Passage Bio to Present at Barclays Gene Editing & Gene Therapy Summit

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PHILADELPHIA, Nov. 06, 2019 (GLOBE NEWSWIRE) -- Passage Bio, a genetic medicines company developing AAV-delivered gene therapies for the treatment of rare monogenic central nervous system diseases, today announced that Jill Quigley, chief operating officer, will present at the Barclays Gene Editing & Gene Therapy Summit on Wednesday, November 13th, 2019 at 9:30 a.m. ET in New York, NY.

About Passage Bio
Passage Bio is a privately-held fully integrated genetic medicines company with a mission to develop a portfolio of life-transforming AAV-delivered therapeutics for the treatment of rare monogenic central nervous system diseases. The company is based in Philadelphia, PA and has a research, collaboration and license agreement with the University of Pennsylvania (Penn) and its Gene Therapy Program (GTP), as well as the Orphan Disease Center at Penn. The GTP conducts IND-enabling preclinical work and Passage Bio conducts all clinical development, regulatory strategy and commercialization activities under the agreement. The company has a development portfolio of six product candidates, with the option to license six more, with lead programs in GM1 gangliosidosis, frontotemporal dementia and Krabbe disease. Since inception, Passage Bio has raised $225.5 million with investments from OrbiMed, Versant Ventures, Frazier Healthcare Partners, Access Biotechnology, Lily Asia Ventures, New Leaf Venture Partners, Vivo Capital, and Boxer Capital of Tavistock Group, among others.

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