Bao Truong, LLC
Bao Truong, B.S.

Bao Truong is a Regulatory Strategic Consultant based in San Diego, CA. She has over 17 years of regulatory experience in oncology product development, in both global pharmaceutical and start-up environment. History of success in developing initial regulatory plan for complex/novel clinical development strategy of 6 NMEs, from Phase 1/proof-of-concept through registration-enabling phase; 5 NMEs subsequently becoming approved products. Bao has broad leadership with extensive experience in leading cross-functional teams.

Prior to being an independent consultant, Bao served as Vice President of Regulatory Affairs at Lassen Therapeutics, Erasca, and IDEAYA Biosciences. Prior to that, she served as Head of Regulatory Affairs at Ignyta, where she helped drive Rozlytrek®'s initial development plan from Phase 1 to registration-enabling Phase 2 supporting both tissue-specific and tissue-agnostic indications; head of regulatory affairs at Seragon Pharmaceuticals and Aragon Pharmaceuticals, helped advance Erleada®'s initial development plan from Phase 1 to initiation of global Phase 3, supporting the first FDA-approved therapy for non-metastatic CRPC using a novel clinical endpoint; and held regulatory lead positions of increasing responsibility at Genentech for multiple oncology products, including Erivedge® (from its Phase 1 to registration-enabling Phase 2 using a composite endpoint, enabling the first FDA-approved therapy for advanced basal cell carcinoma), Kadcyla®, Gazyva®, Tarceva® and Herceptin®. Earlier in her career, Bao worked in oncology research at University of California, San Diego and University of California, San Francisco. Bao earned her Bachelor of Science in Biochemistry, from the University of California, in San Diego.