

Corporate Profile

Precherche Life Sciences, LP is a clinical development organization providing the first complete digital trial management (DTM) platform. Precherche's solutions enable small and medium-sized pharmaceutical, biotechnology, and clinical research organizations to manage clinical studies efficiently and effectively in order to reach evaluation and submission. Our understanding and experience as clinical researchers and technologists translates into patented innovation in digital trial processes that bring significant cost and time savings to our customers, while improving the quality of the trial.

History

Precherche was founded in 2005 with a history that dates back to the late 1990s as an early technology company automating the clinical trial lifecycle. The company is privately financed, with headquarters located in Austin, Texas, and staff in the Washington, D.C. area.

Markets Served

As the average pre-tax cost of clinical trials approaches \$1 billion, pharmaceutical companies struggle to bring drugs to market as quickly as possible within the timeframes and budgets allocated. Successful trials depend on factors that have little to do with the start and administration of the trial. These factors lead to delayed study starts, lost time, overextended budgets, patient data errors, and failed compliance, with no certainty that the trial will reach an evaluative conclusion.

Precherche delivers complete digital trial management service to clinical research organizations, biotechnology companies, and small and medium-sized pharmaceutical companies who need rapid clinical trial management that is visible in real-time, with the ability to foresee the full financial and time requirements of the trial before it ever begins. Precherche clients are in the unique position to plan for the time and cost of the entire life of the trial up front and to make immediate decisions on how to proceed effectively and successfully with their trial while the trial is in progress.

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Digital Trial Management

Precherche's services are available through the InterRx platform and are designed to scale quickly around studies developed for complete digital delivery:

InterRx™ is the first digital trial management solution, going beyond EDC to provide patient, document, and inventory management, data cleaning and handling, workflow scheduling, messaging, and querying between sponsors, monitors, and investigators to encompass all auditable aspects of a clinical trial. Using the Sure Start program, InterRx rapidly moves from approved protocol to study start in 5 business days.

InterRx™ CD combines InterRx with a streamlined study team to create an outsourced clinical research organization, available on a moment's notice. Using the Rapid Trial program, InterRx CD employs the best practices of an experienced team with the innovative technology of InterRx to start trials in 10 business days and to produce quality trial results within budget and on time.

Media Contacts

Art Adair

Innova Design & Advertising
artadair@hotmail.com
713-623-4432 ext. 109

Joi Chevalier

Precherche Life Sciences
jchevalier@precerche.com
512-233-0197