

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

**TÜBINGEN, Germany/BOSTON, USA – October 28, 2024** – CureVac N.V. (Nasdaq: CVAC) (“CureVac”), a global biopharmaceutical company developing a new class of transformative medicines based on messenger ribonucleic acid (“mRNA”), today announced poster presentations at the Society for Immunotherapy of Cancer (SITC) 39<sup>th</sup> Annual Meeting, taking place November 8-10 in Houston, USA.

CureVac will present, among other data, extended preliminary immunogenicity results from Part A of the dose escalation phase of its ongoing Phase 1 CVGBM cancer vaccine study in patients with resected glioblastoma. The presentation will expand on safety, tolerability and immunogenicity results of the CVGBM trial presented last month at the European Society for Medical Oncology (ESMO) Congress, which demonstrated treatment with CVGBM monotherapy successfully induced cancer antigen-specific T-cell responses in 77% of evaluable patients, 84% of which were *de novo*.

### **Details on all poster presentations are below:**

**Abstract: 697**

**Title:** Preliminary immunogenicity results from the dose escalation phase of a first-in-human study of the mRNA-based cancer vaccine CVGBM in patients with newly diagnosed MGMT-unmethylated glioblastoma

**Session type:** Poster Presentation

**Date:** November 8

**Presenting Author:** Sven Koch, Ph.D., Director Immuno-Monitoring, CureVac SE

**Abstract: 380**

**Title:** mRNA vaccination enhances TCRtg T cell therapy efficacy in solid tumors

**Session type:** Poster Presentation

**Date:** November 9

**Presenting Author:** Dr. Johannes Lutz, Director Pre-Clinical Development, CureVac SE

**Abstract: 370**

**Title:** Identification and validation of a T cell receptor recognizing shared HLA-A02:01 presented epitope from TP53 frameshift

**Session type:** Poster Presentation

**Date:** November 9

**Presenting Author:** Katka Franke, Ph.D., Pharm.D., Director Oncology Antigen Discovery and Validation, CureVac Netherlands B.V.

## **About CureVac**

CureVac (Nasdaq: CVAC) is a pioneering multinational biotech company founded in 2000 to advance the field of messenger RNA (mRNA) technology for application in human medicine. In more than two decades of developing, optimizing, and manufacturing this versatile biological molecule for medical purposes, CureVac has introduced and refined key underlying technologies that were essential to the production of mRNA vaccines against COVID-19, and is currently laying the groundwork for application of mRNA in new therapeutic areas of major unmet need. CureVac is leveraging mRNA technology, combined with advanced omics and computational tools, to design and develop off-the-shelf and personalized cancer vaccine product candidates. It also develops programs in prophylactic vaccines and in treatments that enable the human body to produce its own therapeutic proteins. Headquartered in Tübingen, Germany, CureVac also operates sites in the Netherlands, Belgium, Switzerland, and the U.S. Further information can be found at [www.curevac.com](http://www.curevac.com).

## **CureVac Media and Investor Relations Contact**

Dr. Sarah Fakh, Vice President Corporate Communications and Investor Relations  
CureVac, Tübingen, Germany  
T: +49 7071 9883-1298  
M: +49 160 90 496949  
[sarah.fakh@curevac.com](mailto:sarah.fakh@curevac.com)

## **Forward-Looking Statements CureVac**

This press release contains statements that constitute “forward looking statements” as that term is defined in the United States Private Securities Litigation Reform Act of 1995, including statements that express the opinions, expectations, beliefs, plans, objectives, assumptions or projections of CureVac N.V. and/or its wholly owned subsidiaries CureVac SE, CureVac Manufacturing GmbH, CureVac Inc., CureVac Swiss AG, CureVac Corporate Services GmbH, CureVac Belgium SA and CureVac Netherlands B.V. (the “company”) regarding future events or future results, in contrast with statements that reflect historical facts. Examples include discussion of the potential efficacy of the company’s vaccine and treatment candidates and the company’s strategies, financing plans, cash runway expectations, growth opportunities and market growth. In some cases, you can identify such forward-looking statements by terminology such as “anticipate,” “intend,” “believe,” “estimate,” “plan,” “seek,” “project,” or “expect,” “may,” “will,” “would,” “could,” “potential,” “intend,” or “should,” the negative of these terms or similar expressions. Forward-looking statements are based on management’s current beliefs and assumptions and on information currently available to the company. However, these forward-looking statements are not a guarantee of the company’s performance, and you should not place undue reliance on such statements. Forward-looking statements are subject to many risks, uncertainties and other variable circumstances, including negative worldwide economic conditions and ongoing instability and volatility in the worldwide financial markets, ability to obtain funding, ability to conduct current and future preclinical

studies and clinical trials, the timing, expense and uncertainty of regulatory approval, reliance on third parties and collaboration partners, ability to commercialize products, ability to manufacture any products, possible changes in current and proposed legislation, regulations and governmental policies, pressures from increasing competition and consolidation in the company's industry, the effects of the COVID-19 pandemic on the company's business and results of operations, ability to manage growth, reliance on key personnel, reliance on intellectual property protection, ability to provide for patient safety, fluctuations of operating results due to the effect of exchange rates, delays in litigation proceedings, different judicial outcomes or other factors. Such risks and uncertainties may cause the statements to be inaccurate and readers are cautioned not to place undue reliance on such statements. Many of these risks are outside of the company's control and could cause its actual results to differ materially from those it thought would occur. The forward-looking statements included in this press release are made only as of the date hereof. The company does not undertake, and specifically declines, any obligation to update any such statements or to publicly announce the results of any revisions to any such statements to reflect future events or developments, except as required by law.

For further information, please reference the company's reports and documents filed with the U.S. Securities and Exchange Commission (SEC). You may get these documents by visiting EDGAR on the SEC website at [www.sec.gov](http://www.sec.gov).