Roy D. Simmons PhD, MBA, PMP
Consulting Project Manager

Summary. Drug Program management leadership since 1997 delivering superior business results through the application of Project and Portfolio Management best practices. Background includes diverse project management experience in Biopharma from pre-clinical through Phase IV and a unique combination of technical (PhD Chemistry) and business (MBA) expertise. Areas of project management experience and expertise include:

- Biologics, Small-Molecule, and Combination Product Development
- Regulatory Submissions,
- CMC Project Management
- Strong Computer Skills include MSProject, Excel, Visio, and Webex
- Co-Development Alliances and Out-Sourced Project Resources
- Geographically Diverse Cross-functional Teams
- Meeting Facilitation in Complex Environments
- Project Management Office (PMO) Implementation and Management
- Enterprise Project Management Software Implementation and Administration

Professional Experience

Simmons Management Services, LLC
Principal: 2012-present

Integrated Project Management Co., Inc.
Project Management Consultant: 2007 - 2012

Nektar Therapeutics
Associate Director – Program Management: 2004 - 2007

Crompton Corp.
P.A.C.E. Manager (NPD PMO): 1997 - 2003

Education/Certification

1978; BS, Chemistry, Clemson University
1981; MS, Chemistry, Clemson University
1987; PhD, Organic Chemistry, University of New Orleans
1993; MBA, University of Memphis
2006; PMP, Project Management Institute

Drug Development and Regulatory Experience

- Multiple early phase drug development projects for MAbs, polypeptides, and polynucleosides
- Phase I/II management for Novantik (naloxegol)
- Development and commercialization of anti-convulsant auto-injector device for the Dept. of Defense
- Managed clinical supply chain for Pfizer Oncology Grant Studies
- Managed Phase IV CMC activities and supply chain for PEG reagents for Pegasys and Neulasta