

JOB POSTING REQUEST FORM

JOB DETAILS	
Reference #	CP-210-2022
Company Name	Malia Group
Industry	Pharmaceutical
Vacancy	Quality Control Analyst
Number of vacancies for that position	2
Job Type(FT, PT, Contractual, Seasonal)	FT
Major(s)	BS in Chemistry
Degree	Bachelor / Masters
Years of Experience	0 to 1
Location	Nahr Ibrahim
Remuneration & Benefits	
Currency (LBP, Dollar, Lollar)	
Tasks& Responsibilities	<p>On Sampling and Testing</p> <ul style="list-style-type: none"> • Perform sampling, testing and release of raw, packaging material, bulk, semi-finished products and issue Certificate of Analysis to ensure quality of products meet a well-defined set of standards • Perform sampling, testing and approval of manufactured products: Liquid, solid and semi-solid products to ensure quality of products meet set standards • Conduct In-Process control in the Packaging/Manufacturing departments (testing at beginning, during and after stoppage of machines) and analyze test result, ensuring both packaging and manufacturing processes are conforming to quality standards <p>On Material approval</p> <ul style="list-style-type: none"> • Review and approve Certificate of Analysis of materials to be purchased from suppliers • Issue Certificate of Analysis following set template and get approval of Quality Control Manager <p>On System update and documentation</p> <ul style="list-style-type: none"> • Update system with all information included in the Certificate of Analysis and

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upon release of material

- Update system with the bulk's shelf life
- Check status of all materials as per stock inventory sent by Quality Control Manager
- Accurately document the results of the testing performed and properly maintain all documents and test records in a timely manner
- Complete the Batch report with the needed quality information and submit to Quality Control Manager

On Updating of quality procedures

- Assist in the writing and updating of Quality Control Procedures:
 - a. Statistical Procedures
 - b. Analysis Procedures
 - c. Raw Materials analysis
 - d. In process Testing
 - e. Finished products analysis
 - f. Microbiological analysis
 - g. Stability Studies and protocols for raw materials and finished products
 - h. All other Quality Control Procedures including environment
- Participate in the construction and/or revision of SOPs for the quality function

On Equipment monitoring and calibration

- Monitor QC equipment to ensure proper operation and calibration
- Coordinate calibration of all test equipment and fixtures
- Conduct Cleaning and Analytical Validation of equipment and machines to ensure quality standards are implemented

On Inventory

- Monitor Reagents, Glassware and Standards inventory to ensure healthy stock levels
- Execute and fill in Test Method Transfer protocols as per Licensor's requirements

On Customer complaint

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- Participate in investigations following any received complaint, evaluate problems and make initial recommendations for possible corrective action

On Returns

- Handle the recall of products process

On Reporting:

- Prepare reports and submit to Quality Control Manager for review and approval:
 - a. Out of specification Report, OOS
 - b. Out of Trend Report, OOT
 - c. Deviation Incident Report
 - d. Change control or risk assessment report