

Court Square provides professional and strategic consulting services in information technology, business processes and service management using ITIL and PMI best practices. We drive the business of science by providing life science subject matter expertise that spans the drug development cycle with a range of technology, quality and audit services.

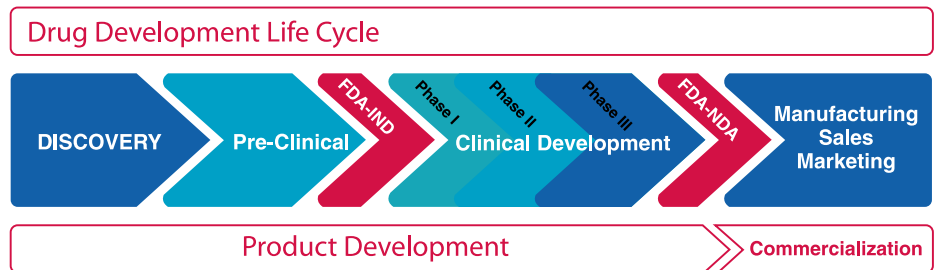


## GSP**bio**

Court Square has been providing disciplined managed services consulting to life sciences firms since 1995. GSP bio™ is specifically designed for new and emerging biotech/pharma companies as they grow through their development and clinical trials life cycle. GSP bio offers infrastructure managed services along with choices in application hosting and support for day to day infrastructure applications or those more specific to life sciences. All of this support is accomplished at Court Square's co-location facility on a qualified infrastructure. GSP bio drives the business of science.

### Multiple levels of GSP**bio** to grow as you grow.

Three levels of GSP bio are available to match the needs in data generation and validation requirements as companies get closer to FDA submission.



### **bio**Launch

Designed for the needs of very young companies with small numbers of employees and limited applications who are likely in the discovery phase. No pressure from the FDA at this point but there will be as this new company progresses toward an IND filing. In the interim, BioLaunch meets this company's needs in basic IT infrastructure and document management.

### **bio**Growth

Designed for companies in phase I to early phase II clinical trials. Life changes dramatically for the company who successfully files an investigational new drug application. The clinical trial phase generates significant amounts of data that is subject to regulations from a number of entities. There are requirements to work in a qualified environment with validated applications. 21CFR11 compliant document management becomes a major part of the strategy.

### **bio**Ready

The final stage for companies with candidates progressing successfully into phase II or III trials and who are preparing for FDA submission and commercialization. Often segments of this work are outsourced to Commercial Research Organizations (CRO) and Contract Manufacturers (CMO). Companies in this stage have IT requirements that increase in sophistication and broaden in information needs. BioReady is specifically for companies whose validated electronic data management needs extend into the areas of quality management, outsource collaboration and possibly adverse event and safety.

For greater details on all the services available in the GSPbio line, visit [www.gspbio.com](http://www.gspbio.com) or [www.courtsquaregroup.com](http://www.courtsquaregroup.com)