Jill C. Bossi
Senior VP Clinical Operations, Founder

jbossi@elitebpconsulting.com

Jill has over twenty-seven years of experience in drug development with a focus in Clinical Operations and Program Management. Jill has expertise in building and leading cross functional project teams throughout the entire clinical trial process, while meeting or exceeding corporate milestones. With a broad history of industry roles in almost all areas of clinical development, she has the extensive knowledge required to successfully execute both early and late phase clinical development programs. Jill has experience with both small emerging biotech and large pharma companies, in a wide variety of therapeutic areas including Oncology, Cell Therapy, Immuno-Oncology, Rare Disease, Ophthalmology, Endocrine/Metabolic, Gastrointestinal, Cardiology, CNS, Analgesia, Dermatology, Infertility, Surgical Device, Dental and Transplant trials.

Jill and her business partners founded Elite BioPharma Consulting in 2019. At Elite, Jill partners with clients to help drive excellence in their clinical development process from IND application through NDA filing. She provides superior Clinical Development and Management Services, offering professional expertise tailored to the client's needs based on the current life cycle of their program. For small biotech companies, she provides operational leadership and essential infrastructure building in the early phases of clinical development, allowing clients the opportunity to run their trials efficiently and in accordance with GCP/ICH guidelines.

Most recently, Jill has served in various consulting roles as Head of Clinical Operations for small emerging biotech companies. Prior to that she was Director of Clinical Operations for Checkmate Pharmaceuticals in Cambridge, MA, spearheading their Phase 1 melanoma program. She provided insight and guidance regarding the necessary infrastructure and continued to provide operational leadership while successfully moving the programs forward in alignment with corporate objectives. Prior to this, she was a successful clinical operations consultant for over eighteen years. During this time, she helped to launch eight novel clinical development programs and collaborated with dozens of clients to fulfill various roles including head of clinical operations, program manager, clinical trial manager, QA auditor, and medical writer. In her career, Jill has been involved in over 40+ trials ranging from single-center PK/PD to global pivotal Phase 3 trials conducted in 11 countries. She also spent a total of three years working for PAREXEL International Corporation first as a clinical research associate and then as a consultant.

Jill earned a Bachelor of Science in Biology, with a minor in Psychology from the University of Vermont. During her time in college, she was also able to participate in a clinical study on phantom limb pain (PLP) and cortical regeneration in amputees. This is what created her passion for clinical research. She resides on the New Hampshire seacoast with her family where she enjoys time in the mountains and at the shore.