FOR IMMEDIATE RELEASE

HURLEY CONSULTING ASSOCIATES LTD. ANNOUNCES APPOINTMENT OF ALBERT T. LEUNG, MD, PhD AS VICE PRESIDENT, DEVELOPMENT

Summit, New Jersey (August 16, 2016) – Hurley Consulting Associates Ltd., a leading international drug development consulting company recognized for its high level of regulatory expertise, announces the appointment of Albert T. Leung, MD, PhD as Vice President of Development. Dr. Leung is responsible for all drug development consulting projects and management of clinical programs. These responsibilities include strategy development, design and oversight of clinical programs, and clinical documentation preparation to support investigational drug applications, marketing applications and health authority meetings.

Dr. Margaret E. Hurley, President and CEO of Hurley Consulting Associates, commented: "Dr. Leung's depth of knowledge with over eighteen years' experience in all phases (Phase 1 to IV) of drug development in multiple therapeutic areas including osteoporosis and metabolic bone diseases, diabetes, woman's health, rare disease and drug delivery systems truly add a great addition to our organization."

Dr. Leung has extensive expertise in the strategy and tactics of drug development, clinical trial design and operations, data collection, pharmacovigilance, regulatory dossier authoring and health authority responses with extensive interactions with worldwide regulatory agencies (FDA, EMA, PMDA) in early and late phase and post marketing drug development, including successful submission and approval of WMA.

Before joining Hurley Consulting, Dr. Leung was Chief Medical Officer at GlySure, Ltd., a U.K based company where he provided leadership and mentorship to the U.S. Clinical Development of intravascular continuous blood glucose monitoring system for critical care patients. As CMO, he built and managed clinical research operation group in the U.S., interfacing with scientific and clinical leaders at the FDA.

Dr. Leung holds a bachelor's degree in chemistry from Washington University, St. Louis, MO, a doctor of philosophy in Physiology and Biophysics and a medical degree from The University of Iowa. He is Board Certified in Internal Medicine and Endocrinology and Metabolism.

"I am very pleased to join the team at Hurley Consulting, a talented group with extensive leadership and success," commented Dr. Leung. "I look forward to adding my clinical and drug development experience to the clinical team at Hurley Consulting."

Based in Summit, NJ, Hurley Consulting Associates Ltd. was founded in 1987. The company specializes in finding solutions for its clients' regulatory and commercial development needs. With unique expertise to prepare global regulatory submission documents and dossiers, Hurley Consulting integrates nonclinical, clinical, and manufacturing and control evaluations, performs data analyses, and develops and implements regulatory strategies. With a proven track record of over twenty-five years' regulatory experience, Hurley Consulting can serve its clients as U.S. agent and authorized representative to FDA for regulated products for the entire IND through NDA process.

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