

## about us

Shenzhen Salubris Pharmaceuticals (Stock Code: 002294), as a leading enterprise of independent innovation in the field of cardiovascular and cerebrovascular disease (Listed China pharmaceutical companies in the top 10), mainly focusing on small molecule drug development and production, and now on Achieving strategic transfer to biopharmaceutical research and development.

Chengdu Genekey Biotech (base in Chengdu) and Salubris Biotherapeutics (base in Maryland), are wholly owned subsidiaries of Salubris Pharmaceuticals. Salubris Biotherapeutics mainly responsible for the early discovery of innovative large molecule drugs. And Genekey Biotech has an R & D team with doctoral, master based 156 people. A 6000 square meters of pilot research and development sites with a total value of nearly \$100 million of advanced equipment supports our three R & D platform (protein drugs, antibody drugs and gene therapy drugs) for the development of the more than 20 projects. And an up-to-date production base (Genemen bio) of biological drugs in line with the requirements of GMP located in Suzhou.



## Open positions

Opening Positions	Job Description
Purification process director (Chengdu)	Manage the purification team
Cell culture process director (Chengdu)	Responsible for the development, expansion and production of large scale culture of mammalian cells, and solve key technical problems
Formulation director (Chengdu)	Be responsible for determining the prescription composition, the use of excipients, the prescription process research, and guide the researcher to formulate the ecific research and development program
Quality research director (Chengdu)	With the comprehensive management ability of Research team, proficient in the use and management of all kinds of experimental equipment
CMC-VP (Chengdu)	Make CMC research plans and project schedule, monitor CMC research process and execution.
Quality management VP (Chengdu)	Establish, develop and keep completing the quality management system.
Director of biological activity (Chengdu)	Responsible for the management of the company's biological activity detection
Structural analysis director (Chengdu)	Responsible for the structural analysis and management
Clinical research director(Beijing)	Assure all clinical research projects are carried under the national regulations, clinical research and the company's SOP
Clinical quality director (Beijing)	Set up the clinical quality assurance system, and supervise its effective operation
RA director (Beijing)	Manage all the registration and application affairs of the company
Production VP (Suzhou)	Overall responsibility for the company's production management
Director of technology transfer (Suzhou)	Responsible for R & D projects from the pilot to the industrialization of technology transfer

If interested, please contact us at [wangliwen@genekeybio.com](mailto:wangliwen@genekeybio.com) (HR)