Process development operator

Ropack Pharma Solutions provides a seamless drug development process for solid oral dosage, from early-stage R&D to commercial manufacturing and packaging, with a continuum of quality integrated throughout. Our extra-mile efforts deliver unwavering quality and reliability and the highest level of customer focus.

We are seeking a Process Development Operator in the Long Island, NY area to service the pharmaceutical industry. The process development operator will be responsible for the set-up, operation, cleaning of solid oral dosage manufacturing equipment, including, but not limited to, tablet presses, encapsulation equipment, aqueous coating equipment, and other processing equipment required in the manufacture of solid oral dosage forms. The Manufacturing Operator will execute the Production Batch Record as written without deviation, and document all specified tasks/actions as required. The Senior Operator will be executing the batch in partnership with the supervising scientist or other Manufacturing Operators. The manufacturing operator will ensure all cGMP’s, SOP’s, OSHA, regulations are followed at all times during the execution of the batch.

Essential duties

- Perform operation, cleaning, set-up and problem solving on manufacturing equipment including tablet presses, coating machines, granulation equipment, and encapsulation equipment.
- Responsibilities may include execution and review of manufacturing and packaging batch records.
- Participate in the investigation and resolution of formulation issues. Monitor the impact of formulation changes and propose improvements as required.
- Perform machine changeovers, cleaning and set up as required.
- Ensure adherence to quality standards during all stages of the manufacturing process.
- Complete manufacturing documents, ensuring accuracy and completeness.
- Maintain compliance at all times with related SOPs, GDP, and cGMPs in the manufacturing process.

Qualifications and Experience

- High School diploma, GED or associates degree;
- Minimum of 5 years working in a solid oral dosage pharmaceutical manufacturing environment;
- Thoroughness, attention to quality, a professional understanding of cGMP practices and concepts are required;
- Computer literacy and solid communication skills are a must;
- Ability to perform basic mathematical calculations;
Mechanically inclined candidates with solid oral dosage manufacturing experience are preferred.

Join a growing dynamic company that offers benefits, continuing education, a dynamic work environment and advancement opportunities.

Send us your application at rh-hr@ropack.com