



## Access the hub – and let amazing things happen.

Your program begins with exclusive access to the hub – complementary innovations you can tap into at any stage of the drug development continuum.

With the hub, you have access to a portfolio of biotech-dedicated solutions to define your journey. Propel forward study by study or with a comprehensive programmatic approach for the ultimate in continuity and time savings. You set the plan and the pace.

### **MarketPlace**

Planning to capitalize on the value of your asset? Now you can gain critical visibility and establish the right connections early with investors and development partners. Finding the right partner may be only a click away via this unique, online networking solution.

### **Early Phase Development Solutions**

Need to move swiftly through IND/CTA and into the clinic? Now you can do it 30% faster by choosing a prospective plan that gives you unprecedented continuity. Your asset is surrounded with a cross-functional team of experts and proven efficient processes that save you time and money.

### **Phase I Sites and Adaptive Trial Designs**

Not sure where to start with your First-in-Human trial? Things just got easier with access to Phase I clinical sites and an experienced team. Gain earlier clinical insights with adaptive trial designs that accelerate timelines, enable decision making and improve success rates at this early stage.

### **Clinical Trials**

Seeking a more collaborative, flexible approach to your clinical trials? Look no further than the boutique trial approach of Chiltern, a Covance company. Gain insights from a devoted team of scientific, regulatory and therapeutic experts that specialize in a variety of indications, including oncology and rare and orphan diseases.

### **Central Laboratory Services**

Does your study call for streamlined safety and esoteric testing? Choose to work with a biotech-dedicated study management team. Leverage trial-specific specimen collection kits, unmatched logistics support and easy access to scientific expertise.

### **Biomarkers and Companion Diagnostics**

Want to reach patients who will benefit most from your novel drug? You can do it successfully through the power of precision medicine. From pathway selection and biomarker development to validation, you'll get an informed regulatory and commercial strategy for your companion diagnostic.

### **Advanced Informatics and Analytics**

Is finding rare patients a roadblock for your trial? Streamline trial design, speed investigator and patient recruitment and enhance your monitoring practices with a combination of advanced analytics and LabCorp's opt-in patient database. Find the right patients anywhere, no matter how rare the disease.

### **Regulatory and Commercial Consulting**

Need to advance your nonclinical and clinical strategies, navigate regulatory requirements or identify the right approach? Start with the end in mind and a solid Target Product Profile. Then, pave the way to commercial success with insights from early-stakeholder and market research.

Determining your approach is up to you. Planning for success is possible via the hub. Visit: [Covance.com/TheBioExperience](https://Covance.com/TheBioExperience)