



# go further

On-going investments in cell and gene therapy (CGT) fuel growth

As of 2020, 16 out of 20 of the largest biopharma manufacturers have added cell and gene therapies to their portfolios.<sup>1</sup>

 **2,700**

The number of CGT therapies in development as of April 2022

 **\$67.4B**

The expected worth of the global CGT manufacturing market by 2030<sup>2</sup>



# Realize the potential of your CGT

The road ahead is complex and uncertain. PPD is here to help you overcome challenges and get ahead of unknowns—advancing your novel therapy across every aspect of the development life cycle. From early strategic development to value and access consultation, we can partner with you to help identify and operationalize your path forward.

**> 130**

clinical trials in the last 5 years

**> 75**

gene therapy clinical trials

adenovirus/AAV • lentivirus • retrovirus •  
virus-based immunotherapies

**> 55**

cell therapy clinical trials

CAR-NK cells • adoptive T-Cell • antigen-  
presenting cells • primitive stem cells

**> 300**

peri- & post-approval projects

value demonstration • market access •  
commercial strategy

**> 20**

years of bioanalytical lab  
GCT experiences

millions of samples analyzed • 300+ CGT  
assays • over 170 scientists and staff

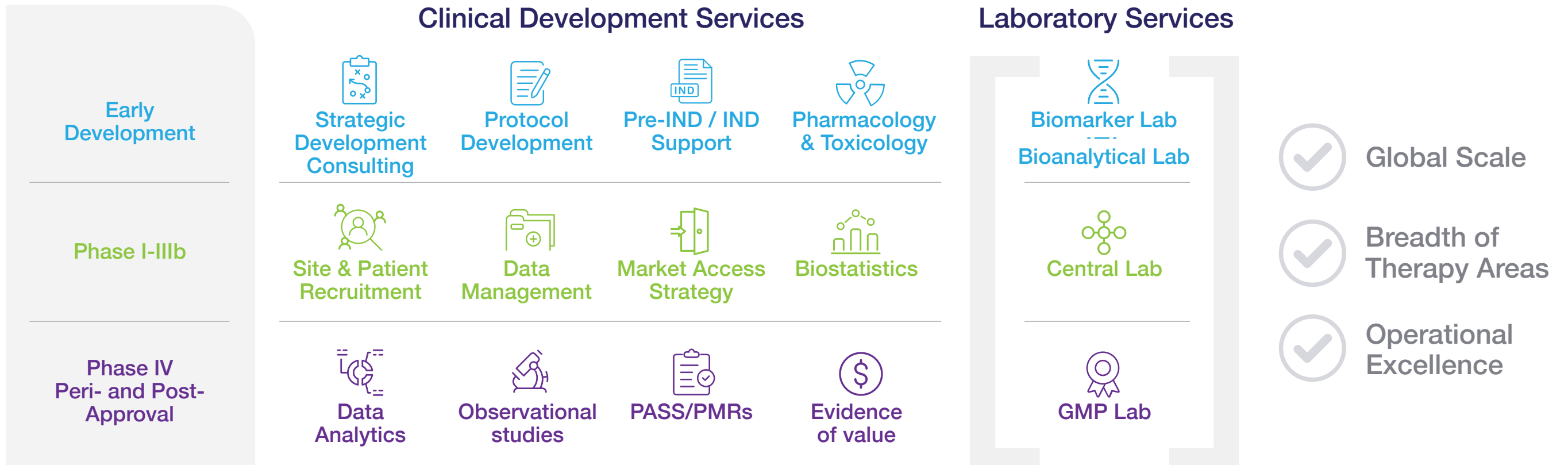
**Partner with the CRO that  
supported 100% of marketed  
transformative therapies.**

**Let's go →**



# Novel solutions for novel therapies

To optimize success, you need a CRO that truly gets CGT drug development and is ready with solutions at every phase of the journey. Our comprehensive end-to-end CGT solutions support the distinct needs of patients, sites, and sponsors with the ease of working with a single partner.





# Backed by an industry-leading CGT team

Working under constantly evolving regulatory guidance, accessing dispersed patient populations, and leading long-term follow-up studies and registries takes a team with equal parts skill and passion. With us, you have a partner with access to the experience and expertise to put your CGT on a clear path to changing lives.

**100+**

regulatory and market access consultants and strategists

**700+**

clinical and real-world data management staff globally

**2,900+**

clinical operations staff

**600+**

multidisciplinary peri- and post-approval scientists globally

**12+**

hours of formalized, comprehensive training for all CGT staff

Put our 20+ years of CGT experience to work for you.

**Let's go →**

