

METIS

LABORATORY SERVICES

Streamlining Your Path to Pharmaceutical Success

Metis Consulting Services goes beyond regulatory expertise to provide comprehensive laboratory services that support your entire pharmaceutical development journey. Whether you are developing small molecules or biologics, we offer a suite of solutions to streamline your process and ensure success.

Small Molecule Development Services:

- Formulation Development: Expertise in designing formulations for oral, topical, and parenteral delivery, ensuring stability and bioavailability.
- Analytical Method Development: Specialized techniques for quantifying and characterizing small molecules, ensuring quality and compliance.
- Regulatory Support: Navigating complex regulatory pathways, including FDA submissions and compliance with international standards.
- Scale-up and Manufacturing: Transitioning from lab- to production while maintaining product quality and efficiency.

Biologics Development Services:

- Process Development: Designing and optimizing manufacturing processes, including purification, and formulation.
- Analytical Characterization: Advanced techniques for analyzing biologics, such as ELISA and chromatography, ensuring purity and potency.

- Regulatory Strategy: Developing comprehensive regulatory strategies for biologics, including BLA submissions and compliance with ICH guidelines.

Cross-Functional Expertise:

- Project Management: Coordinating multidisciplinary teams to streamline development timelines and ensure project milestones are met.
- Quality Assurance: Support for implementing robust quality systems to maintain compliance with cGMP standards and regulatory requirements.
- Technology Transfer: Facilitating seamless transfer of processes and knowledge between development, manufacturing, and quality control.
- Supply Chain Management: Optimizing supply chain logistics to ensure timely delivery of materials and minimize risks of product shortages.
- Lifecycle Management: Supporting post-approval activities, including process optimization, variation filings, and life cycle extension strategies.

Do not let regulatory complexities or laboratory hurdles hinder your pharmaceutical development journey. Contact Metis Consulting Services today to unlock the full potential of your product and achieve regulatory success.

At METIS, our mission is to ensure quality, drug safety, pharmacovigilance, clinical, and data management have the tools to support innovation pathways, with a balance of humanity and lightheartedness along the way.



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AUDITS | DATA MANAGEMENT | PHARMACOVIGILANCE | QUALITY MANAGEMENT | REMS