CONSULTING SERVICES

Clinical Research-Clinical Affairs Services

South 6 Life Sciences offers a complete suite of clinical affairs and clinical research services, from individual consultants to project teams.
Our consultants can fill knowledge gaps on current teams, or work independently under our clients' direction as biostatisticians, statistical programmers, clinical data managers and professionals, study designers, and more.
Our consultants have extensive hands-on experience working in FDA-regulated industries including pharmaceutical, medical device, biotechnology, and related life science organizations.

South 6 recruits highly skilled consultants with expertise within the following areas:

Biometrics
Biostatistics, statistical programming, clinical data management, clinical applications development, document control, and clinical programming

Clinical Operations
Study design, protocol development, and all clinical research monitoring and duties, project management, clinical trial management, managing CROs, vendors and budget, close-out of studies, and working phases I, II, III, and IV

Medical Affairs
Medical program coordination, adverse event reporting, writing medical documentation, internal and submission level documents, and SOPs, quality assurance auditing, and regulatory affairs reporting and labeling

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