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BVMA member since 2014
Audit passed in 2014

## HISTORY AND MISSION

From its beginnings in 1985 as a one-person consulting firm in Maryland to its position today as one of the world´s leading contract research organisations, PPD has offices in 46 countries and approximately 13,000 professionals worldwide. Four themes drive PPD´s success: engaged employees, medical and scientific expertise, quality execution and global reach. Our mission is to help our clients and partners in maximizing returns on their R&D investments.


## HIGHLIGHTS

- Development Services since 1985
- Worked with 46 out of the top 50 pharmaceutical companies (according to 2013 R&D spend) and more than 600 biotechnology companies
- 20+ years experience with government trials
- Experience managing studies in all medical device phases
- PPD ranked best for reputation, innovation in improving site relationships and use of technology in 2013 CenterWatch Global Site Survey
- PharmaTimes´ 2014 International Clinical Research Team of the year

## SERVICES

**Drug Discovery**
- Range of integrated drug discovery services, as well as access to innovative technology platforms

**Early Development**
- 25 + years of dedication to phase 1 studies, with strong reputation for innovative and complex studies
- 300-bed-healthy volunteer clinic in Austin, Texas, with 160 studies/year

**Clinical Development**
- Biostatistics, program development, randomization methods
- Global medical writing with literature reviews and integrated clinical reports
- Pharmacovigilance with 24/7 safety and medical management coverage
- Regulatory Affairs to address local, regional and global development needs
- Quality Assurance including audits, mock inspections and computer validation
- Project management to ensure efficiency and accuracy throughout projects
- Data Management to ensure consistent high-quality data
- Feasibility studies and comprehensive patient recruitment services
- Clinical Trial Monitoring, Source Data Verification in phase I, II, III and IV trials

**Laboratories**
- Central and Bioanalytical Lab with analysis of clinical samples for safety, efficacy, hematology, chemistry, biomarker and genomic testing, analysis of biological samples for content of drug, metabolite and biomarker. cGMP and Vaccines and Biologics Lab.

**Post Approval**
- Global pharmacovigilance including individual case safety reporting and medical review, global medical communications by contact center services, real-world outcomes

**Consulting**
- Physicians, scientists, regulatory professionals and biostatisticians with strong scientific and medical expertise across therapeutic areas and functional disciplines