Crown Laboratories, along with our affiliated companies, has a long-standing heritage in skin care, and we are passionate about providing consumers and clinicians with innovative solutions for the skin. As our mission states, we are committed to developing and providing a diverse portfolio of pharmaceutical and consumer products that improve the quality of life for our customers.

With consumer brands like Blue Lizard Australian Sunscreen, Vita Liberata, and more, we offer best-in-class sunscreens, self-tanning lotions, moisturizers, cleansers and oral care products. Our prescription products provide healthcare professionals with solutions for managing a variety of skin conditions. With the recent addition of Bellus Medical, we are now a leading innovator in aesthetics. We are excited about our future, as we grow to become an innovative leader in the dermatology, skin care, and aesthetics industries while enriching the lives we touch.

We are currently recruiting for **Quality Specialist** for our night shift operation which would be responsible for performing Quality compliance activities on the plant floor and ensuring that all activities are in compliance with company and regulatory guidelines and procedures. In addition, the QA Specialist is also responsible for assisting in activities directly related to batch release, batch record & procedure review, revision and approval.

Responsibilities:

- Perform audits/checks of manufacturing, filling and packaging processes to include QA weight checks, line clearance activities, record review and signoff, cleaning verification label verification and reconciliation verification.
- Approve minor deviations on the operation floor.
- Review and approve executed batch records for completeness and accuracy for batch release.
- Review, revise and write batch records, procedures and policies in support of continuous Quality Improvement.
- Verify and ensure compliance with regulatory and company procedures and guidelines.
- Perform investigations as required.
- Assist in other quality related functions as requested, (sampling, component receipt and inspection, etc.)
- Additional tasks as assigned by Quality Management.

Qualifications/Requirements:

- Bachelor's degree in a scientific or related discipline from an accredited college or university OR
- Minimum of 3-7 years' experience in the Pharmaceutical or regulated industry with prior experience in plant quality assurance.
- Demonstrated knowledge of cGMPs, ICH and other regulatory requirements for the manufacture, testing and release of pharmaceutical products.
- Strong proficiency in Microsoft Office suite of products, (Word, Excel, PowerPoint)
- Excellent verbal and written communication skills.
- Excellent interpersonal, communications and organizational skills as well as the ability to work independently.
- Occasionally lift 15-25 lbs. Must be able to stand for long periods of time.

NOTE: This job profile is intended to provide an overview of expected job duties and requirements. It is not intended to be a contract of employment, explicit or implicit. All contents are subject to change at the sole discretion of the company. Cooperation is expected of all employees. Other duties may be assigned as needed.

We offer a competitive benefits package including vacation, medical, dental, vision, short/long term disability and 401(k) w/match.

an Equal Opportunity Employer

For other career opportunities, please visit our careers page:
https://www.crownlaboratories.com/employment-opportunities

To learn more about our company and our brands, please visit our websites:
www.crownlaboratories.com
www.bluelizard.net
www.vitaliberata.com
www.nkskn.com
www.bellusmedical.com

Apply here: https://www.crownlaboratories.com/application-for-employment