Position:VP of Quality

Reports to: CEO

Job Responsibilities:

- Responsible for the formulation and implementation of the company's quality strategy, quality policy and quality objectives;
- Responsible for the establishment and improvement of the company's quality management system, and ensuring that all activities in drug life cycle, such as research and development, production, inspection, and non-clinical studies, meet the requirements of domestic and foreign laws and regulations;
- Responsible for the establishment and implementation of the company's GXP training management system, and the promotion and implementation of quality policies;
- Establish quality risk management system, Corrective and Preventive Action system, data integrity management system, supplier quality management system, adverse event reporting system and other quality management systems to ensure the effectiveness of implementation strategies;
- Responsible for organizing internal self-inspection and accepting external audit, coordinating various departments of the company to cooperate in the inspection, and implementing the rectification of non-conformities;
- Review and approve standard operating procedures, deviation reports, stability study reports, production inspection, validation plans and documents, monitoring equipment maintenance plans and other quality-related documents;
- Responsible for checking the release of materials and products to ensure the compliance of release procedures; Responsible for evaluating and approving material suppliers, and responsible for supplier management and making supplier annual audit plan;
- Responsible for daily management within the department and coordination between departments within the company;
- Responsible for quality related government affairs to establish and maintain good working relationships.

Job Requirements:

- Master degree or above, major in biology, pharmacy, chemistry, with rich experience in biopharmaceutical production or quality management.
- Familiar with biological product process such as antibody and/or ADC;
- Good English listening, speaking, reading, and writing skills, skilled in using office softwares;
- Careful and meticulous with good team communication skills and team spirit;
- Familiar with system, equipment, instrument verification/verification, familiar with computer system validation;
- Internationalization experience in biopharmaceutical industry;
- Familiar with GMP, GLP, and other norms of drug overall life cycle management, familiar with national pharmaceutical laws and regulations, understand the domestic and foreign development status and trend of biological products industry.

Company Background

BlissBio, founded in Hangzhou, Zhejiang Province in December 2017, is a clinical-stage biopharmaceutical company focusing on discovery, development, and commercialization of biotherapeutics with independent intellectual properties. BlissBio's core team has successful experience in product development and commercialization in this field, as well as a variety of technology platforms. BlissBio has close collaborations with domestic and foreign biopharmaceutical companies to develop internationally competitive BIC and FIC biotherapeutics.

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