Job Description

Quality Control

The Company:

Yapan Bio is a Biotechnology Company established to serve the rapidly expanding Custom Development and Manufacturing (CDMO) market for Vaccines and Biologics/Bio-therapeutics. Yapan Bio has established state-of-the-art facilities for process development and GMP manufacturing for human clinical studies, including both Drug Substance (DS) and Drug Product (DP - Liquid Vials), at Genome Valley in Hyderabad. The facility is built as per the quality and regulatory expectation of the global regulatory authorities. The current facility includes two manufacturing suites, including one Bio Safety Level 2+ (BSL-2+) suite to develop and manufacture high containment product classes.

As we expand our team, we are looking for talented and experienced individuals, with can- do attitude, who are self-starters with excellent communications & leadership skills, who enjoy working on the details without losing site of the larger organizational goals, and are adaptable to a dynamic "startup" environment, to join our team.

At Yapan Bio, you will get a supportive, collaborative, and empowering work environment to make a significant impact along with ownership as we drive the growth of the company together. This is an excellent opportunity to be a part of the team to establish a new biotechnology company in India, with global impact.

The Role:

The incumbent will be part of the team responsible for establishing and managing the analytical and quality control for the organization. The incumbent will support the Quality Head to deliver as per the corporate goals, projects, plans and timelines.

Duties and Responsibilities:

- Manage and oversee all Quality Related Testing right from IPQC to Final Testing.
- To work with cross functional teams to develop and update work progress, instructions and processes for quality assurance and management.
- Perform testing of the samples and document the same following quality requirements for internal and external customers.
- Calibration of pH meter/analytical balance other analytical instruments.
- Manage and request for inventory of all associated reagents, chemicals and consumables used for quality control testing.
- To follow all procedures, policies, guidelines, and quality systems defined by the management to support the smooth flow of operations.

Education and Experience Required:

Post graduate degree in Science/Engineering required; advanced degree is preferred.

3 to 5 years of Analytical/Quality Control experience in the vaccines and/or biologics industry.

Skills:

- Experience in quality control and GMP documentation activities.
- Hands on experience on SDS-PAGE, western blot, ELISA, protein estimation methods.
- Experience on BET testing,
- Cell culture based assays, and mycoplasma testing,
- Host cell proteins, Host cell DNA, PCR& RT-PCR methods is desirable.
- Preparation of SOPs, STPs and QC testing related documents
- Should be able to work in shifts and independently.
- Clear capability in team building and in being a team player.
- Demonstrated capability to think strategically and flexibly to achieve business goals.
- Demonstrated ability and willingness to handle ambiguities and pressures of a start-up environment.

The Next Steps:

If the above role fits your experience and expertise and you would like to build and grow your career with a dynamic start-up, working on the next generation products and technologies, please share your CV at hr@yapanbio.com.