4DMT Announces Presentations at 43rd Annual Scientific Meeting of the American Society of Retina Specialists

July 24, 2025

EMERYVILLE, Calif., July 24, 2025 (GLOBE NEWSWIRE) -- 4D Molecular Therapeutics (Nasdaq: FDMT, 4DMT or the Company), a leading late-stage biotechnology company advancing durable and disease-targeted therapeutics with potential to transform treatment paradigms and provide unprecedented benefits to patients, today announced podium presentations at the 43rd Annual Scientific Meeting of the American Society of Retina Specialists (ASRS) being held in Long Beach, CA, from July 30 – August 2, 2025.

ASRS 2025 Presentation Details:

Title:	Interim Results from the SPECTRA Phase 2a Clinical Trial Evaluating Intravitreal 4D-150 in Adults with Diabetic Macular Edema*
Date/Time:	Thursday, July 31, 2025 (11:35 – 11:38 a.m. PT)
Presenter:	David Almeida, MD, MBA, PhD, DABO, FRCSC, FASRS, Erie Retina Research, PA

*Includes 52-week primary endpoint analysis and 60-week analysis (all patients have reached 60 weeks as of the data cutoff date of May 2, 2025)

Title:	PRISM Phase 2b Clinical Trial Evaluating Intravitreal 4D-150 in Adults with Neovascular Age-related Macular Degeneration: 52-week Results
Date/Time:	Friday, August 1, 2025 (11:24 – 11:27 a.m. PT)
Presenter:	John A. Wells, MD, FACS, Palmetto Retina Center, SC

The presentations will also be available on the 4DMT website: https://4dmoleculartherapeutics.com/pipeline/#posters-and-publications

About 4DMT

4DMT is a leading late-stage biotechnology company advancing durable and disease-targeted therapeutics with potential to transform treatment paradigms and provide unprecedented benefits to patients. The Company's lead product candidate 4D-150 is designed to be a backbone therapy forming the foundation of treatment of blinding retinal vascular diseases by providing multi-year sustained delivery of anti-VEGF (aflibercept and anti-VEGF-C) with a single, safe, intravitreal injection, which substantially reduces the treatment burden associated with current bolus injections. The Company's lead indication for 4D-150 is wet age-related macular degeneration, which is currently in Phase 3 development, and second indication is diabetic macular edema. The Company's second product candidate is 4D-710, which is the first known genetic medicine to demonstrate successful delivery and expression of the CFTR transgene in the lungs of people with cystic fibrosis after aerosol delivery. 4D Molecular Therapeutics™, 4DMT™ Therapeutic Vector Evolution™, and the 4DMT logo are trademarks of 4DMT.

All of the Company's product candidates are in clinical or preclinical development and have not yet been approved for marketing by the U.S. Food and Drug Administration or any other regulatory authority. No representation is made as to the safety or effectiveness of the Company's product candidates for the therapeutic uses for which they are being studied.

Learn more at www.4DMT.com and follow us on LinkedIn.

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