Heather Knight, Pharm.D.
Vice President, Regulatory Strategy

Dr. Heather Knight is a Vice President of Regulatory Strategy at Advyzom. Heather provides strategic regulatory guidance and acts as a US Agent to FDA for domestic and international biopharmaceutical companies. Heather's areas of expertise include oncology/hematology, virology, immunology, diabetes, CNS, obesity, urology, radiation medicine and pulmonary drugs, biologics, generics, imaging and medical countermeasures. Heather has hands-on experience in successfully obtaining BLA, NDA and sNDA approvals. Heather manages multiple complex projects simultaneously, including FDA meetings, IND submissions, amendments, responses to FDA requests, advisory committee preparation and critical issue management. She brings strong expertise to clients through her role as US Agent and Global Regulatory Leader, providing strategic advice to small and large clients.

Heather has over 20 years of experience in the pharmaceutical industry. Prior to joining Advyzom, Heather was the former Senior Vice President and Head of Regulatory Affairs at Aerium Therapeutics leading health authority interactions on monoclonal antibodies for COVID-19. Prior to Aerium, she was Vice President of Regulatory Affairs and Quality Assurance at Chimerix Inc., where she led the submission and approval of TEMBEXA™ for the treatment of smallpox and the clearance of an IND for Acute Lung Injury in patients with COVID-19. Earlier in her career, Heather was at Bristol Myers Squibb where she held roles in Global, U.S. and Europe. She led the global and US approvals of YERVOY, the first immuno-oncology agent. Heather also was an Associate Director in Drug Regulatory Affairs at Hoffmann-La Roche, Inc., wherein her work included antiviral and oncology drugs.

Heather earned her Doctor of Pharmacy and a Bachelor of Arts in Biology from West Virginia University. She is a Registered Pharmacist in West Virginia.