

# BioNTech to Present Data from BNT311 (GEN1046) and BNT131 (SAR441000) Programs at SITC 35th Annual Meeting

October 14, 2020

- Data updates from key oncology collaborations to be presented
- Preliminary data from the first-in-human trial from intratumoral immunotherapy program BNT131 (SAR441000) in collaboration with Sanofi, to be presented in an e-poster
- Preliminary Phase 1/2 and preclinical data of DuoBody®-PD-L1x4-1BB, developed in collaboration with Genmab, to be presented in e-posters

MAINZ, Germany, October 14, 2020 (GLOBE NEWSWIRE) – BioNTech SE (Nasdaq: BNTX, "BioNTech" or "the Company") today announced that results from three clinical and preclinical studies have been accepted for presentation at the Society for Immunotherapy of Cancer (SITC) 35<sup>th</sup> Annual Meeting. The data represent the first clinical results from BioNTech's ongoing collaborations with Genmab and Sanofi from both its mRNA and antibody drug class portfolios. Preliminary data from a Phase 1/2 study of DuoBody-PD-L1x4-1BB (BNT311/GEN1046) in advanced solid tumors as well as preclinical data from the program have been accepted for eposter presentations. In addition, preliminary data from the first in-human dose escalation trial of intratumoral immunotherapy BNT131 (SAR441000) in collaboration with Sanofi will be shared as e-poster presentation. The full abstracts are scheduled to be available on the SITC website on November 9, 2020.

#### **Poster Presentation Details:**

Title: First-in-human phase I/IIa trial to evaluate the safety and initial clinical activity of DuoBody<sup>®</sup>-PD L1×4-1BB (GEN1046) in patients with advanced solid tumors

Poster Presentation Date & Time: 11-14 Nov 2020, 9 am - 5 pm

Abstract Number: 630 Presenter: Ignacio Melero

**Title:** DuoBody<sup>®</sup>-PD-L1×4-1BB (GEN1046) induces superior immune-cell activation, cytokine production and cytotoxicity by combining PD-L1 blockade with conditional 4-1BB co-stimulation

Poster Presentation Date & Time: 11-14 Nov 2020, 9 am - 5 pm

Abstract Number: 561
Presenter: Alexander Muik

**Title:** A first-in-human study of intratumoral SAR441000, an mRNA mixture encoding IL-12sc, interferon alpha2b, GM-CSF and IL-15sushi as monotherapy and in combination with cemiplimab in advanced solid tumors

Presentation Date & Time: 11-14 Nov 2020, 9 am - 5 pm

Abstract Number: 391 Presenter: Oliver Bechter

## **About BioNTech**

Biopharmaceutical New Technologies is a next generation immunotherapy company pioneering novel therapies for cancer and other serious diseases. The Company exploits a wide array of computational discovery and therapeutic drug platforms for the rapid development of novel biopharmaceuticals. Its broad portfolio of oncology product candidates includes individualized and off-the-shelf mRNA-based therapies, innovative chimeric antigen receptor T cells, bi-specific checkpoint immuno-modulators, targeted cancer antibodies and small molecules. Based on its deep expertise in mRNA vaccine development and in-house manufacturing capabilities, BioNTech and its collaborators are developing multiple mRNA vaccine candidates for a range of infectious diseases alongside its diverse oncology pipeline. BioNTech has established a broad set of relationships with multiple global pharmaceutical collaborators, including Genmab, Sanofi, Bayer Animal Health, Genentech, a member of the Roche Group, Genevant, Fosun Pharma, and Pfizer.

For more information, please visit www.BioNTech.de.

# **BioNTech's Forward-Looking Statements**

This press release contains "forward-looking statements" of BioNTech within the meaning of the Private Securities Litigation Reform Act of 1995. These forward-looking statements may include, but may not be limited to, statements concerning: availabliity of clinical trial data for BioNTech's program candidates, including BNT311 (GEN1046) and BNT131 (SAR441000). Any forward-looking statements in this press release are based on BioNTech management's current expectations and beliefs of future events, and are subject to a number of risks and uncertainties that could cause actual results to differ materially and adversely from those set forth in or implied by such forward-looking statements. For a discussion of these and other risks and uncertainties, see BioNTech's Annual Report on Form 20-F filed with the SEC on March 31, 2020, which is available on the SEC's website <a href="https://www.sec.gov">www.sec.gov</a>. All information in this press release is as of the date of the release, and BioNTech undertakes no duty to update this information unless required by law.

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