

ABOUT

PharmaNet/i3, the inVentiv Health clinical segment, is recognized as a leading provider of global drug development services to pharmaceutical, biotechnology, generic drug, and medical device companies, offering therapeutically specialized capabilities for Phase I-IV clinical development, bioanalytical services, and staffing from a single clinical professional to an entire functional team. For intelligent solutions needed to accelerate high quality product development programs of all sizes around the world, *PharmaNet/i3 works for you.*

CAPABILITIES

Phase I-IIa

For proof-of-concept, First in Human (FIH) or bioequivalency studies, our extensive database of study participants and relationships with leading hospitals ensure rapid recruitment. In addition to a dedicated project manager for each study, we deploy a team of experts, and specialists in quality assurance, and scientific and regulatory affairs to custom-fit a program to your needs.

Phase IIb-III

PharmaNet/i3 combines deep therapeutic knowledge with a commitment to quality and proven operational expertise to meet our clients' product development goals. Our project teams approach their work with a deep understanding of the specific needs of each therapeutic category. We offer expertise and services for clinical trial design and a full range of clinical trial services, including biostatistics, clinical monitoring, data management, global safety and pharmacovigilance, regulatory consulting, medical writing, project management, and full-service patient recruitment and retention. Our exclusive access to UnitedHealth Care claims data allows you to have real-time, robust insight into the right patient populations and investigators. We will then consult with clients to apply the data and develop a protocol to set a strong foundation for an efficient, safe and cost-effective study. PharmaNet/i3's global footprint and therapeutic expertise progress drugs and medical devices closer to regulatory submission.

Phase IV

PharmaNet/i3 has a dedicated suite of services, including strategic and operational planning, observational studies and patient registries, health economics and outcomes research, safety/risk management and epidemiology, traditional interventional studies, and a team of experts to guide clients through the post-approval environment.

Strategic Partnerships/FSP

We draw on a set of unique tools and strategic partnership experiences to create a team that is tailored exclusively to our clients' business goals. This customer-focused team can handle a single function across several drug programs, a range of functions within a single therapeutic area, a complete staff lift-out or any other defined need—serving as a consistent, scalable resource. We currently have 32 active partnerships dating back to 1999.

Staffing

Whether a client needs a single clinical project manager to cover a spike in activity or an experienced team to complete a specific therapeutic study, we can help you find the right clinical development professionals, when needed. Powered by the capabilities of i3 Pharma Resourcing, MedFocus, and Smith Hanley, and more than 40 years of combined experience and success in the field, our staffing and business process outsourcing solutions are simply unmatched.

Bioanalytical

PharmaNet/i3 offers bioanalytical support through our two state-of-the-art GLP-compliant bioanalytical laboratories, an extensive list of validated assays, knowledgeable scientists, and skilled technicians. Our experts develop, optimize, and validate analytical methods and rapidly process sample analysis of drugs from toxicokinetic, PK, bioavailability, bioequivalence, and all stages of clinical studies for both small and large molecules. *continued on next page*

Technology

PharmaNet/i3 provides comprehensive software support services for a variety of clinical trial data management and EDC systems. Our technology services are supported by fully validated business continuity plans, redundant data storage and timely backup procedures to ensure the integrity of your data. In addition, we offer a complete IVRS/IWRS and eDiary solution that is easy to deploy, scalable and 21 CFR Part 11 compliant.

Corporate Responsibility

As a pharmaceutical company, corporate responsibility and legal and ethical conduct shape how we conduct business. Our commitment consists of integrated policies and practices for ethical conduct, regulatory compliance, internal controls, risk assessment and prevention, continual improvement and communication.

THERAPEUTIC AREAS OF EXPERTISE

- Oncology
- Neurosciences
- Cardiovascular
- Endocrinology/Metabolism
- Rheumatology
- Pain Management
- Infectious Diseases/Vaccines
- Respiratory/Pulmonology
- Dermatology
- Ophthalmology
- Urology
- Nephrology
- Women's Health

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