Our company is searching for experienced candidates for the position of quality specialist senior. Please review the list of responsibilities and qualifications. While this is our ideal list, we will consider candidates that do not necessarily have all of the qualifications, but have sufficient experience and talent.

Responsibilities for quality specialist senior

- Develops, implements and ensures compliance of quality assurance plans for production processes to ensure product specifications (quality and safety) are met
- Identify and implement corrective actions to reverse negative trends in manufacturing related consumer complaints
- Contribute and support internal and external team Sharepoint sites
- Perform specialty audits, as requested by management
- Execute on supplier qualification activities, such as Supplier Corrective Action Requests, performing supplier audits, and writing audit reports
- Interact with lab operations personnel to ensure compliance with company quality systems policies and procedures
- Support final review and release of material produced in-house and at contract manufacturers
- Develop/Execute automated tests, identify critical bugs quickly and work with Engineers to resolve them
- Develop effective tests to cover all scenarios for both iOS and Android Mobile applications
- Stay on top of all bugs, help triage bugs and driving issues to resolution and keeping everyone involved aware of QA status

Qualifications for quality specialist senior

- Minimum 5 8 years' experience in pharmaceutical GMP-regulated industries (Biotech preferred) and minimum 5 years in Quality Assurance or Compliance
- Hands-on working experience with GMP documentation and/or training systems is a must
- BS/MS in Chemistry, Biology or related science in a relevant discipline
- Requires ability to work independently and on teams, managing multiple projects/tasks simultaneously with competing priorities
- Experience with Pilgrim SmartSolve and TrackWise QMS preferred