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| ANTABIO Developing tomorrow's antibacterials | **Clinical Pharmacologist** |
| | June 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 beta-lactamase inhibitor program for the treatment of resistant Gram-negative infections towards the clinic and is seeking a **Clinical Pharmacologist** to join our R&D team to set up and co-ordinate a Phase 1 clinical trial, reporting to Head of Clinical Development.

**Requirements**

- MD with in-depth experience of clinical pharmacology, including clinical pharmacology principles, PK, PK/PD, drug metabolism, and drug interactions
- ≥ 5 years' experience in the pharmaceutical and/or biotech industry
- Experience in the design, analysis, and reporting of clinical pharmacology studies, including experience with FIH studies for NCEs
- Infectious diseases experience would be beneficial
- Experience with regulatory submissions, including CTA and IMPD
- 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 travel to Antabio (Toulouse) on an occasional basis, as required

**Responsibilities**

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