

## Taysha Gene Therapies Provides Update on Deprioritized Pipeline Programs

DALLAS, Feb. 15, 2024 (GLOBE NEWSWIRE) -- Taysha Gene Therapies, Inc. (Nasdaq: TSHA) ("Taysha" or "the Company"), a clinical-stage gene therapy company focused on developing and commercializing AAV-based gene therapies for the treatment of monogenic diseases of the central nervous system (CNS), today provided an update on its deprioritized pipeline programs as part of an ongoing effort to help support their further potential development.

Taysha has been working to find ways to advance its deprioritized programs. On November 13, 2023, Taysha terminated its existing loan and security agreement and entered into a new loan and security agreement that provides consent to allow the Company to transfer intellectual property (IP) for several deprioritized programs to third parties in a more efficient manner. The Company's new loan and security agreement also extended its cash runway into 2026.

Recent progress on previously deprioritized pipeline programs includes:

- o TSHA-120: The Company initiated the transfer of the United States Food and Drug Administration Investigational New Drug (IND) application and investigational clinical trial material for TSHA-120 in giant axonal neuropathy (GAN) to clinical trial collaborator National Institute of Neurological Disorders and Stroke (NINDS), creating an opportunity for continued clinical evaluation of TSHA-120 in GAN. Additionally, the Company entered discussions with the originating advocacy organization regarding TSHA-120 in an effort to transfer rights back to the advocacy organization to move the program forward.
- TSHA-101: The Company transferred rights back to Queen's University (Queen's) for TSHA-101 in GM2 gangliosidosis, resulting in Queen's regaining exclusive IP to the program.
- TSHA-104 and TSHA-112: The Company transferred rights back to the originating institution for select programs, including TSHA-104 in SURF1-associated Leigh syndrome and TSHA-112 in APBD.
- TSHA-118: The Company provided investigational clinical trial material for TSHA-118 in CLN1 to support an individualpatient investigator-initiated IND request from RUSH University Medical Center for the treatment of a patient with CLN1 disease.

"Today's announcement demonstrates meaningful progress to advance important development work for several deprioritized programs. Creating options for these programs has been a focus since the Company completed a management change in December 2022, and the new loan and security agreement afforded the flexibility to implement certain opportunities," said Sean P. Nolan, Chairman and Chief Executive Officer of Taysha. "As we continue to focus on advancing our lead TSHA-102 program for the treatment of Rett syndrome, we are pleased that we can ensure these programs are provided to the right advocates, clinicians and scientific experts who can potentially move these programs forward for the benefit of patients."

The Company continues to explore potential partnerships and opportunities for its other deprioritized programs to help support their further potential development.

#### **About Taysha Gene Therapies**

Taysha Gene Therapies (Nasdaq: TSHA) is a clinical-stage biotechnology company focused on advancing adeno-associated virus (AAV)-based gene therapies for severe monogenic diseases of the central nervous system. Its lead clinical program TSHA-102 is in development for Rett syndrome, a rare neurodevelopmental disorder with no approved disease-modifying therapies that address the genetic root cause of the disease. With a singular focus on developing transformative medicines, Taysha aims to address severe unmet medical needs and dramatically improve the lives of patients and their caregivers. The Company's management team has proven experience in gene therapy development and commercialization. Taysha leverages this experience, its manufacturing process and a clinically and commercially proven AAV9 capsid in an effort to rapidly translate treatments from bench to bedside. For more information, please visit <a href="https://www.tayshagtx.com">www.tayshagtx.com</a>.

#### Forward-Looking Statements

This press release contains forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995. Words such as "anticipates," "believes," "expects," "intends," "projects," "plans," and "future" or similar expressions are intended to identify forward-looking statements. Forward-looking statements concerning Taysha's cash runway and potential partnerships and opportunities for Taysha's deprioritized programs. Forward-looking statements are based on management's current expectations and are subject to various risks and uncertainties that could cause actual results to differ materially and adversely from those expressed or implied by such forward-looking statements. Accordingly, these forward-looking statements do not constitute guarantees of future performance, and you are cautioned not to place undue reliance on these forward-looking statements. Risks regarding our business are described in detail in our Securities and Exchange Commission ("SEC") filings, including in our Annual Report on Form 10-K for the full-year ended December 31, 2022, and our Quarterly Report on Form 10-Q for the quarter ended September 30, 2023, both of which are available on the SEC's website at <a href="https://www.sec.gov">www.sec.gov</a>. Additional information will be made available in other filings that we make from time to time with the SEC. These forward-looking statements speak only as of the date hereof, and we disclaim any obligation to update these statements except as may be required by law.

### **Company Contact:**

Hayleigh Collins
Director, Head of Corporate Communications and Investor Relations
Taysha Gene Therapies, Inc.
<a href="mailto:hcollins@tayshagtx.com">hcollins@tayshagtx.com</a>

# Media Contact:

Carolyn Hawley
Inizio Evoke
Carolyn.hawley@inizioevoke.com



Source: Taysha Gene Therapies, Inc.