Replimune to Present at the 39th Annual Meeting of the Society for Immunotherapy of Cancer (SITC)

October 30, 2024

Late-breaking abstract featuring IGNYTE clinical trial data, including subgroup and initial biomarker analyses, selected for oral presentation

Data from ARTACUS clinical trial of RP1 monotherapy in solid organ transplant patients with advanced cutaneous malignancies selected for encore poster presentation

WOBURN, Mass., Oct. 30, 2024 (GLOBE NEWSWIRE) -- Replimune Group, Inc. (NASDAQ: REPL), a clinical stage biotechnology company pioneering the development of novel oncolytic immunotherapies, today announced that a late-breaking abstract featuring the IGNYTE clinical trial primary analysis has been selected for oral presentation at the 39th Annual Meeting of the Society for Immunotherapy of Cancer (SITC 2024) being held November 6-10, 2024, in Houston, Texas. Data from the ARTACUS clinical trial of RP1 monotherapy in solid organ transplant patients with advanced cutaneous malignancies will also be shared in an encore poster presentation.

Presentation Details

Title: Primary analysis of the registration-intended cohort of patients with anti-PD1 failed melanoma from the IGNYTE trial of RP1 plus nivolumab, including clinical subgroup and initial biomarker data
Abstract Number: 1504
Presentation Session Title: Session 204 – Clinical Late-Breaking Abstract Session
Date and Time: Saturday, November 9, 2024, 11:45 am – 12:45 pm CST
Presenter: Michael K. Wong, MD, PhD, University of Texas MD Anderson Cancer Center

Title: Safety and efficacy results from an open-label phase 1b/2 study of RP1 oncolytic immunotherapy in solid organ transplant recipients with advanced cutaneous malignancies (ARTACUS)
Abstract Number: 693
Date and Time: Friday, November 8, 2024
Presenter: Diwakar Davar, MD, UPMC Hillman Cancer Center

About Replimune
Replimune Group, Inc., headquartered in Woburn, MA, was founded in 2015 with the mission to transform cancer treatment by pioneering the development of novel oncolytic immunotherapies. Replimune's proprietary RPx platform is based on a potent HSV-1 backbone intended to maximize immunogenic cell death and the induction of a systemic anti-tumor immune response. The RPx platform is designed to have a unique dual local and systemic activity consisting of direct selective virus-mediated killing of the tumor resulting in the release of tumor derived antigens and altering of the tumor microenvironment to ignite a strong and durable systemic response. The RPx product candidates are expected to be synergistic with most established and experimental cancer treatment modalities, leading to the versatility to be developed alone or combined with a variety of other treatment options. For more information, please visit www.replimune.com.

Forward Looking Statements
This press release contains forward looking statements within the meaning of Section 27A of the Securities Act of 1933, as amended, and Section 21E of the Securities Exchange Act of 1934, as amended, including statements regarding the design and advancement of our clinical trials, the timing and sufficiency of our clinical trial outcomes to support potential approval of any of our product candidates, our goals to develop and commercialize our product candidates, patient enrollments in our existing and planned clinical trials and the timing thereof, and other statements identified by words such as "could," "expects," "intends," "may," "plans," "potential," "should," "will," "would," or similar expressions and the negatives of those terms. Forward-looking statements are not promises or guarantees of future performance, and are subject to a variety of risks and uncertainties, many of which are beyond our control, and which could cause actual results to differ materially from those contemplated in such forward-looking statements. These factors include risks related to our limited operating history, our ability to generate positive clinical trial results for our product candidates, the costs and timing of operating our in-house manufacturing facility, the timing and scope of regulatory approvals, the availability of combination therapies needed to conduct our clinical trials, changes in laws and regulations to which we are subject, competitive pressures, our ability to identify additional product candidates, political and global macro factors including the impact of the coronavirus as a global pandemic and related public health issues and the Russian-Ukrainian and Israel-Hamas political and military conflicts, and other risks as may be detailed from time to time in our Annual Reports on Form 10-K and Quarterly Reports on Form 10-Q and other reports we file with the Securities and Exchange Commission. Our actual results could differ materially from the results described in or implied by such forward-looking statements. Forward-looking statements speak only as of the date hereof, and, except as required by law, we undertake no obligation to update or revise these forward-looking statements.

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