



## **Kymera Therapeutics Appoints Karen Weisbach as Vice President, People and Culture and Jolly Bhatia as Vice President, Quality**

July 29, 2021

WATERTOWN, Mass., July 29, 2021 (GLOBE NEWSWIRE) -- Kymera Therapeutics, Inc. (NASDAQ: KYMR), a clinical-stage biopharmaceutical company advancing targeted protein degradation to deliver novel small molecule protein degrader medicines, today announced the appointments of Karen Weisbach, as Vice President, People and Culture, and Jolly Bhatia as Vice President, Quality.

"We are excited to welcome Karen and Jolly in these important roles as we continue to build the capabilities to establish Kymera as a leading, fully integrated degrader medicines company," said Nello Mainolfi, PhD, Co-Founder, President and CEO, Kymera Therapeutics. "Karen's experience in driving organizational and talent strategies in dynamic R&D organizations will be instrumental in supporting our continued growth, while Jolly's considerable experience in quality assurance for clinical and commercial-stage products will be invaluable as we continue to advance and broaden Kymera's pipeline of novel protein degraders."

### **Karen Weisbach, Vice President, People and Culture**

Ms. Weisbach joins Kymera with over 15 years of experience in human resources and operations, designing values-based initiatives to enhance company culture, improve employee experience and manage change in dynamic, innovative organizations in the biopharmaceutical industry. Prior to Kymera, she served in human resource leadership roles at bluebird bio, Inc., including building and leading the global People Partner function responsible for driving talent and organizational strategies with an emphasis on employee engagement, development, and performance.

### **Jolly Bhatia, Vice President, Quality**

Mr. Bhatia joins Kymera with approximately 25 years in quality oversight of manufacturing and clinical trial lifecycle activities including process and product development, clinical and commercial supply chains and new product launch. Prior to Kymera, he served in quality leadership roles at X4 Pharmaceuticals, Inc., Alnylam Pharmaceuticals, Inc., and Genzyme Corporation. He holds a Bachelor of Pharmacy degree from Birla Institute of Technology and Science and a Master's degree in Industrial Pharmacy from St. John's University.

### **About Kymera Therapeutics**

Kymera Therapeutics (Nasdaq: KYMR) is a clinical-stage biopharmaceutical company founded with the mission to discover, develop, and commercialize transformative therapies while leading the evolution of targeted protein degradation, a transformative new approach to address previously intractable disease targets. Kymera's Pegasus™ platform enables the discovery of novel small molecule degraders designed to harness the body's natural protein recycling machinery to degrade disease-causing proteins, with a focus on undrugged nodes in validated pathways currently inaccessible with conventional therapeutics. Kymera's initial programs are IRAK4, IRAK1MiD, and STAT3, each of which addresses high impact targets within the IL-1R/TLR or JAK/STAT pathways, providing the opportunity to treat a broad range of immune-inflammatory diseases, hematologic malignancies, and solid tumors. Kymera's goal is to be a fully integrated biopharmaceutical company at the forefront of this new class of protein degrader medicines, with a pipeline of novel degrader medicines targeting disease-causing proteins that were previously intractable.

Founded in 2016, Kymera is headquartered in Watertown, Mass. Kymera has been named a "Fierce 15" biotechnology company by FierceBiotech and has been recognized by the Boston Business Journal as one of Boston's "Best Places to Work." For more information about our people, science, and pipeline, please visit [www.kymeratx.com](http://www.kymeratx.com) or follow us on Twitter or LinkedIn.

### **Cautionary Note Regarding Forward-Looking Statements**

*This press release contains forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995, as amended, including, without limitation, implied and express statements regarding its: strategy, business plans and objectives for the IRAK4, IRAK1MiD and STAT3 degrader programs; and plans and timelines for the clinical development of Kymera Therapeutics' product candidates, including the therapeutic potential and clinical benefits thereof. The words "may," "might," "will," "could," "would," "should," "expect," "plan," "anticipate," "intend," "believe," "expect," "estimate," "seek," "predict," "future," "project," "potential," "continue," "target" and similar words or expressions are intended to identify forward-looking statements, although not all forward-looking statements contain these identifying words. Any forward-looking statements in this press release are based on management's current expectations and beliefs and are subject to a number of risks, uncertainties and important factors that may cause actual events or results to differ materially from those expressed or implied by any forward-looking statements contained in this press release, including, without limitation, risks associated with: the impact of COVID-19 on countries or regions in which we have operations or do business, as well as on the timing and anticipated results of our current preclinical studies and future clinical trials, strategy and future operations; the delay of any current preclinical studies or future clinical trials or the development of Kymera Therapeutics' drug candidates; the risk that the results of current preclinical studies may not be predictive of future results in connection with future clinical trials; Kymera Therapeutics' ability to successfully demonstrate the safety and efficacy of its drug candidates; the timing and outcome of the Company's planned interactions with regulatory authorities; and obtaining, maintaining and protecting its intellectual property. These and other risks and uncertainties are described in greater detail in the section entitled "Risk Factors" in the Quarterly Report on Form 10-Q for the period ended March 31, 2021, expected to be filed on or about August 5, 2021, as well as discussions of potential risks, uncertainties, and other important factors in Kymera Therapeutics' subsequent filings with the Securities and Exchange Commission. In addition, any forward-looking statements represent Kymera Therapeutics' views only as of today and should not be relied upon as representing its views as of any subsequent date. Kymera Therapeutics explicitly disclaims any obligation to update any forward-looking statements. No representations or warranties (expressed or implied) are made about the accuracy of any such forward-looking statements.*

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