Agilis Biotherapeutics Presents at Sachs Associates 15th Annual Biotech in Europe Forum

Cambridge, MA, September 30, 2015 -- Agilis Biotherapeutics, LLC (Agilis), a biotechnology company advancing innovative DNA therapeutics for rare genetic diseases that affect the central nervous system (CNS), announced today that Mark Pykett, President and CEO of Agilis, presented at the Sachs Associates' 15th Annual Biotech in Europe Forum and participated in the panel session "Orphan & Rare Diseases" in Basel, Switzerland. Dr. Pykett's presentation provided an update on Agilis' development progress in its gene therapy programs for Friedreich's Ataxia and Angelman Syndrome, and recent scientific advances in the use Reelin supplementation in disorders of cognitive impairment. Dr. Pykett's panel session covered an overview of treatment approaches utilizing enzyme replacement therapy and gene therapy, as well as other novel therapeutic interventions for rare diseases. Additional topics included investors' perspectives on rare diseases, entrepreneurial best practices in the space, and financing and partnering for businesses focused on rare diseases. Dr. Pykett discussed Agilis' gene therapy efforts in rare diseases and the potential for innovative gene therapy technologies to provide new treatment paradigms that address the significant unmet need often found in orphan-category diseases, particularly in rare disorders of the central nervous system.

The Sachs Associates' Annual Biotech in Europe Forum is recognized as a leading international stage for investing and partnering in the biotech and life science industry. The Forum draws a cross-section of early-stage/pre-IPO, late-stage and public companies and leading investors, analysts, money managers and licensing executives. The conference hosted over 600 delegates and 100 presenting companies, facilitated two days of one-to-one meetings, and featured twelve plenary panels covering a range of therapeutic areas.

About Agilis Biotherapeutics, LLC

Agilis is advancing innovative DNA therapeutics designed to provide long-term efficacy for patients with debilitating, often fatal, rare genetic diseases that affect the central nervous system. Our therapies are engineered to impart sustainable clinical benefits, and potentially a functional cure, by inducing persistent expression of a therapeutic gene. The Company's technology is aimed at the precise targeting and restoration of lost gene function, while avoiding unintended off-target effects. Our integrated strategy increases the efficiency of developing DNA therapeutics into safe, targeted gene therapies that achieve long-term efficacy and enable patients to remain asymptomatic without continuous invasive treatment. Agilis' rare disease programs are focused on gene therapy for Friedreich's Ataxia, Angelman Syndrome and Fragile X Syndrome, rare genetic diseases that include severe neurological deficits and result in physically debilitating conditions. Friedreich's Ataxia is the most common hereditary ataxia with an estimated 5,000 to 10,000 patients in the U.S. There are an estimated 10,000 to 15,000 people living with Angelman Syndrome in the US. Fragile X Syndrome is the most common known cause of inherited intellectual disability with an estimated 64,000 patients living in the U.S.

We invite you to visit our website at www.agilisbio.com

Safe Harbor Statement

Some of the statements made in this press release are forward-looking statements. These forward-looking statements are based upon our current expectations and projections about future events and generally relate to our plans, objectives and expectations for the development of our business. Although management believes that the plans and objectives reflected in or suggested by these forward-looking statements are reasonable, all forward-looking statements involve risks and uncertainties and actual future results may be materially different from the plans, objectives and expectations expressed in this press release.

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