ArQule to Present at the 8th Annual Leerink Partners Global Healthcare Conference on February 28, 2019

February 21, 2019

BURLINGTON, Mass.--(BUSINESS WIRE)--Feb. 21, 2019-- ArQule, Inc. (Nasdaq: ARQL) today announced that Paolo Pucci, Chief Executive Officer, and Brian Schwartz, M.D., Chief Medical Officer and Head of Research and Development, of ArQule will present at the 8th Annual Leerink Partners Global Healthcare Conference on February 28, 2019, at 11:30 a.m. ET at the Lotte New York Palace in New York City.

The live webcast of the presentation will be available via the “Investors & Media” section of ArQule’s website, www.arqule.com, under “Events & Presentations.” A replay of the webcast will be available shortly after the conclusion of the presentation.

About ArQule
ArQule is a biopharmaceutical company engaged in the research and development of targeted therapeutics to treat cancers and rare diseases. ArQule’s mission is to discover, develop and commercialize novel small molecule drugs in areas of high unmet need that will dramatically extend and improve the lives of our patients. Our clinical-stage pipeline consists of five drug candidates, all of which are in targeted, biomarker-defined patient populations, making ArQule a leader among companies our size in precision medicine. ArQule’s pipeline includes: ARQ 531, an orally bioavailable, potent and reversible inhibitor of both wild type and C481S-mutant BTK, in Phase 1 for patients with B-cell malignancies refractory to other therapeutic options; Miransertib (ARQ 092), a selective inhibitor of the AKT serine/threonine kinase, in a Phase 1/2 company-sponsored study for Overgrowth Diseases, in a Phase 1 study for ultra-rare Proteus syndrome conducted by the National Institutes of Health (NIH), and in Phase 1b in combination with the hormonal therapy, anastrozole, in patients with advanced endometrial cancer; ARQ 751, a next generation AKT inhibitor, in Phase 1 for patients with AKT1 and PI3K mutations; Derazantinib, a multi-kinase inhibitor designed to preferentially inhibit the fibroblast growth factor receptor (FGFR) family, in a registrational trial for iCCA; and ARQ 761, a β-lapachone analog being evaluated as a promoter of NQO1-mediated programmed cancer cell necrosis, in Phase 1/2 in multiple oncology indications in partnership with the University of Texas Southwestern Medical Center. ArQule’s current discovery efforts are focused on the identification and development of novel kinase inhibitors, leveraging the Company’s proprietary library of compounds.

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