUCATION REQUIREMENTS:**

- Bachelor's degree in Chemical/Biology/Pharmacology (Preferred) or closely related field

**HELPFUL SKILLS:**

- Experience working in food manufacturing or familiarity with FDA and cGMP is a plus
- Detail oriented and ability to thrive in a team environment
- Good knowledge of Microsoft Office applications
- Proficient in technology and general office equipment
- Excellent verbal and written communication skills
- Strong organizational and time management skills
- Ability to work efficiently with minimal supervision

Apply today!

<https://vestanutra.com/careers/>