

Board of Directors Bio



Shamim Ruff, SVP, Head of Regulatory Affairs and Quality Assurance, Stoke Therapeutics

Shamim Ruff is the Senior Vice President, Head of Regulatory Affairs and Quality Assurance, at Stoke Therapeutics. She has extensive knowledge of drug development with more than 25 years in the biopharmaceutical industry, working with a diverse range of therapeutics. She has expertise in both domestic and international regulatory affairs spanning rare diseases, oncology, hematology and antivirals.

Prior to joining Stoke, Shamim was the Chief Regulatory Affairs Officer and SVP of Quality at Sarepta Therapeutics. She was previously Vice President, Head of Oncology, Global Regulatory Affairs at Sanofi- Genzyme where she was responsible for leading the Global, European and CMC Regulatory Affairs teams. Shamim has also held senior positions at Amgen Inc., Abbott Laboratories Inc., and AstraZeneca PLC, where she oversaw the development and filings of multiple successful regulatory approvals across the world. Shamim holds a B.Sc. in Chemistry and Biology and an MSc. in Analytical Chemistry.