ANTABIO Developing tomorrow's antibacterials	**Clinical Pharmacologist** **(Full or Part-time)**
	January 2021

Antabio is a Labège, France-based biopharmaceutical company dedicated to the discovery of first-in-class small molecule compounds to treat severe infections caused by antibiotic-resistant bacterial pathogens. Antabio's novel compounds block the drug-resistance mechanisms and the virulence of pathogenic bacteria by targeting key bacterial pathways. Using its patented technology, antibacterial discovery expertise, and partnering-oriented business strategy, Antabio advances novel drugs that will offer solutions to the shortfall of the current antibacterial arsenal. Antabio is advancing its Pseudomonas elastase inhibitor (PEi) program for the treatment of *Pseudomonas aeruginosa* infections in Cystic Fibrosis patients towards the clinic which is funded through a grant from CARB-X. This is an innovative inhaled approach targeting a key Pseudomonas virulence factor, intended to be used as an adjunct to antibiotic treatments.

Antabio is seeking a **Clinical Pharmacologist** to join our R&D team to set up and co-ordinate future clinical trials under the responsibility of our Head of Clinical Development. The position could be full or part-time, depending on the experience of the applicant and the scope of the role they are able to undertake.

Requirements

- MD with in-depth experience of clinical pharmacology, including development of novel respiratory medicines
- ≥ 5 years' experience in the pharmaceutical and/or biotech industry
- Experience in the design, analysis, and reporting of clinical pharmacology studies, including experience with first-in-human studies for NCEs/NBEs
- Experience with pulmonary infectious diseases would be beneficial
- Experience with regulatory submissions including CTAs, INDs, and/or IMPDs
- Experience in applying regulatory guidances (FDA and EMA) to the design and interpretation of Phase 1 trials
- Fluent Oral and written English mandatory
- Able to work in globally distributed company environment
- Flexible with regards to work/home location, with regular meetings onsite in Toulouse required
- Excellent oral and written communications skills, including proven history of positive interactions with Phase 1 units, CROS, and ethics committees as the medical representative of the sponsor

Responsibilities

- Design of clinical development strategy for the PEi program
- Develop, oversee, and integrate PK/PD plans into overall drug development programs
- Responsible for leading the development of protocols, informed consents, clinical study reports, and investigator brochures
- Hands-on management and responsibility for all CRO activities with respect to development of protocols, case report forms, data management plans, safety monitoring, and oversee the conduct of Phase 1 studies
- Lead the development and oversight of the clinical pharmacology component of development programs
- Effectively collaborate as a member of cross functional project teams
- High quality review and interpretation of study results and incorporation of these data into overall strategy
- Oversee and supervise the activities of the clinical research scientist(s) responsible for Phase 1 studies