March 14, 2018

Abstract on AG200-15 (Twirla®) Accepted for Presentation at the APhA 2018 Annual Meeting & Exposition

PRINCETON, N.J., March 14, 2018 (GLOBE NEWSWIRE) -- Agile Therapeutics, Inc., (Nasdaq:AGRX), a women's healthcare company today announced that an abstract related to the Phase 3 SECURE study for AG200-15 (Twirla®), an investigational, once-weekly, low-dose hormonal contraceptive patch, has been accepted as a poster presentation at the upcoming APhA 2018 (American Pharmacists Association) Annual Meeting & Exposition, which will be held from March 16 — 19, 2018 in Nashville, TN.

The poster will include a summary of efficacy and safety along with the results of the bleeding and wearability/tolerability profile from the SECURE study as previously reported.

The presentation details are as follows:

Poster Title: Results from the SECURE trial, a Phase 3 Study of the AG200-15 Investigational Transdermal Contraceptive Patch
Poster ID: 231
Room: Exhibit Hall C-D, Music City Center
Date and Time: Saturday, March 17, 9:00am-6:00pm and Sunday, March 18, 9:00am-3:00pm

The Phase 3 SECURE study was a multicenter, single-arm, open-label, 13 cycle trial designed to evaluate the efficacy, safety and tolerability of AG200-15, also known as Twirla, in 2032 healthy women, aged 18 years and over, at 102 investigational sites across the United States. The SECURE study design included a number of stringent elements, including exclusion of treatment cycles for use of back-up contraception and for lack of sexual activity. The study also had broad entry criteria, placed no limitations on BMI or other demographic factors during enrollment, and enrolled a large and diverse patient population in order to allow efficacy to be assessed across different, real-world groups, as requested by the FDA. These entry criteria resulted in the inclusion of a substantial number of women with a high BMI, who have frequently been underrepresented in past contraceptive studies.

About Agile Therapeutics, Inc.
Agile Therapeutics is a forward-thinking women's healthcare company dedicated to fulfilling the unmet health needs of today's women. Our product candidates are designed to provide women with contraceptive options that offer freedom from taking a daily pill, without committing to a longer-acting method. Our lead product candidate, Twirla® (levonorgestrel/ethinyl estradiol transdermal system) or AG200-15, is a non-daily, prescription contraceptive. Twirla is based on our proprietary transdermal patch technology, called Skinfusion®, which is designed to allow drug delivery through the skin while optimizing patch adhesion and comfort for the patient.

For more information, please visit the company website at www.agiletherapeutics.com. The Company may occasionally disseminate material, nonpublic information on the Company's website. Follow Agile on LinkedIn and Twitter**:** @AgileTher.

Forward-Looking Statement
Certain information contained in this press release includes "forward-looking statements" related to the Company's regulatory submissions. We may, in some cases use terms such as "predicts," "believes," "potential," "continue," "anticipates," "designed," "estimates," "expects," "plans," "intends," "may," "could," "might," "will," "should" or other words that convey uncertainty of the future events or outcomes to identify these forward-looking statements. Our forward-looking statements are based on current beliefs and expectations of our management team that involves risks, potential changes in circumstances, assumptions, and uncertainties. Any or all of the forward-looking statements may turn out to be wrong or be affected by inaccurate assumptions we might make or by known or unknown risks and uncertainties. Our statements about the results of our clinical trial could be affected by the potential that there are changes in the interpretation of the data by the FDA (for example, the FDA continues to question the number of pregnancies included in our results and they may adjudicate additional pregnancies); our statements about the potential commercial opportunity could be affected by potential labeling restrictions, the potential that our product does not receive regulatory approval, does not receive reimbursement by third party payors, or a commercial market for the product does not develop because of any of the risks inherent in the commercialization of contraceptive products. For all these reasons, actual results and developments could

be materially different from those expressed in or implied by our forward-looking statements. All forward-looking statements are subject to risks detailed in our filings with the U.S. Securities and Exchange Commission, including the Company's Annual Report on Form 10-K and our Quarterly Reports on Form 10-Q. You are cautioned not to place undue reliance on these forward-looking statements, which are made only as of the date of this press release. We undertake no obligation to publicly update such forward-looking statements to reflect subsequent events or circumstances.

SOURCE: Agile Therapeutics, Inc.

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