RH Biopharma

Job Posting

Date May 14, 2013

COMPANY INTRODUCTION

The business scope of Ruenhuei Biotech Corp. Inc. (潤惠生技股份有限公司) is new pharmaceutical product development and commercialization in medical fields. The company chairperson, Dr. Jane Hsiao (許照惠) and the president, Dr. Sherry Ku (顧曼芹) together with 潤泰集團 established the company to source Taiwan drug research candidates and develop these for clinical evaluation and commercialization. The initial projects are spin-offs from TWi biotechnology, Inc. (安成生技) where Dr. Ku served as the President and Chief Scientific Officer. The 3 projects are the FLT-3 multiple kinase inhibitor for Acute Myelogenous Leukemia from Taiwan Industrial Technology Research Institute ("ITRI"), Cytotoxic Fibrillar Protein for Ovary Cancer from Academia Sinica ("AS") and Respiratory Syncytial Virus Vaccine from National Health Research Institute ("NHRI"). These company research partnerships are established in 2012 and continued the business strategy focusing on sourcing research candidates with feasibility of early clinical proof of concept while meeting unmet medical needs in humans.

JOB DESCRIPTION

1. Clinical Research Associates/Manager/Director
   Plan, conduct, and manage clinical studies inclusive of country-specific clinical feasibility evaluation, global clinical protocol development, investigator relationship, clinical study conducting and reporting.

2. Preclinical Research Associates/Manager/Director
   Plan, conduct and manage together with partner institutions and Contract Research Organizations, preclinical studies inclusive of in-vitro binding, cell line and in-vivo animal model studies, pharmacology, pharmacokinetic, tissue distribution, metabolism, immunogenicity, and toxicology studies.

3. Regulatory Associates/Manager/Director
   Plan, and execute regulatory strategies and prepare regulatory filing documents. Perform electronic Common Technical Document Filings for IND, bio-IND, IMPD, CTC, PL, NDA, ANDA. Assess impact of changes to drug regulations and standards in the industry and communicate with various regulatory authorities in preclinical and clinical development strategies.

CANDIDATE QUALIFICATION

Ph.D with 3-year job experience or M.S. with 10-year experience in related fields including biochemistry, biotechnology, bioengineering, pharmacology, pharmaceutical and regulatory sciences. Fluent in English writing and proficient in computer and literature search skills.

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