

Achieve Life Sciences Announces Granting of New Hire Inducement Awards

SEATTLE and VANCOUVER, British Columbia, Oct. 02, 2025 (GLOBE NEWSWIRE) -- Achieve Life Sciences, Inc. (Nasdaq: ACHV), a late-stage specialty pharmaceutical company focused on the global development and commercialization of cytisinicline for treatment of nicotine dependence for smoking cessation, today announced that effective October 1, 2025, Achieve's board of directors approved stock option grants to purchase an aggregate of 161,000 shares of its common stock for five new employees under Achieve's 2024 Equity Inducement Plan.

The stock options have an exercise price equal to the closing price of Achieve's common stock on October 1, 2025. Each stock option has a ten-year term and will vest as to 25% of the shares underlying the stock option on the first anniversary following commencement of employment, and the remaining 75% of the shares underlying the stock option will vest in 36 equal monthly installments thereafter, subject to the employee's continued service through each applicable vesting date.

The stock options are subject to the terms and conditions of the 2024 Equity Inducement Plan, as well as the terms and conditions of the stock option agreement covering the grant and were made as an inducement material to the individual entering into employment with Achieve in accordance with Nasdaq Listing Rule 5635(c)(4).

About Achieve Life Sciences, Inc.

Achieve Life Sciences is a late-stage specialty pharmaceutical company committed to addressing the global smoking health and nicotine dependence epidemic through the development and commercialization of cytisinicline. In September 2025, the company announced that its New Drug Application, submitted to the U.S. Food and Drug Administration (FDA) in June 2025, had been accepted for review. The FDA has assigned a Prescription Drug User Fee Act (PDUFA) date of June 20, 2026. The NDA is for cytisinicline to be used as a treatment of nicotine dependence for smoking cessation in adults, based on two successfully completed Phase 3 studies and its open-label safety study. Additionally, the company has completed a Phase 2 study with cytisinicline in vaping cessation and conducted a successful end-of-Phase 2 meeting with the FDA for a future vaping indication.

About Cytisinicline

There are approximately 29 million adults in the United States who smoke combustible cigarettes. Tobacco use is currently the leading cause of preventable death that is responsible for more than eight million deaths worldwide and nearly half a million deaths in the United States annually. More than 87% of lung cancer deaths, 61% of all pulmonary disease deaths, and 32% of all deaths from coronary heart disease are attributable to smoking and exposure to secondhand smoke.

In addition, there are approximately 17 million adults in the United States who use ecigarettes, also known as vaping. In 2024, approximately 1.6 million middle and high school students in the United States reported using e-cigarettes. There are no FDA-approved treatments indicated specifically as an aid to nicotine e-cigarette cessation. Cytisinicline has been granted Breakthrough Therapy designation by the FDA to address this critical unmet need.

Cytisinicline is a plant-derived alkaloid with a high binding affinity to the nicotinic acetylcholine receptor. It is believed to aid in treating nicotine dependence for smoking cessation and e-cigarette cessation by interacting with nicotine receptors in the brain, reducing the severity of nicotine craving symptoms, and reducing the reward and satisfaction associated with nicotine products. Cytisinicline is a new investigational product candidate being developed for the treatment of nicotine dependence for smoking cessation and has not been approved by the Food and Drug Administration for any indication in the United States.

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Source: Achieve Life Sciences