# MaxCyte Signs Strategic Platform License Agreement with Legend Biotech to Accelerate Cell Therapy Discovery and Development

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**Legend Biotech obtains license to use MaxCyte's Flow Electroporation® technology and ExPERT™ platform to support its non-viral engineered pipeline portfolio across a variety of cell types including T cells, Gamma-Delta T cells and NK Cells.**

ROCKVILLE, Md., May 22, 2024 (GLOBE NEWSWIRE) -- MaxCyte, Inc., (Nasdaq: MXCT; LSE: MXCT), a leading, cell-engineering focused company providing enabling platform technologies to advance the discovery, development, and commercialization of next-generation cell-based therapeutics and innovative bioprocessing applications, today announced it has entered into a strategic platform license (SPL) agreement with Legend Biotech, a global leader in cell therapy.

Under the terms of the SPL, Legend Biotech obtains a non-exclusive worldwide license to use MaxCyte's Flow Electroporation® technology and ExPERT™ platform in connection with the research, clinical development and commercialization of cell-based therapeutical products (Licensed Products). In return, MaxCyte will be eligible to receive annual licensing fees and milestones from Legend Biotech during clinical development and, upon successful commercialization, is eligible to receive licensing fees and royalties on net sales of licensed products.

"We are looking forward to supporting Legend Biotech's non-viral engineered cell therapy program as they expand their portfolio with new delivery modalities. As a leading provider of cell-engineered platform technologies for drug developers, our global infrastructure allows us to provide Legend Biotech with technical, scientific, and regulatory support to advance its non-viral engineered therapeutic pipeline across all major regions," said **Maher Masoud, President and CEO of MaxCyte**. "We are thrilled to enable Legend Biotech in scaling and optimizing its manufacturing process to meet their expanding clinical and commercial needs."

"MaxCyte's clinical manufacturing platform, non-viral cell engineering technology, and regulatory expertise will support the development of our product pipeline across a wide variety of cell types and modalities," said **Ying Huang PhD, Chief Executive Officer of Legend Biotech**. "Our goal is to transform the treatment landscape by creating a broad portfolio of cell therapies to help strengthen patients' immune systems and fight disease."

MaxCyte's ExPERT™ instrument portfolio is the next generation of leading, clinically-validated electroporation technology for complex and scalable cell engineering. By delivering high transfection efficiency, seamless scalability and enhanced functionality, the ExPERT™ platform delivers the high-end performance essential to enabling the next wave of biological and cellular therapeutics. Legend Biotech is MaxCyte's 28 $^{th}$ clinical / commercial partnership overall, each partnership generates pre-commercial milestone revenue, the vast majority of which includes program-related revenue.

**About MaxCyte**

At MaxCyte, we pursue cell engineering excellence to maximize the potential of cells to improve patients' lives. We have spent more than 20 years honing our expertise by building best-in-class platforms, perfecting the art of the transfection workflow, and venturing beyond today's processes to innovate tomorrow's solutions. Our ExPERT™ platform, which is based on our Flow Electroporation® technology, has been designed to support the rapidly expanding cell therapy market and can be utilized across the continuum of the high-growth cell therapy sector, from discovery and development through commercialization of next-generation, cell-based medicines. The ExPERT family of products includes: four instruments, the ATx™, STx™, GTx™ and VLx ™; a portfolio of proprietary related processing assemblies or disposables; and software protocols, all supported by robust worldwide intellectual property portfolio. By providing our partners with the right technology platform, as well as scientific, technical and regulatory support, we aim to guide them on their journey to transform human health. Learn more at maxcyte.com and follow us on X (formerly Twitter) and LinkedIn.

**Forward-Looking Statements**

This press release contains "forward-looking statements" within the meaning of the "safe harbor" provisions of the Private Securities Litigation Reform Act of 1995. These statements about us and our industry involve substantial known and unknown risks, uncertainties, and assumptions, including those described in Item 1A under the heading "Risk Factors" and elsewhere in our report on Form 10-K, that may cause our actual results, performance or achievements to be materially different from any future results, performance or achievements expressed or implied by the forward-looking statements. All statements other than statements of historical facts contained in this press release, including statements regarding our future results of operations or financial condition, business strategy and plans and objectives of management for future operations, are forward-looking statements. Forward-looking statements include, but are not limited to, statements about the Company's expectations regarding annual licensing revenue, pre-commercial milestone revenue, including program related revenue and royalties on sales of products. In some cases, you can identify forward-looking statements because they contain words such as "may," "might," "will," "could," "would," "should," "expect," "plan," "anticipate," "intend," "believe," "expect," "estimate," "seek," "predict," "future," "project," "potential," "continue," "contemplate," "target," the negative of these words and similar words or expressions. These statements are inherently uncertain, and investors are cautioned not to unduly rely on these statements. The forward-looking statements contained in this press release, include, without limitation, statements concerning the following: our expected future growth and success of our business model; the size and growth potential of the markets for our products, and our ability to serve those markets, increase our market share, and achieve and maintain industry leadership; our ability to expand our customer base and enter into additional SPL partnerships; our expectation that our partners will have access to capital markets to develop and commercialize their cell therapy programs; our financial performance and capital requirements; the adequacy of our cash resources and availability of financing on commercially reasonable terms; our expectations regarding our ability to obtain and maintain intellectual property protection for our products, as well as our ability to operate our business without infringing the intellectual property rights of others; our expectations regarding general market and economic conditions that may impact investor confidence in the biopharmaceutical industry and affect the amount of capital such investors provide to our current and potential partners; and our use of available capital resources.

These and other risks and uncertainties are described in greater detail in Item 1A , entitled "Risk Factors," in our Annual Report on Form 10-K for the year ended December 31, 2023, filed with the Securities and Exchange Commission ("SEC") on March 12, 2024, as well as in discussions of potential risks, uncertainties, and other important factors in the other filings that we make with the Securities and Exchange Commission from time to time, including in our Form 10-Q for the quarter ended March 31, 2024, filed with the SEC on May 7, 2024. These documents are available through the Investor Menu, Financials section, under "SEC Filings" on the Investors page of our website at http://investors.maxcyte.com. Any forward-looking

statements in this press release are based on our current beliefs and opinions on the relevant subject based on information available to us as of the date of such press release, and you should not rely on forward-looking statements as predictions of future events. We undertake no obligation to update any forward-looking statements made in this press release to reflect events or circumstances after the date of this press release or to reflect new information or the occurrence of unanticipated events, except as required by law.

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