MA

CONTACT

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San Francisco, CA

SKILLS

- cGMP Compliance
- Cleanroom Commissioning
- Process Validation
- Team Collaboration
- Risk Assessment
- Regulatory Standards

LANGUAGES

- English
- Spanish
- French

EDUCATION

BACHELOR OF SCIENCE IN CHEMICAL ENGINEERING, STATE UNIVERSITY

ACHIEVEMENTS

- Successfully led a project that received the 'Best New Facility' award in 2020.
- Implemented a new validation process that reduced time to market by 10%.
- Recognized for outstanding contributions to safety with a company award in 2021.

MICHAEL ANDERSON

COMMISSIONING ENGINEER

PROFILE

I am a Chemical Commissioning Engineer with 8 years of experience in the pharmaceutical industry, specializing in the commissioning of cleanroom facilities and bioprocessing plants. My background in chemical engineering has equipped me with a thorough understanding of regulatory requirements and quality assurance protocols. I have successfully managed the commissioning of multiple projects, ensuring compliance with cGMP standards while optimizing processes to enhance product quality.

EXPERIENCE

COMMISSIONING ENGINEER

PharmaTech Solutions

2016 - Present

- Managed the commissioning of a new biopharmaceutical facility, achieving FDA compliance on first inspection.
- Developed and executed commissioning protocols for cleanroom environments, ensuring adherence to industry standards.
- Collaborated with QA teams to validate processes and maintain product integrity.
- Trained staff on equipment operation and safety procedures in a controlled environment.
- Implemented process improvements that reduced commissioning time by 15%.
- Maintained comprehensive documentation for all commissioning activities and outcomes.

PROCESS VALIDATION ENGINEER

BioLife Manufacturing

2014 - 2016

- Executed process validation protocols for production equipment, ensuring compliance with cGMP.
- Conducted risk assessments to identify potential process deviations and implement corrective actions.
- Collaborated with engineering teams to design and optimize bioprocessing systems.
- Developed training materials for staff on validation practices and regulatory requirements.
- Performed audits of internal procedures to ensure alignment with industry best practices.
- Presented findings and recommendations to senior management, driving continuous improvement initiatives.