PPD®

Helping Our Customers Deliver Life-Changing Therapies

Bending the Cost and Time Curve of Drug Development and Optimizing Value for Our Customers

PPD accelerates our customers' medicines from early development through regulatory approval and market access. We are committed to global operational excellence and innovative solutions that deliver more patients to fewer sites, more patient-centric trials to recruit and retain patients faster, and generating evidence faster and at lower cost to increase return on investment.

Laboratories

- Bioanalytical
- Biomarkers
- Central Lab
- GMP Lab
- Vaccine Sciences

Early Development

- Chemistry, manufacturing, and control (CMC) consulting
- Manufacturing and controls
- Nonclinical development and chemistry
- Pharmacology and toxicology
- Phase I clinic (for healthy volunteers)
- Phase I patient network
- Translational medicine

Clinical Development

- Accelerated Enrollment Solutions
- Biostatistics
- Clinical supplies
- Clinical trial monitoring
- Data management
- Feasibility studies
- IVR
- Medical communications
- Medical writing
- Patient recruitment
- Pharmacokinetics and pharmacodynamics (PK, PD)
- Project management
- Quality and compliance
- Regulatory affairs
- Study startup

Real-World Research

- Real-world evidence (database analytics, registries and other observational studies)
- Interventional studies (IIIB/IV, EAP/CUP, XAP)
- Risk evaluation and mitigation strategies (REMS) and risk management plans
- Pragmatic/adaptive studies
- Patient-centered research
- Market access
- Modeling and meta research
- Epidemiology
- Health economics
- Global pharmacovigilance
- Medical communications
- Consumer health

Consulting

- Adaptive Trial Design
- Biosimilars
- Gap Analyses and Due Diligence
- Pediatrics
- Product Development
- Protocol & Trial Optimization
- Rare Diseases
- Regulatory Consulting
- Strategic Development Planning

Our end-to-end services expedite the development of our customer's life-changing therapies from early development through market access

Early Development

Expertise and infrastructure to deliver custom early phase strategies.

Phase I clinical services deployed globally through our extensive network of clinical pharmacology sites.

Clinical Development

Full-service clinical studies for multinational regulatory submissions.

Proven expertise in biostatistics and data management, clinical supplies, medical communications, regulatory affairs, pharmacovigilance, and site intelligence and activation.

Real-time data and analysis delivered by Preclarus®.

Real-World Research

Evidence of safety, value and effectiveness to optimize market access.

Innovative, comprehensive strategies to achieve regulatory approval and speed market access through Evidera.

PPD® Laboratories

Global lab services spanning large and small molecules, integrated across the spectrum of drug development.

Central lab, GMP lab, bioanalytical lab, vaccine sciences lab and biomarkers.

PPD® Consulting

Industry-leading experts with diverse expertise across therapeutic areas and functional disciplines.

Product development, biosimilars, adaptive trial design, pediatrics, rare disease, cardiovascular outcomes, medical devices, market access and more.

30+ YEARS
Helping Deliver Life-Changing Therapies

21,000+ employees

93 Offices

48 Countries

WORKED ON DEVELOPMENT PROGRAMS FOR

270+
biotech & small pharma companies

49 of the top **50**
pharmaceutical companies

INVOLVED IN 66 DRUG APPROVALS IN 2018:

21 By FDA
21 By EMA
24 By NMPA

+1 877 643 8773 | + 1 919 456 5600