



Phone: (555) 234-5678

Email: michael.anderson@email.com

Address: San Francisco, CA

Website: www.michaelanderson.com

## **EXPERTISE SKILLS**

- Sterile Manufacturing
- ISO Standards
- Data Review
- Internal Audits
- Method Validation
- Production Support

## **LANGUAGES**

- English
- Spanish
- French

## **CERTIFICATION**

- Bachelor of Science in Biotechnology, University of California, Davis, 2014

## **REFERENCES**

### **John Smith**

Senior Manager, Tech Corp  
john.smith@email.com

### **Sarah Johnson**

Director, Innovation Labs  
sarah.j@email.com

### **Michael Brown**

VP Engineering, Solutions Inc  
mbrown@email.com

# MICHAEL ANDERSON

## QUALITY CONTROL SUPERVISOR

Highly skilled Biotechnology Quality Control Analyst with over six years of dedicated experience in the field of biopharmaceutical manufacturing.

Recognized for a comprehensive understanding of quality control methodologies and regulatory requirements, particularly in the context of sterile product production. Expertise in performing complex analytical techniques and inspections to ensure product safety and effectiveness.

## **PROFESSIONAL EXPERIENCE**

### **MedImmune**

*Mar 2018 - Present*

Quality Control Supervisor

- Supervised quality control activities, ensuring compliance with ISO and FDA standards.
- Reviewed and approved laboratory data, ensuring accuracy and reliability of results.
- Implemented training programs for staff on quality control processes and regulatory requirements.
- Conducted internal audits to assess compliance with quality management systems.
- Managed the investigation of quality deviations and implemented corrective actions.
- Collaborated with production teams to resolve quality-related issues in real-time.

### **Amgen**

*Dec 2015 - Jan 2018*

Quality Control Analyst

- Conducted routine laboratory testing of pharmaceuticals to ensure compliance with specifications.
- Participated in method validation and stability testing for new products.
- Maintained accurate records of laboratory activities in accordance with GLP.
- Collaborated with quality assurance teams to prepare for regulatory inspections.
- Analyzed data for trends and anomalies to support quality initiatives.
- Provided technical support during production scale-up and validation processes.

## **ACHIEVEMENTS**

- Achieved a 20% reduction in non-conformance reports through enhanced quality control measures.
- Played a pivotal role in obtaining ISO certification for the quality management system.
- Recognized as 'Employee of the Year' for outstanding contributions to quality improvement efforts.