

Full and independent support by applying three core principles on every project to ensure success:

CREATIVITY | CONSISTENCY | COLLABORATION

<u>collaborations@qPharmetra.com</u> www.qPharmetra.com Broad Clinical Pharmacology expertise throughout all stages of drug development

"It's great just to be able to pick up the phone and get an immediate, scientifically sound answer to my question"

Extensive Clinical Pharmacology experience including Neuroscience, Oncology, Respiratory, Cardiovascular Diseases, Infectious Diseases, Vaccines & Immunology

Creative solutions, Collaborative mind-set Consistent processes and deliverables

CAPABILITIES:

- Dose selection from FIH to Phase 3 using pharmacometric approaches
- Develop and strategize the clinical pharmacology plan including FIH (SAD, MAD), BE, food effect, DDI, TQT, special populations PK and ADME studies
- Provide PK support for biopharmaceutical decisions throughout drug development
- Contribute to regulatory submissions and meetings
- Support pediatric drug development
- Innovative clinical trial design and clinical trial simulations
- PK analysis, interpretation and reporting of PK results from clinical trials



Applying quantitative methods to deliver the right dose to the right patient

Increase efficiency within drug development

Scientifically sound application of regulatory guidelines

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