Jesse Hampton

Manufacturing Manager and Engineer

Ambler, PA - Email me on Indeed: indeed.com/r/Jesse-Hampton/8bb212b324b4174d

A department manager and chemical engineer with over nine years direct cGMP experience in pharmaceuticalmanufacturing, packaging, and supply chain management. Demonstrated success in working collaborativelywith team members to drive quality and productivity improvement and to encourage team members to achievecompany objectives.• Production & Supply Chain Management: 11 • Lean Six Sigma Tools: Utilization of practical lean,years' experience in the pharmaceutical industry in 5s, and six sigma tools to drive continuousthe areas of production and supply chain improvement initiatives in the areas of productivity,management, technical operations, process CAPAs, and Quality.engineering, change control, and risk management. • Serialization: Led project management of• Production Tracking: Develop automated pharmaceutical serialization in collaboration withproduction tracking tools to perform statistical the Engineering department and equipmentanalysis and have a record of improving partners.productivity in three different organizations. • Software: Advanced proficiency in MRP/ERP• Investigations: Author and approve investigations (SAP/JDE) and in statistical analysis. Expert inand complete group root cause analysis as primary utilizing Excel (macro creation) to provide stafftechnical contact for seven years spanning across with the necessary tools for success.three different organizations.OPERATIONS EXPERIENCELANNETT COMPANY, INC., Philadelphia, PennsylvaniaManufacturing Manager […] packaging operations and SAP logistics for raw material and packaging components for the portfolio of 55active products with varying strengths and fill sizes, submission batches, and support product launches. Directedmanagement of 30+ personnel including supervisors, chem weigh technicians, packers, line leaders and materialhandlers. Supported technician execution of electronic batch records. Served as VAWD® representative supervisor forthe facility including schedule II vault and schedule III-V cage with inventory managed using SAP. Conducted thepreparation and annual inventory destruct of all controlled drug waste for packaging, manufacturing, and quality. Usedmaterial bulk handling equipment and forklift truck (certified).• Led startup of 2nd shift operations with minimal staff to support production growth as annual production planincreased from 400 million to 600 million annual doses.• FDA audits resulted in zero, 483 area observations attributed to both responsible departments.• Served as packaging representative for two successful DEA audits.• Led comprehensive effort to revise key operation SOPs to increase procedure robustness and to improve qualitysystem control Using feedback from the field• Developed an automated packaging production tool that tracks performance down to both the equipment andoperator levels, which led to decreased downtime by 30%; results were used for staffing and equipmentproposals.• Led capital acquisition projects for new equipment including equipment for serialization.• Led project management of serialization implementation project using MS Project to track progress andmaintain timelines.• Prepared first/second level performance reviews for entire staff and initiated performance management whennecessary.GLAXOSMITHKLINE (GSK), Clifton, New JerseyFirst Line Manager […] ten direct reports, compounders and pre-weigh associates. Conducted daily walkthroughs to sustain 5s andsafety improvements and performed risk assessments as appropriate. Used scoreboards and specialized planningsoftware to plan manufacturing operations. Designated department quality champion. Served as process engineeringrepresentative in executing six sigma kaizen events and in post-kaizen issue resolution for continuous improvementinitiatives.• Reduced process and testing errors by 75% affecting "Batch Not Right First Time" attributed to human error inthree months by designing and implementing a new performance management and incentive program.• Championed a program, "Quality Champion Program, " to give select associates opportunity for feedback onquality issue resolution within the department and to develop value added JDIs and CAPAs.• Led writing of investigations and change controls.• Performed daily 5s reviews of equipment downtime and maintenance status boards.• Managed shift training compliance and coached employees to maintain quality and performance.• Designed dynamic electronic supervisor carry over form.• Designed an excel spreadsheet that automatically trended weekly and

yearly downtime and performance,immediately printable for presentation.• Performed IQ/OQ/PQ activities for newly designed batch recipes.WATSON PHARMACEUTICALS, Carmel, New JerseyPackaging Supervisor […] as direct supervisor of 25 production employees. Wrote investigations and change controls. to provideall process engineering support for packaging.• Successfully planned and maintained production goals for one billion annual doses plan.• Updated preventative maintenance (PM) frequency requirements and configured automative PM system toschedule PMs.Manufacturing Engineer Associate […] as primary technical contact for validation and special projects. Supervised maintenance and productionpersonnel. Scheduled maintenance and GMP equipment procurement.• Developed consolidation scenarios for bottle size and recommended and tested the scenario that saved the […] Recognized as, "Best of Class, " for the JDI MRP implementation team.• Replaced packaging performance monitoring system with a unified system that semi-automatically monitoredproduction's individual and collective average, variance, and % variance of 5-Line / 4-Shift operation againststandard.• Authored the Standard Operating Procedure for Preventative Maintenance and unscheduled repairs for thepackaging department.

WORK EXPERIENCE

Manufacturing Manager

Lannett - Philadelphia, PA - July 2011 to Present

Managed Packaging and Chem-Weigh Operations for pharmaceutical drug manufacturing.

First Line Manager

GSK - Consumer Healthcare - Clifton, NJ - March 2009 to July 2011

Assistant

HOSPIRA - McPherson, KS - 2005 to 2007

Validated method for analytical test methods for injectable products per ICH and USP guidelines.

• Processed engineering validation for a solvent extraction process to separate API from the formulation.

• Tested a variety of product contact materials used in manufacturing to demonstrate API recovery rates for parenteral products.

EDUCATION

chemical engineering

university of oklahoma

2000 to 2004

LINKS

https://www.linkedin.com/in/HampJJ