UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
Washington, D.C. 20549

FORM 6-K

REPORT OF FOREIGN PRIVATE ISSUER PURSUANT TO RULE 13a-16 OR 15d-16
UNDER THE SECURITIES EXCHANGE ACT OF 1934

FOR THE MONTH OF MAY 2025

COMMISSION FILE NUMBER 001-39081

BioNTech SE
(Translation of registrant's name into English)

An der Goldgrube 12
D-55131 Mainz
Germany
+49 6131-9084-0
(Address of principal executive offices)

Indicate by check mark whether the registrant files or will file annual reports under cover Form 20-F or Form 40-F: Form 20-F ☒ Form 40-F ☐

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(1): ☐

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(7): ☐

DOCUMENTS INCLUDED AS PART OF THIS FORM 6-K

On May 5, 2025, BioNTech SE (the "Company") issued a press release announcing its first quarter 2025 financial results and corporate update and details of a conference call to be held at 8:00 am EDT on May 5, 2025 to discuss the results. The press release and the conference call presentation are attached as Exhibits 99.1 and 99.2, respectively, and incorporated by reference herein.

The information contained in Exhibits 99.1 and 99.2 shall not be deemed "filed" for purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the "Exchange Act"), or incorporated by reference in any filing under the Securities Act of 1933, as amended, or the Exchange Act, unless expressly set forth by specific reference in such a filing.

SIGNATURE

Pursuant to the requirements of the Exchange Act, the registrant has duly caused this report to be signed on its behalf by the undersigned, thereunto duly authorized.

BioNTech SE

By: /s/ Jens Holstein
Name: Jens Holstein
Title: Chief Financial Officer

By: /s/ Dr. Sierk Poetting
Name: Dr. Sierk Poetting
Title: Chief Operating Officer

Date: May 5, 2025

EXHIBIT INDEX

Exhibit	Description of Exhibit
99.1	BioNTech Announces First Quarter 2025 Financial Results and Corporate Update
99.2	First Quarter 2025: Corporate Update and Financial Results

BioNTech Announces First Quarter 2025 Financial Results and Corporate Update

- *Continued oncology pipeline advancement with a strategic focus on two priority pan-tumor programs: next-generation immunomodulator BNT327, a bispecific antibody targeting PD-L1 and VEGF-A[1], and mRNA cancer immunotherapies*
- *Presented multiple clinical updates across oncology pipeline underlining BioNTech's combination strategy in oncology with first data presented for the novel combination of BNT327 plus antibody-drug conjugates ("ADCs")*
- *Development and commercial preparation for a 2025/2026 season variant-adapted COVID-19 vaccine*
- *First quarter 2025 revenues of €0.2 billion[2], net loss of €0.4 billion and basic and diluted loss per share of €1.73 ($1.82[3])*
- *Maintained strong financial position with €15.9 billion in cash, cash equivalents and security investments as of March 31, 2025[4]*
- *Full year 2025 financial guidance confirmed*

Conference call and webcast scheduled for May 5, 2025, at 8:00 a.m. EDT (2:00 p.m. CEST)

MAINZ, Germany, May 5, 2025 (GLOBE NEWSWIRE) -- BioNTech SE (Nasdaq: BNTX, "BioNTech" or "the Company") today reported financial results for the three months ended March 31, 2025 and provided an update on its corporate progress.

"In the first quarter of 2025, we demonstrated continued execution against our strategic focus areas, highlighted by data updates for our PD-L1xVEGF-A bispecific antibody candidate BNT327 and the progress in clinical evaluation of our focus programs and combination treatment approaches," **said Prof. Ugur Sahin, M.D., CEO and Co-Founder of BioNTech**. "We will continue to focus on our key strategic programs as we remain steadfast in our vision to translate our science into survival for patients in need."

Financial Review for First Quarter 2025

in millions €, except per share data	First Quarter 2025	First Quarter 2024
Revenues	182.8	187.6
Net loss	(415.8)	(315.1)
Basic and diluted loss per share	(1.73)	(1.31)

Revenues reported were €182.8 million for the three months ended March 31, 2025, compared to €187.6 million for the comparative prior year period. Revenues during the first quarter of 2025 were mainly driven by revenues derived from BioNTech's COVID-19 vaccine collaboration.

Cost of sales were €83.8 million for the three months ended March 31, 2025, compared to €59.1 million for the comparative prior year period. The change was mainly due to a positive impact of an inventory revaluation in the first quarter of 2024.

Research and development ("R&D") expenses were €525.6 million for the three months ended March 31, 2025, compared to €507.5 million for the comparative prior year period. The increase was mainly driven by progressing late-stage clinical studies for candidates in BioNTech's ADC and antibody portfolio.

Sales, general and administrative ("SG&A") expenses[5], in total, amounted to €120.6 million for the three months ended March 31, 2025, compared to €132.6 million for the comparative prior year period. The decrease was primarily driven by a reduction in external services.

Income taxes were realized with an amount of €29.6 million in tax income for the three months ended March 31, 2025, compared to €16.7 million in realized tax income for the comparative prior year period.

Net loss was €415.8 million for the three months ended March 31, 2025, compared to a net loss of €315.1 million for the comparative prior year period.

Cash and cash equivalents plus security investments as of March 31, 2025, reached €15,854.4 million, comprising €10,184.9 million in cash and cash equivalents, €3,542.0 million in current security investments and €2,127.5 million in non-current security investments.

Basic and diluted loss per share was €1.73 for the three months ended March 31, 2025, compared to a basic and diluted loss per share of €1.31 for the comparative prior year period.

Shares outstanding as of March 31, 2025, were 240,392,622, excluding 8,159,578 shares held in treasury.

"Our revenues for the first quarter reflect the seasonal demand for COVID-19 vaccines and are in line with our expectations," **said Jens Holstein, CFO of BioNTech**. "BioNTech's robust financial position empowers us to pursue our strategic goal of evolving into a leading biotech company with multiple oncology products by 2030."

2025 Financial Year Guidance Confirmed[6]

Total revenues for the 2025 financial year	**€1,700 million - €2,200 million**

BioNTech expects its revenues for the full 2025 financial year to be in the range of €1,700 - €2,200 million and revenue phasing similar to 2024, primarily concentrated in the last three to four months, driving the full year revenue figure. The revenue guidance assumes: relatively stable vaccination rates, pricing levels and market share compared to 2024; estimated inventory write-downs and other charges by BioNTech's collaboration partner Pfizer that negatively influence BioNTech's revenues; anticipated revenues from a pandemic preparedness contract with the German government; and anticipated revenues from the BioNTech Group service businesses. Potential changes to the law or governmental policy, including tariffs and public health policy, and evolving public sentiment worldwide, could further negatively impact our anticipated revenues and expenses.

Planned 2025 Financial Year Expenses and Capex

R&D expenses	**€2,600 million - €2,800 million**
SG&A expenses	**€650 million - €750 million**
Capital expenditures for operating activities	**€250 million - €350 million**

BioNTech expects to continuously focus investments on R&D and scaling the business for late-stage development and commercial readiness in oncology, while remaining cost-disciplined. Strategic capital allocation will continue to be a key driver of the Company's trajectory. As part of BioNTech's strategy, the Company may continue to evaluate appropriate corporate development opportunities with the aim of driving sustainable long-term growth and create future value.

The full interim unaudited condensed consolidated financial statements can be found in BioNTech's Report on Form 6-K for the period ended March 31, 2025, filed today with the United States Securities and Exchange Commission ("SEC") and available at www.sec.gov.

Endnotes

[1] All abbreviations are compiled in an abbreviation directory at the end of this press release.
[2] All numbers in this press release have been rounded.
[3] Calculated applying the average foreign exchange rate for the three months ended March 31, 2025, as published by the German Central Bank (Deutsche Bundesbank).
[4] A settlement payment of $400 million related to a contractual dispute with the University of Pennsylvania is expected to be reflected in the Company's second quarter 2025 financial results. In connection with this and another settlement with the National Institutes of Health ("NIH"), BioNTech expects to be reimbursed approximately $535 million by its collaboration partner during 2025 and 2026. Reimbursement payments have begun with the first payment received by BioNTech in the first quarter of 2025.
[5] Sales, general and administrative expenses ("SG&A") include sales and marketing expenses as well as general and administrative expenses.
[6] Financial guidance excludes external risks that are not yet known and/or quantifiable, including, but not limited to the effects of ongoing and/or future legal disputes and related activities, certain potential one-time effects and charges related to portfolio prioritization. It includes effects identified from licensing arrangements, collaborations or potential M&A transactions to the extent disclosed and may be subject to update. The Company does not expect to report a positive net income figure for the 2025 financial year.

Operational Review for the First Quarter 2025, Key Post Period-End Events and 2025 Outlook

Selected Oncology Pipeline Updates

Next-Generation Immunomodulators

BNT327, formerly also known as PM8002, is a bispecific antibody candidate combining PD-L1 checkpoint inhibition with VEGF-A neutralization.

- In March 2025, at the European Lung Cancer Congress ("ELCC"), preliminary data from two Phase 2 clinical trials conducted in China in first-line extensive-stage small cell lung cancer ("ES-SCLC") and second-line small cell lung cancer ("SCLC") were presented.
    - Preliminary data from the ongoing single-arm, open-label Phase 2 clinical trial (NCT05844150) evaluating BNT327 in combination with chemotherapy as a first-line treatment in patients with ES-SCLC showed anti-tumor activity and an acceptable safety profile with no new safety signals beyond those typically described for chemotherapy agents and anti-PD-(L)1 and anti-VEGF monotherapies. These data were the first presented for BNT327 as a potential first-line treatment in ES-SCLC supporting the ongoing global randomized Phase 3 clinical trial ROSETTA Lung-01 (NCT06712355).
    - Preliminary data from the ongoing Phase 2 clinical trial (NCT05879068) evaluating BNT327 in combination with chemotherapy as a second-line treatment in patients with SCLC showed anti-tumor activity, which was observed regardless of prior immuno-oncology treatment, and an acceptable safety profile.
- In April 2025, at the American Association for Cancer Research ("AACR") Annual Meeting 2025, first data were presented for the novel combination of the PD-L1xVEGF-A bispecific antibody candidate BNT327 with various ADC candidates.
    - Interim data from the ongoing Phase 1/2 clinical trial (NCT05438329) evaluating BNT327 with BNT325/DB-1305, a TROP2-targeting ADC candidate being developed

in collaboration with Duality Biologics (Suzhou) Co. Ltd. (“DualityBio”), in patients with advanced/metastatic solid tumors showed a manageable safety profile and early signs of anti-tumor activity in a cohort with patients with platinum-resistant ovarian cancer (“PROC”). Across the 13 efficacy evaluable patients with PROC, seven patients achieved partial response and three stable disease. Responses were also observed in patients with non-small cell lung cancer (“NSCLC”) or triple-negative breast cancer (“TNBC”).

- First data from an ongoing Phase 2 clinical trial (NCT05918107) in first-line mesothelioma and two trial-in-progress posters for the ongoing global Phase 3 and Phase 2/3 clinical trials, ROSETTA Lung-01 (NCT06712355) and ROSETTA Lung-02 (NCT06712316) respectively, will be presented at the American Society for Clinical Oncology (“ASCO”) Annual Meeting taking place from May 30 to June 3, 2025 in Chicago, Illinois.
    - *Abstract title:* First report of efficacy and safety results from a Phase 2 trial evaluating BNT327/PM8002 plus chemotherapy (chemo) as first-line treatment (1L) in unresectable malignant mesothelioma
    - *Abstract title:* A global Phase III, double-blind, randomized trial of BNT327/PM8002 plus chemotherapy (chemo) compared to atezolizumab plus chemo in patients (pts) with first-line (1L) extensive-stage small cell lung cancer (ES-SCLC)
    - *Abstract title:* A global Phase 2/3, randomized, open-label trial of BNT327/PM8002 in combination with chemotherapy (chemo) in first-line (1L) non-small cell lung cancer (NSCLC)

mRNA Cancer Immunotherapies

BNT116 is based on BioNTech’s fully owned, off-the-shelf FixVac platform, and is designed to elicit an immune response to six tumor-associated antigens that were identified to be frequently expressed in NSCLC.

- In April 2025, at the AACR Annual Meeting 2025, preliminary data from a cohort with frail patients were presented from the ongoing Phase 1 clinical trial LuCa-MERIT-1 (NCT05142189) evaluating BNT116 in combination with cemiplimab in patients with PD-L1 positive (TPS≥1%) unresectable Stage III or metastatic Stage IV NSCLC who are not eligible for chemotherapy as first-line treatment. The preliminary data showed anti-tumor activity, consistent immune response induction and a manageable safety profile.

Antibody-Drug Conjugates

BNT324/DB-1311 is an B7H3-targeted ADC candidate that is being developed in collaboration with DualityBio. The program has received Fast Track designation from the U.S. Food and Drug Administration (“FDA”) for the treatment of patients with advanced castration-resistant prostate cancer (“CRPC”) whose disease has progressed while undergoing or after undergoing standard systemic regimens. In addition, the program has received Orphan Drug designation from the FDA for the treatment of patients with advanced esophageal squamous cell carcinoma.

- Preliminary data from the ongoing Phase 1/2 clinical trial (NCT05914116) evaluating BNT324/DB-1311 in patients with advanced solid tumors, including patients with previously treated castration-resistant prostate cancer (“CRPC”), will be presented at the 2025 ASCO Annual Meeting.

- *Abstract title:* DB-1311/BNT324 (a novel B7H3 ADC) in patients with heavily pretreated castrate-resistant prostate cancer (CRPC)

BNT325/DB-1305 is a TROP2-targeted ADC candidate that is being developed in collaboration with DualityBio. BNT325/DB-1305 received Fast Track designation from the FDA for the treatment of patients with platinum-resistant ovarian epithelial cancer, fallopian tube cancer, or primary peritoneal cancer who have received one to three prior systemic treatment regimens.

- In March 2025, at the Society of Gynecologic Oncology ("SGO") Annual Meeting, preliminary data were presented from an ongoing Phase 1/2 clinical trial (NCT05438329) evaluating BNT325/DB-1305 in signal-seeking cohorts across various cancer indications, including PROC. The data from a cohort of patients with PROC showed a manageable safety profile and early signs of anti-tumor activity.

Corporate and Commercial Update for the First Quarter 2025

- Earlier today, BioNTech announced that the Supervisory Board has appointed Ramón Zapata-Gomez to the Management Board as Chief Financial Officer ("CFO") effective July 1, 2025. He will join BioNTech from Novartis AG's global biomedical research organization where he has been serving as CFO since 2022. Ramón Zapata will succeed Jens Holstein, who, as previously planned and announced, will retire at the end of his term on June 30, 2025.

- In February 2025, BioNTech completed the acquisition of Biotheus, obtaining full global rights to BNT327 and to all other candidates from Biotheus' pipeline, as well as to its in-house antibody generation platform and bispecific ADC capability. The transaction amounted to an upfront consideration of $800 million, plus additional performance-based payments of up to $150 million.

- BioNTech and Pfizer developed, manufactured and delivered their JN.1 and KP.2-adapted COVID-19 vaccines, which have received multiple regulatory approvals, including full approvals, authorizations for emergency or temporary use, or marketing authorizations, in more than 40 countries and regions. BioNTech is now focused on preparing for variant strain vaccine adaptation to be ready for commercial launch ahead of the upcoming 2025/2026 vaccination season, pending approvals.

Upcoming Investor and Analyst Events

- Annual General Meeting: May 16, 2025
- Innovation Series R&D Day: November 11, 2025

Conference Call and Webcast Information

BioNTech invites investors and the general public to join a conference call and webcast with investment analysts today, May 5, 2025, at 8:00 a.m. EDT (2:00 p.m. CEST) to report its financial results and provide a corporate update for the first quarter of 2025.

To access the live conference call via telephone, please register via this link. Once registered, dial-in numbers and a PIN number will be provided.

The slide presentation and audio of the webcast will be available via this link.

Participants may also access the slides and the webcast of the conference call via the "Events & Presentations" page of the Investor section of the Company's website at www.BioNTech.com. A replay

of the webcast will be available shortly after the conclusion of the call and archived on the Company's website for 30 days following the call.

About BioNTech

Biopharmaceutical New Technologies (BioNTech) is a global next generation immunotherapy company pioneering novel investigative therapies for cancer and other serious diseases. BioNTech exploits a wide array of computational discovery and therapeutic modalities with the intent of rapid development of novel biopharmaceuticals. Its diversified portfolio of oncology product candidates aiming to address the full continuum of cancer includes mRNA cancer immunotherapies, next-generation immunomodulators and targeted therapies such as antibody-drug conjugates (ADCs) and innovative chimeric antigen receptor (CAR) T cell therapies. Based on its deep expertise in mRNA development and in-house manufacturing capabilities, BioNTech and its collaborators are researching and 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 and specialized pharmaceutical collaborators, including Duality Biologics, Fosun Pharma, Genentech, a member of the Roche Group, Genevant, Genmab, MediLink, OncoC4, Pfizer and Regeneron.

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

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, but not limited to, statements concerning: BioNTech's expected revenues and net profit/(loss) related to sales of BioNTech's COVID-19 vaccine in territories controlled by BioNTech's collaboration partners, particularly for those figures that are derived from preliminary estimates provided by BioNTech's partners; the rate and degree of market acceptance of BioNTech's COVID-19 vaccine and, if approved, BioNTech's investigational medicines; expectations regarding anticipated changes in COVID-19 vaccine demand, including changes to the ordering environment and expected regulatory recommendations to adapt vaccines to address new variants or sublineages; the initiation, timing, progress, results, and cost of BioNTech's research and development programs, including BioNTech's current and future preclinical studies and clinical trials, including statements regarding the expected timing of initiation, enrollment, and completion of studies or clinical trials and related preparatory work and the availability of results, and the timing and outcome of applications for regulatory approvals and marketing authorizations; BioNTech's expectations regarding potential future commercialization in oncology, including goals regarding timing and indications; the targeted timing and number of additional potentially registrational clinical trials, and the registrational potential of any clinical trial BioNTech may initiate; discussions with regulatory agencies; BioNTech's expectations with respect to intellectual property; the impact of BioNTech's collaboration and licensing agreements; the development, nature and feasibility of sustainable vaccine production and supply solutions; the deployment of AI across BioNTech's preclinical and clinical operations; BioNTech's expectations with respect to tariff policy; BioNTech's estimates of revenues, research and development expenses, selling, general and administrative expenses and capital expenditures for operating activities; BioNTech's expectations regarding upcoming payments relating to litigation settlements; BioNTech's expectations for upcoming scientific and investor presentations; and BioNTech's expectations of net profit / (loss). In some cases, forward-looking statements can be identified by terminology such as "will," "may," "should," "expects," "intends," "plans," "aims," "anticipates," "believes," "estimates," "predicts," "potential," "continue," or the negative of these terms or other comparable terminology, although not all forward-looking statements contain these words.

The forward-looking statements in this press release are based on BioNTech's current expectations and beliefs of future events, and are neither promises nor guarantees. You should not place undue reliance on these forward-looking statements because they involve known and unknown risks,

uncertainties, and other factors, many of which are beyond BioNTech's control and which could cause actual results to differ materially and adversely from those expressed or implied by these forward-looking statements. These risks and uncertainties include, but are not limited to: the uncertainties inherent in research and development, including the ability to meet anticipated clinical endpoints, commencement and/or completion dates for clinical trials, projected data release timelines, regulatory submission dates, regulatory approval dates and/or launch dates, as well as risks associated with preclinical and clinical data, including the data discussed in this release, and including the possibility of unfavorable new preclinical, clinical or safety data and further analyses of existing preclinical, clinical or safety data; the nature of the clinical data, which is subject to ongoing peer review, regulatory review and market interpretation; BioNTech's pricing and coverage negotiations regarding its COVID-19 vaccine with governmental authorities, private health insurers and other third-party payors; the future commercial demand and medical need for initial or booster doses of a COVID-19 vaccine; the impact of tariffs and escalations in trade policy; competition from other COVID-19 vaccines or related to BioNTech's other product candidates, including those with different mechanisms of action and different manufacturing and distribution constraints, on the basis of, among other things, efficacy, cost, convenience of storage and distribution, breadth of approved use, side-effect profile and durability of immune response; the timing of and BioNTech's ability to obtain and maintain regulatory approval for its product candidates; the ability of BioNTech's COVID-19 vaccines to prevent COVID-19 caused by emerging virus variants; BioNTech's and its counterparties' ability to manage and source necessary energy resources; BioNTech's ability to identify research opportunities and discover and develop investigational medicines; the ability and willingness of BioNTech's third-party collaborators to continue research and development activities relating to BioNTech's development candidates and investigational medicines; the impact of COVID-19 on BioNTech's development programs, supply chain, collaborators and financial performance; unforeseen safety issues and potential claims that are alleged to arise from the use of products and product candidates developed or manufactured by BioNTech; BioNTech's and its collaborators' ability to commercialize and market BioNTech's COVID-19 vaccine and, if approved, its product candidates; BioNTech's ability to manage its development and related expenses; regulatory and political developments in the United States and other countries; BioNTech's ability to effectively scale its production capabilities and manufacture its products and product candidates; risks relating to the global financial system and markets; and other factors not known to BioNTech at this time. You should review the risks and uncertainties described under the heading "Risk Factors" in BioNTech's Report on Form 6-K for the period ended March 31, 2025 and in subsequent filings made by BioNTech with the SEC, which are available on the SEC's website at www.sec.gov. These forward-looking statements speak only as of the date hereof. Except as required by law, BioNTech disclaims any intention or responsibility for updating or revising any forward-looking statements contained in this press release in the event of new information, future developments or otherwise.

CONTACTS

Investor Relations
Michael Horowicz
Investors@biontech.de

Media Relations
Jasmina Alatovic
Media@biontech.de

Target Overview

Anti-PD-(L)1	Anti-programmed cell death protein (death-ligand) 1
PD-L1	Programmed death-ligand 1
TROP2	Trophoblast cell-surface antigen 2
VEGF-A	Vascular endothelial growth factor A

Interim Condensed Consolidated Statements of Profit or Loss

	Three months ended March 31,	
	2025	**2024**
(in millions €, except per share data)	*(unaudited)*	*(unaudited)*
Revenues	182.8	187.6
Cost of sales	(83.8)	(59.1)
Research and development expenses	(525.6)	(507.5)
Sales and marketing expenses	(13.7)	(15.6)
General and administrative expenses	(106.9)	(117.0)
Other operating expenses	(48.5)	(23.9)
Other operating income	61.6	28.3
Operating loss	**(534.1)**	**(507.2)**
Finance income	122.6	180.1
Finance expenses	(33.9)	(4.7)
Loss before tax	**(445.4)**	**(331.8)**
Income taxes	29.6	16.7
Net loss	**(415.8)**	**(315.1)**
Loss per share		
Basic and diluted loss per share	(1.73)	(1.31)

Interim Condensed Consolidated Statements of Financial Position

(in millions €)	March 31, 2025 *(unaudited)*	December 31, 2024
Assets		
Non-current assets		
Goodwill	375.9	380.6
Other intangible assets	1,511.2	790.4
Property, plant and equipment	1,026.1	935.3
Right-of-use assets	241.9	248.1
Contract assets	7.9	9.8
Other financial assets	2,285.8	1,254.0
Other non-financial assets	22.6	26.3
Deferred tax assets	85.5	81.7
Total non-current assets	**5,556.9**	**3,726.2**
Current assets		
Inventories	254.4	283.3
Trade and other receivables	956.5	1,463.9
Contract assets	10.0	10.0
Other financial assets	3,924.9	7,021.7
Other non-financial assets	234.3	212.7
Income tax assets	60.5	50.0
Cash and cash equivalents	10,184.9	9,761.9
Total current assets	**15,625.5**	**18,803.5**
Total assets	**21,182.4**	**22,529.7**
Equity and liabilities		
Equity		
Share capital	248.6	248.6
Capital reserve	1,447.4	1,398.6
Treasury shares	(8.2)	(8.6)
Retained earnings	18,682.2	19,098.0
Other reserves	(1,443.4)	(1,325.5)
Total equity	**18,926.6**	**19,411.1**
Non-current liabilities		
Lease liabilities, loans and borrowings	237.5	214.7
Other financial liabilities	150.9	46.9
Provisions	20.8	20.9
Contract liabilities	182.9	183.0
Other non-financial liabilities	85.1	87.5
Deferred tax liabilities	44.1	42.4
Total non-current liabilities	**721.3**	**595.4**
Current liabilities		
Lease liabilities, loans and borrowings	55.5	39.5
Trade payables and other payables	443.8	426.7
Other financial liabilities	443.4	1,443.4
Income tax liabilities	5.4	4.5
Provisions	121.9	144.8
Contract liabilities	294.5	294.9
Other non-financial liabilities	170.0	169.4
Total current liabilities	**1,534.5**	**2,523.2**
Total liabilities	**2,255.8**	**3,118.6**
Total equity and liabilities	**21,182.4**	**22,529.7**

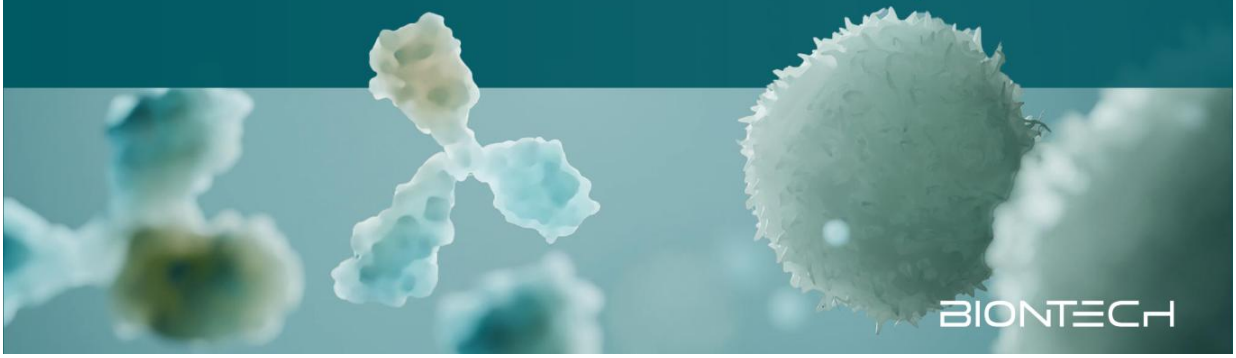
Interim Condensed Consolidated Statements of Cash Flows

(in millions €)	Three months ended March 31, 2025 *(unaudited)*	2024 *(unaudited)*
Operating activities		
Net loss	(415.8)	(315.1)
Income taxes	(29.6)	(16.7)
Loss before tax	**(445.4)**	**(331.8)**
Adjustments to reconcile profit before tax to net cash flows:		
Depreciation and amortization of property, plant, equipment, intangible assets and right-of-use assets	42.8	38.3
Share-based payment expenses	22.1	16.3
Net foreign exchange differences	48.3	(28.7)
(Gain) / Loss on disposal of property, plant and equipment	(0.1)	—
Finance income excluding foreign exchange differences	(122.6)	(174.9)
Finance expense excluding foreign exchange differences	7.9	4.7
Government grants	(14.5)	(9.1)
Other non-cash (income) / loss	(14.3)	—
Unrealized (gain) / loss on derivative instruments at fair value through profit or loss	(11.3)	1.7
Working capital adjustments:		
Decrease in trade and other receivables, contract assets and other assets	520.7	498.2
Decrease in inventories	33.8	12.3
Decrease in trade payables, other financial liabilities, other liabilities, contract liabilities, refund liabilities and provisions	(971.0)	(288.0)
Interest received and realized gains from cash and cash equivalents	118.6	199.4
Interest paid and realized losses from cash and cash equivalents	(3.1)	(3.7)
Income tax paid	(12.2)	(258.8)
Share-based payments	(3.6)	(2.4)
Government grants received	23.2	9.2
Net cash flows used in operating activities	**(780.7)**	**(317.3)**
Investing activities		
Purchase of property, plant and equipment	(48.9)	(58.5)
Proceeds from sale of property, plant and equipment	0.5	—
Purchase of intangible assets and right-of-use assets	(569.2)	(78.4)
Acquisition of subsidiaries and businesses, net of cash acquired	(78.5)	—
Investment in other financial assets	(2,507.7)	(4,895.1)
Proceeds from maturity of other financial assets	4,450.6	2,727.6
Net cash flows from / (used in) investing activities	**1,246.8**	**(2,304.4)**
Financing activities		
Repayment of loans and borrowings	(4.5)	—
Payments related to lease liabilities	(9.3)	(7.8)
Net cash flows used in financing activities	**(13.8)**	**(7.8)**
Net increase / (decrease) in cash and cash equivalents	452.3	(2,629.5)
Change in cash and cash equivalents resulting from exchange rate differences	(16.1)	6.8
Change in cash and cash equivalents resulting from other valuation effects	(13.2)	(64.4)
Cash and cash equivalents at the beginning of the period	9,761.9	11,663.7
Cash and cash equivalents as of March 31	**10,184.9**	**8,976.6**

Exhibit 99.2

May 5th, 2025

1st Quarter 2025
Financial Results & Corporate Update



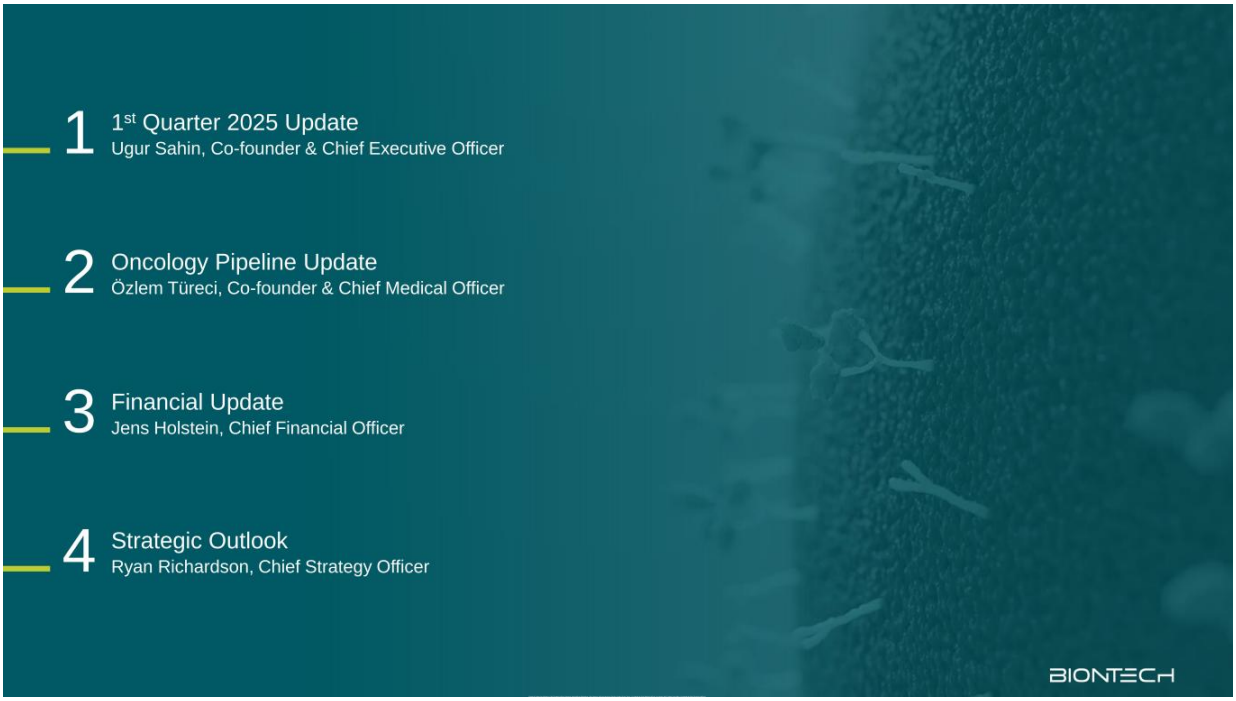
This Slide Presentation Includes Forward-Looking Statements

This presentation contains forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995, as amended, including, but not limited to, statements concerning: BioNTech's expected revenues and net profit/(loss) related to sales of BioNTech's COVID-19 vaccine, referred to as COMIRNATY where approved for use under full or conditional marketing authorization, in territories controlled by BioNTech's collaboration partners, particularly for those figures that are derived from preliminary estimates provided by BioNTech's partners; the rate and degree of market acceptance of BioNTech's COVID-19 vaccine and, if approved, BioNTech's investigational medicines; expectations regarding anticipated changes in COVID-19 vaccine demand, including changes to the ordering environment and expected regulatory recommendations to adapt vaccines to address new variants or sublineages; the initiation, timing, progress, results, and cost of BioNTech's research and development programs, including BioNTech's current and future preclinical studies and clinical trials, including statements regarding the expected timing of initiation, enrollment, and completion of studies or trials and related preparatory work and the availability of results, and the timing and outcome of applications for regulatory approvals and marketing authorizations; BioNTech's expectations regarding potential future commercialization in oncology, including goals regarding timing and indications; the targeted timing and number of additional potentially registrational trials, and the registrational potential of any trial BioNTech may initiate; discussions with regulatory agencies; BioNTech's expectations with respect to intellectual property; the impact of BioNTech's collaboration and licensing agreements; the development, nature and feasibility of sustainable vaccine production and supply solutions; the deployment of AI across BioNTech's preclinical and clinical operations; BioNTech's expectations with respect to tariff policy; BioNTech's estimates of revenues, research and development expenses, selling, general and administrative expenses, and capital expenditures for operating activities; BioNTech's expectations regarding upcoming payments relating to litigation settlements; BioNTech's expectations for upcoming scientific and investor presentations; and BioNTech's expectations of net profit / (loss). In some cases, forward-looking statements can be identified by terminology such as "will," "may," "should," "expects," "intends," "plans," "aims," "anticipates," "believes," "estimates," "predicts," "potential," "continue," or the negative of these terms or other comparable terminology, although not all forward-looking statements contain these words.

The forward-looking statements in this presentation are based on BioNTech's current expectations and beliefs of future events and are neither promises nor guarantees. You should not place undue reliance on these forward-looking statements because they involve known and unknown risks, uncertainties, and other factors, many of which are beyond BioNTech's control, and which could cause actual results to differ materially and adversely from those expressed or implied by these forward-looking statements. These risks and uncertainties include, but are not limited to: the uncertainties inherent in research and development, including the ability to meet anticipated clinical endpoints, commencement and/or completion dates for clinical trials, projected data release timelines, regulatory submission dates, regulatory approval dates and/or launch dates, as well as risks associated with preclinical and clinical data, including the data discussed in this release, and including the possibility of unfavorable new preclinical, clinical or safety data and further analyses of existing preclinical, clinical or safety data; the nature of the clinical data, which is subject to ongoing peer review, regulatory review and market interpretation; BioNTech's pricing and coverage negotiations regarding its COVID-19 vaccine with governmental authorities, private health insurers and other third-party payors; the future commercial demand and medical need for initial or booster doses of a COVID-19 vaccine; the impact of tariffs and escalations in trade policy; competition from other COVID-19 vaccines or related to BioNTech's other product candidates, including those with different mechanisms of action and different manufacturing and distribution constraints, on the basis of, among other things, efficacy, cost, convenience of storage and distribution, breadth of approved use, side-effect profile and durability of immune response; the timing of and BioNTech's ability to obtain and maintain regulatory approval for its product candidates; the ability of BioNTech's COVID-19 vaccines to prevent COVID-19 caused by emerging virus variants; BioNTech's and its counterparties' ability to manage and source necessary energy resources; BioNTech's ability to identify research opportunities and discover and develop investigational medicines; the ability and willingness of BioNTech's third-party collaborators to continue research and development activities relating to BioNTech's development candidates and investigational medicines; the impact of COVID-19 on BioNTech's development programs, supply chain, collaborators and financial performance; unforeseen safety issues and potential claims that are alleged to arise from the use of products and product candidates developed or manufactured by BioNTech; BioNTech's and its collaborators' ability to commercialize and market BioNTech's COVID-19 vaccine and, if approved, its product candidates; BioNTech's ability to manage its development and related expenses; regulatory and political developments in the United States and other countries; BioNTech's ability to effectively scale its production capabilities and manufacture its products and product candidates; risks relating to the global financial system and markets; and other factors not known to BioNTech at this time. You should review the risks and uncertainties described under the heading "Risk Factors" in BioNTech's Report on Form 6-K for the period ended March 31, 2025, and in subsequent filings made by BioNTech with the SEC, which are available on the SEC's website at www.sec.gov. These forward-looking statements speak only as of the date hereof. Except as required by law, BioNTech disclaims any intention or responsibility for updating or revising any forward-looking statements contained in this presentation in the event of new information, future developments or otherwise.

Furthermore, certain statements contained in this presentation relate to or are based on studies, publications, surveys and other data obtained from third-party sources and BioNTech's own internal estimates and research. While BioNTech believes these third-party sources to be reliable as of the date of this presentation, it has not independently verified, and makes no representation as to the adequacy, fairness, accuracy or completeness of, any information obtained from third-party sources. In addition, any market data included in this presentation involves assumptions and limitations, and there can be no guarantee as to the accuracy or reliability of such assumptions. While BioNTech believes its own internal research is reliable, such research has not been verified by any independent source. In addition, BioNTech is the owner of various trademarks, trade names and service marks that may appear in this presentation. Certain other trademarks, trade names and service marks appearing in this presentation are the property of third parties. Solely for convenience, the trademarks and trade names in this presentation may be referred to without the ® and TM symbols, but such references should not be construed as any indicator that their respective owners will not assert, to the fullest extent under applicable law, their rights thereto.

An abbreviation directory of defined terms can be found at the end of the presentation.


1 1st Quarter 2025 Update
Ugur Sahin, Co-founder & Chief Executive Officer
2 Oncology Pipeline Update
Özlem Türeci, Co-founder & Chief Medical Officer
3 Financial Update
Jens Holstein, Chief Financial Officer
4 Strategic Outlook
Ryan Richardson, Chief Strategy Officer
BIONTECH


1
1st Quarter 2025 Updates
Ugur Sahin, Co-founder & Chief Executive Officer
BIONTECH


Building a
Global Immunotherapy Powerhouse
Translating Science into Survival
BIONTECH

Progress in Q1 2025 Towards Our Strategic Goals

Execution in Oncology	
BNT327	Presented Phase 2 data[1] for BNT327 in 1L SCLC Reported first BNT327+ADC combo data[1] with TROP2- targeting ADC, BNT325/DB-1305[2]
mRNA Cancer Immunotherapies	Reported Phase 1 data[1] for BNT116[3] in NSCLC Published two manuscripts for autogene cevumeran[4] in Nature and Nature Medicine
BNT323/DB-1303[2]	Preparing for regulator discussions with planned BLA submission by end of 2025, pending regulatory feedback
COVID-19 Leadership	
COMIRNATY	Maintained >50% global COVID-19 vaccine[5] market share
Corporate Update	
Corporate Development	Completed acquisition of Biotheus, securing global control of BNT327 Appointed Ramón Zapata to Management Board as Chief Financial Officer effective July 1, 2025
Financials	Strong balance sheet : **~€ 15.9 bn** total cash and cash equivalents plus security investments[6]

1. Phase 1 data for BNT116 (AACR); BNT327 data included: Phase 2 data in ES-SCLC and SCLC (ELCC) and Phase 1/2 data (AACR); 2. Partnered with DualityBio; 3. In collaboration with Regeneron; 4. Partnered with Genentech, a member of the Roche Group; 5. Partnered with Pfizer; 6. Cash and cash equivalents plus security investments as of March 31, 2025, reached €15,854.4 million, comprising €10,184.9 million cash and cash equivalents, €3,542.0 million current security investments and €2,127.5 million non-current security investments, respectively. A settlement payment of $400 million related to a contractual dispute with the University of Pennsylvania is expected to be reflected in the Company's second quarter 2025 financial results. In connection with this and another settlement with the NIH, BioNTech expects to be reimbursed approximately $535 million by its collaboration partner during 2025 and 2026. Reimbursement payments have begun to be received in the first quarter of 2025.

BIONTECH

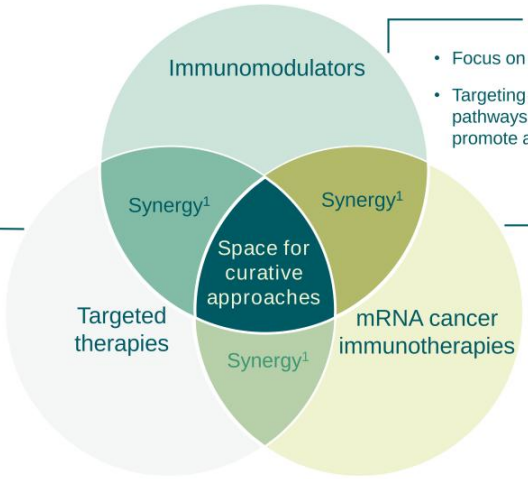
We are Uniquely Positioned to Combine Approaches to Transform Cancer Care

Immunomodulators

- Focus on the critical IO pathways
- Targeting different complementary pathways in cancer immunity cycle may promote a durable anti-tumor effect

Targeted therapies

- Precise and potent modalities for fast onset tumor reduction
- ADC as potential “augmenters” of immunomodulators and mRNA cancer immunotherapies
- Focus on HER2, HER3, TROP2, B7H3 ADCs as combination partners



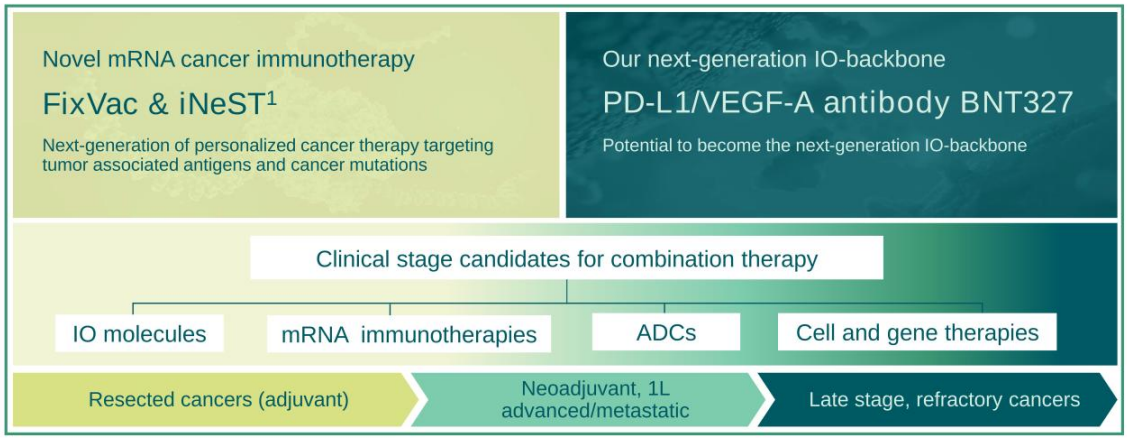
mRNA cancer immunotherapies

- Eliminate polyclonal residual disease with multi-antigen and individualized approaches
- Polyspecific activity by targeting multiple antigens at once
- Establish long-lasting immunological memory to prevent relapses

1. Synergistic potential.

BIONTECH

Our Priorities are Novel mRNA Cancer Immunotherapy and Next-Generation IO-Backbone



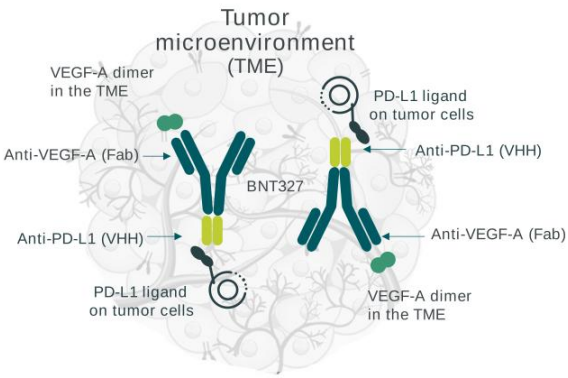
1. Partnered with Genentech, a member of the Roche Group.


2
Oncology Pipeline Update
Özlem Türeci, Co-founder & Chief Medical Officer
BIONTECH

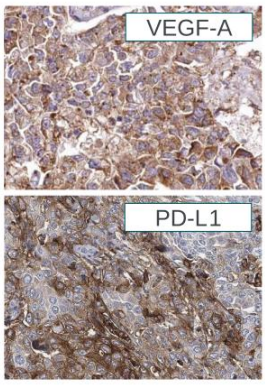
Advancing Towards Commercial Stage in Oncology

1. Partnered with DualityBio.

BNT327: Synergistic Targeting of PD-L1 and VEGF



NSCLC IHC[1]



Bispecific MOA

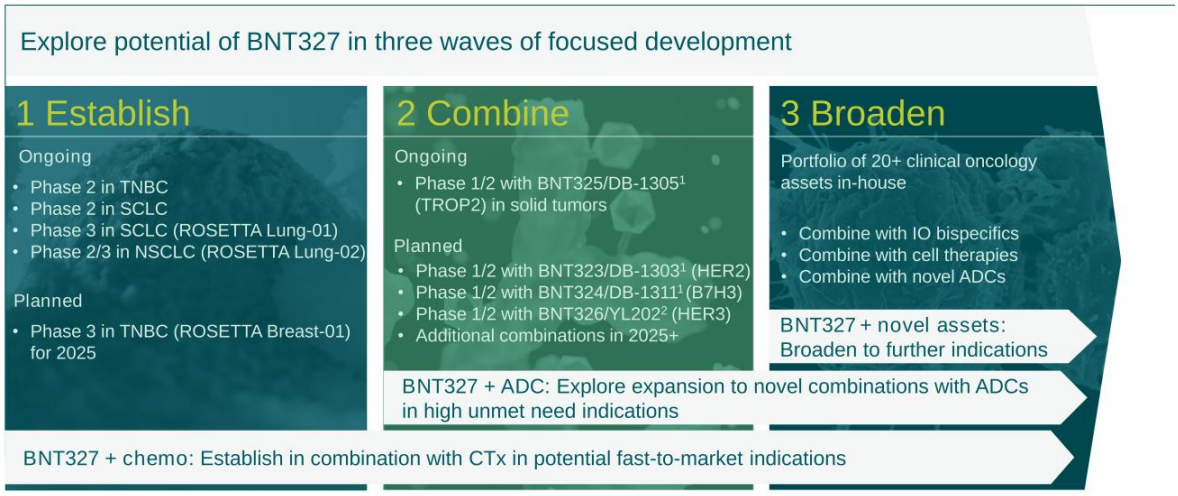
Local neutralization of angiogenic and immunosuppressive VEGF-A effects

Targeting the TME and blockade of PD-1/PD-L1 signaling

1. IHC data: Human Protein Atlas.

BIONTECH

Accelerating BNT327 Global Clinical Development



Partnered with: 1. DualityBio; 2. MediLink.

BIONTECH

BNT327 Combined With Chemotherapy Indicated Encouraging Efficacy in 1L TNBC Irrespective of PD-L1 Status in Phase 1/2 Study

Unmet medical need remains high for patients with TNBC

Patients with Stage IV TNBC[1] have a 5-year survival rate of 10%

Phase 1/2 Study (NCT05918133): Interim overall survival
Jiong Wu et al. presented at SABCS 2024

ITT population (n=42)	
Confirmed ORR (95% CI)	73.8 % (58.0, 86.1)
Median PFS (95% CI)	13.5 months (9.4, 19.3)
12-month OS rate (95% CI)	80.8 % (65.3, 89.9)
18-month OS rate (95% CI)	69.7 % (52.7, 81.6)

A manageable safety profile was observed, with no new safety signals beyond those typically described for nab-paclitaxel and anti- PD-1/PD-L1 and anti-VEGF monotherapies.

Benchmark[2] comparator data by PD-L1 expression level (Keynote-355)
Cortes, J, et al., New England Journal of Medicine, 2022

	1L TNBC (CPS <10)[4,5]	1L TNBC (CPS ≥ 10)
Benchmark regimen	Chemo	Pembro + Chemo
ORR	35 %	53 %
Median PFS	5.7 months	9.7 months
Median OS	15.2 months	23.0 months

The above data are not based on a head-to-head study comparing BioNTech's investigational products with other products/candidates - no conclusions can be drawn.

We believe BNT327 has the potential to become a first-line treatment option for patients with TNBC[3], including those currently not addressed by existing IO therapies

1. Incidence from SEER (US); Zentrum für Krebsregisterdaten (DE); Globocan (ES); Sante Publique (FR); AIOM (IT); Cancer Research UK . 2 Benchmark study: KEYNOTE-355 as reported in Cortes, J, et al. New England Journal of Medicine,. 2022. 3. The above information is not based on head-to-head trials between BioNTech's investigational candidates and other products or product candidates. Furthermore, definitive conclusions cannot be drawn from cross-trial comparisons or anticipated data, as they may be confounded by various factors, and should be interpreted with caution. 4. Obtained from subgroup analysis. 5. mPFS for CPS < 10 subgroup from Cortes, J. et al. Lancet, 2020.

BIONTECH

BNT327 Combined With Chemotherapy Indicated Encouraging Efficacy in 1L ES-SCLC in Phase 2 Study

Unmet medical need remains high for patients with ES-SCLC

Patients with ES-SCLC[1] have a 5-year survival rate of 3%

Phase 2 Study (NCT05844150): Emerging efficacy profile
Ying Cheng et al. presented at ELCC 2025

ITT population (n=48)	
Confirmed ORR (95% CI)	85.4 % (72.2, 93.9)
Median PFS (95% CI)	6.9 months (4.34, 8.21)
Median OS (95% CI)	16.8 months (14.3, --)
OS events, n (%)	17 (35.4)
12-month OS rate (95% CI)	72.7 % (57.6, 83.1)

A manageable safety profile was observed, with no new safety signals beyond those typically described for chemotherapy agents and anti-PD-(L)1 and anti-VEGF monotherapies.

Benchmark[2] comparator data (IMpower133)
L. Horn et al., New England Journal of Medicine, 2018

	1L ES-SCLC
Benchmark regimen	Atezo + Chemo
ORR	60%
Median PFS	5.2 months
Median OS	12.3 months

The above data are not based on a head-to-head study comparing BioNTech's investigational products with other products/candidates - no conclusions can be drawn.

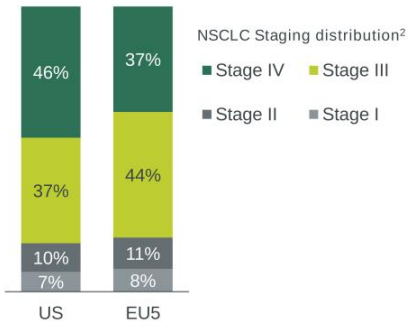
We believe BNT327 has the potential to become a new first-line treatment option for patients with ES-SCLC[3]

1. Incidence from: SEER data for diagnosed SCLC incidence in US; Cancer Research UK; Zentrum für Krebsregisterdaten; Sante Publique; AIOM; EPDATA. 2. Benchmark study: IMpower133 as reported in L. Horn et al., New England Journal of Medicine, 2018 3. The above information is not based on head-to-head trials between BioNTech's investigational candidates and other products or product candidates. Furthermore, definitive conclusions cannot be drawn from cross-trial comparisons or anticipated data, as they may be confounded by various factors, and should be interpreted with caution.

BIONTECH

Non-Small Cell Lung Cancer is One of the Highest Incidence Cancers Globally[1]

2030 U.S., EU4, U.K. NSCLC projected incidence[1] **~415k**



Treatment outcomes vary based on histology and PD-L1 levels in 1L NSCLC patients without actionable genomic alterations

	Non-squamous (~ 70%)[3]	Squamous (~ 30%)[3]
PD-**L1 ≥ 50%** (~ 25 - 30%)[4,5]	5-year OS: 30% (KN-189)[6]	5-year OS: 23% (KN-407)[7]
PD-L1 1 - 49% (~ 30 - 40%)[4,5]	5-year OS: 20% (KN-189)[6]	5-year OS: 21% (KN-407)[7]
PD-L1 < 1% (~ 30 - 40%)[4,5]	5-year OS: 10% (KN-189)[6]	5-year OS: 11% (KN-407)[7]

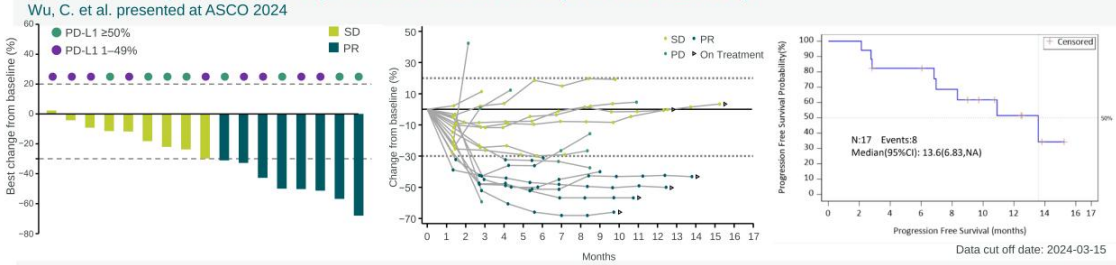
1. Globocan – Cancer Tomorrow; 2. CancerMPact® 2024 Treatment Architecture EU5 and US; Note that 5-year survival reported includes all comer NSCLC population ie including with actionable genetic alterations; 3. Ganti AK, et al., Update of Incidence, Prevalence, Survival, and Initial Treatment in Patients With Non-Small Cell Lung Cancer in the US. JAMA Oncology, 2021 Dec; 4. Mansour MSI et al., PD-L1 Expression in Non-Small Cell Lung Cancer Specimens: Association with Clinicopathological Factors and Molecular Alterations. International Journal of Molecular Sciences, 2022 Apr 19;23(9):4517; 5. Saez de Gordoa, K. et al. PD-L1 Expression in Non-Small Cell Lung Cancer: Data from a Referral Center in Spain. Diagnostics 2021, 11, 1452; 6. Garassino MC, et al. Pembrolizumab Plus Pemetrexed and Platinum in Nonsquamous Non-Small-Cell Lung Cancer: 5-Year Outcomes From the Phase 3 KEYNOTE-189 Study. Journal of Clinical Oncology. 2023 Apr 10;41(11):1992-1998; 7. Silvia Novello et al., Pembrolizumab Plus Chemotherapy in Squamous Non–Small-Cell Lung Cancer: 5-Year Update of the Phase III KEYNOTE-407 Study. Journal of Clinical Oncology, 41, 1999-2006(2023).

BIONTECH

BNT327 Indicates Single Agent Activity in 1L NSCLC in Phase 1b/2a Study

Phase 1b/2a (NCT05918445); Cohort 1: 1L NSCLC (EGFR & ALK WT)

Wu, C. et al. presented at ASCO 2024



Data cut off date: 2024-03-15

BNT327 indicated manageable safety in this patient population. Safety events were consistent with those described for anti-PD-L1 and anti-VEGF monotherapy.

Benchmark[1] comparator data

Indication	Benchmark[2] regimen	ORR	mPFS	mOS
1L NSCLC (PD-L1 ≥ 50%)	Pembrolizumab monotherapy	45%	7.7 months	26.3 months

The above data are not based on a head-to-head study comparing BioNTech's investigational products with other products/candidates - no conclusions can be drawn.

1L NSCLC mono tx (cohort 1, n=17): ORR 47%, DCR 100%, mPFS 13.6 months
Comparable ORR in PD-L1 1–49% (n=9) and PD-L1 ≥50% (n=8)

1. Benchmark study: KEYNOTE-024 as reported in Reck, M. et al. New England Journal of Medicine, 2016; 2. The above information is not based on head-to-head trials between BioNTech's investigational candidates and other products or product candidates. Furthermore, definitive conclusions cannot be drawn from cross-trial comparisons or anticipated data, as they may be confounded by various factors, and should be interpreted with caution

BIONTECH

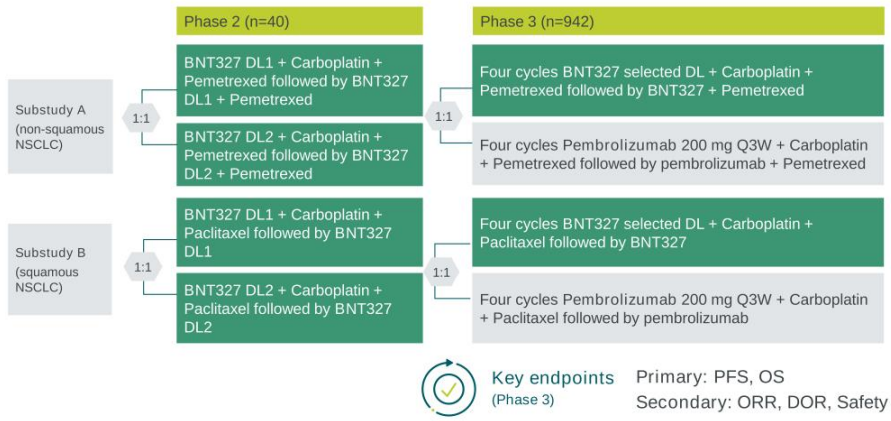
ROSETTA Lung-02 – Our First Global Registrational Trial in NSCLC

ROSETTA LUNG-02

A Phase 2/3, multisite, randomized global trial of BNT327 in combination with chemotherapy in first-line non-small cell lung cancer (NCT06712316)

Key eligibility criteria
- Treatment naïve Stage IIIB/IIIC or IV NSCLC
- RECIST 1.1 measurable disease
- ECOG PS 0 or 1
- PD-L1 all-comers

Stratification factors Phase 3
- PD-L1 (negative, 1-49%, ≥50%)
- Brain metastasis (Y/N)
- Region (East Asia vs RoW)



Key endpoints (Phase 3)
Primary: PFS, OS
Secondary: ORR, DOR, Safety

BIONTECH

BioNTech at ASCO 2025

2025 ASCO ANNUAL MEETING

- Across portfolio
- Data for making informed decisions about the direction of further development

Related Program	Indication	Content
BNT327	1L NSCLC	Phase 2/3 TiP (ROSETTA Lung-02)
BNT327	1L SCLC	Phase 3 TiP (ROSETTA Lung-01)
BNT327	1L Mesothelioma	Phase 2 data
BNT316[1]	2L+ Melanoma	Phase 2 data
BNT316[1]	2L+ CRPC	Phase 1 data
BNT324/DB-1311[2]	2L+ CRPC	Phase 2 data
BNT142	CLDN6+ Solid Tumors	Phase 1/2 data

1. Partnered with 1. OncoC4; 2. DualityBio.


3
Financial Update
Jens Holstein, Chief Financial Officer
BIONTECH

Q1 2025 Financial Results

(in millions €, except per share data)[1]	Three months ended March 31 2025	2024
Total Revenues	183	188
Cost of sales	(84)	(59)
Research and development expenses	(526)	(508)
Sales and marketing expenses	(14)	(16)
General and administrative expenses	(107)	(117)
Other operating result	14	5
Operating loss	(534)	(507)
Finance result	89	175
Income taxes	29	17
Net loss	(416)	(315)
Loss per share		
Basic and diluted loss per share	(1.73)	(1.31)

Balance Sheet as of March 31, 2025	
Cash and cash equivalents plus security investments[2]	€15.9 bn

1. Numbers have been rounded; numbers presented may not add up precisely to the totals and may have been adjusted in the table. Presentation of the consolidated statements of profit or loss has been condensed. More information can be found in BioNTech's Report on Form 6-K for the three months ended March 31, 2025, filed today with the United States Securities and Exchange Commission and available at https://www.sec.gov/; 2. Cash and cash equivalents plus security investments as of March 31, 2025, reached €15,854.4 million, comprising €10,184.9 million cash and cash equivalents, €3,542.0 million current security investments and €2,127.5 million non-current security investments, respectively. A settlement payment of $400 million related to a contractual dispute with the University of Pennsylvania is expected to be reflected in the Company's second quarter 2025 financial results. In connection with this and another settlement with the NIH, BioNTech expects to be reimbursed approximately $535 million by its collaboration partner during 2025 and 2026. Reimbursement payments have begun to be received in the first quarter of 2025.

2025 Financial Year Guidance Confirmed[1]

		FY 2025 Guidance
Planned FY 2025 revenues	Total revenues	**€1,700 – €**2,200 m
Planned FY 2025 expenses and capex[4]	R&D expenses	**€2,600 – €2,800** m
	SG&A expenses	**€650 – €750 m**
	Capital expenditure for operating activities	**€250 – €350 m**
Guidance considerations	• Our revenue guidance assumes relatively stable vaccination rates, pricing and market share as compared to 2024. We also anticipate a revenue phasing similar to 2024 with the last 3-4 months driving the full year revenue figure. However, potential changes to the law or governmental policy, including tariffs and public health policy, and evolving public sentiment worldwide, could further negatively impact our anticipated revenues and expenses. • Inventory write-downs and other charges are estimated to be ~15% of BioNTech's share of gross profit from COVID-19 vaccines sales in Pfizer's territory • Anticipated revenues related to service businesses include InstaDeep, JPT Peptide and IMFS as well as revenues from the German pandemic preparedness agreement	

1. Financial guidance excludes external risks that are not yet known and/or quantifiable, including, but not limited to the effects of ongoing and/or future legal disputes and related activities, certain potential one-time effects and charges related to portfolio prioritization. It includes effects identified from licensing arrangements, collaborations or potential M&A transactions to the extent disclosed and may be subject to update. The Company does not expect to report a positive net income figure for the 2025 financial year.

4 Strategic Outlook

Ryan Richardson, Chief Strategy Officer

Strategic Priority Areas in 2025

mRNA Cancer Immunotherapy

- Expect first randomized data in the adjuvant setting (CRC)
- Continue to execute 7 ongoing Phase 2 trials and first novel combination trials

BNT327

- Advance 3 global registration-enabling trials in potential fast-to-market indications
- Generate first BNT327+ ADC combination datasets

Commercial Readiness in Oncology

- Advance BNT323/DB-1303[1] towards BLA submission
- Continue to build targeted AI-enabled commercialization team in key markets

COVID-19 Vaccine[2]

- Maintain global COVID-19 vaccine market leadership
- Advance next-gen and combination vaccine programs

Partnered with: 1. DualityBio; 2. Pfizer.


BIONTECH
Save the date
Annual General Meeting
May 16, 2025
Innovation Series R&D Day
November 11, 2025
BIONTECH

Thank you

Appendix

Selected Pipeline Milestones in 2025 and Beyond

	Program	Indication	2025+ Milestone
Next-generation immunomodulator	BNT327	1L SCLC	China Phase 2 data
		1L/2L SCLC	Global Phase 2 dose optimization data
		1L/2L TNBC	Global Phase 2 dose optimization data
	BNT327 + BNT325/DB-1305[1]	Solid tumors	Global Phase 1 data
mRNA cancer immunotherapy	Autogene cevumeran (BNT122 / RO7198457)[2]	ctDNA+ adj. CRC	Phase 2 data
	BNT111[3]	2L+ melanoma	Phase 2 data
	BNT116[3]	PD-L1 > 1% NSCLC	Phase 1 data
Targeted therapy	BNT323[1]	2L+ HER2 EC	Phase 2 data
			Regulatory submission

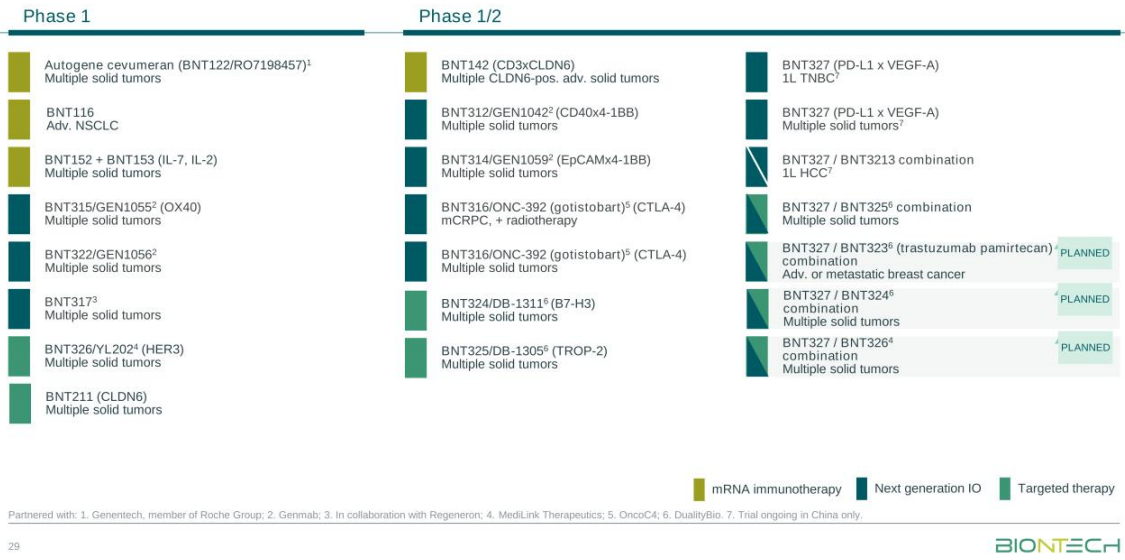
Partnered with: 1. DualityBio; 2. Genentech, a member of the Roche Group; 3. In collaboration with Regeneron.

BioNTech's Oncology Pipeline – Phase 2 and Phase 3 Clinical Trials

Partnered with: 1. Genentech, member of Roche Group; 2. Genmab; 3. MediLink Therapeutics; 4. OncoC4; 5. DualityBio. 6. In collaboration with Regeneron; 7. Trial ongoing in China only.

BIONTECH

BioNTech's Oncology Pipeline – Phase 1 and Phase 1/2 Clinical Trials



Partnered with: 1. Genentech, member of Roche Group; 2. Genmab; 3. In collaboration with Regeneron; 4. MediLink Therapeutics; 5. OncoC4; 6. DualityBio. 7. Trial ongoing in China only.

BIONTECH

Abbreviation Directory

n L	nth line	EU4(5)	Includes Germany, France, Italy, Spain (UK)	OX40	CD134
AACR	American Association for Cancer Research	Fab	Fragment antigen binding	PD	Progressive disease
ADC	Antibody-drug conjugate	FixVac	Fixed Antigen Vaccine	PDAC	Pancreatic ductal adenocarcinoma
adj.	Adjuvant	FY	Fiscal year	PD-(L)1	Programmed cell death protein (ligand) 1
AI	Artificial intelligence	HCC	Hepatocellular carcinoma	PFS	Progression-free survival
AIOM	Associazione Italiana di Oncologia Medica	HER2 (or 3)	Human epidermal growth factor receptor 2 (or 3)	PR	Partial response
ALK	Anaplastic large-cell lymphoma kinase	HNC	Head and neck cancer	PROC	Platinum-resistant ovarian cancer
ASCO	American Society of Clinical Oncology	HPV	Human papilloma virus	QxW	Every x week(s)
BLA	Biologics License Applications	HR	Hormone receptor	R&D	Research and development
CAPEX	Capital expenditures	IHC	Immunohistochemistry	RECIST	Response Evaluation Criteria in Solid Tumors
CD-x	Cluster of differentiation	IMFS	BioNTech Innovative Manufacturing Services	RoW	Rest of world
CI	Confidence interval	iNeST	Individualized NeoAntigen-Specific Therapy	R/R	Relapsed/refractory
CLDN6	Claudin 6	IO	Immuno-oncology	SABCS	San Antonio Breast Cancer Symposium
CPS	Combined positive score	ITT	Intention to treat	(ES)SCLC	(Extensive stage) small cell lung cancer
CRC	Colorectal cancer	JAMA	Journal of the American Medical Association	SD	Stable disease
CRPC	Castration resistant prostate cancer	m	Median	SEC	U.S. Securities and Exchange Commission
ctDNA	Circulating tumor DNA	M&A	Merger and acquisitions	SEER	Surveillance, epidemiology, and end results
CTLA	Cytotoxic T-lymphocyte-associated protein	MIUC	Muscle-invasive urothelial carcinoma	SG&A	Selling, general and administrative expenses
CTx	Chemotherapy	MOA	Mechanism of Action	TKI	Tyrosine kinase inhibitor
DCR	Disease control rate	MPM	Malignant pleural mesothelioma	TME	Tumor microenvironment
DL	Dose level	mRNA	Messenger ribonucleic acid	TNBC	Triple-negative breast cancer
DOR	Duration of response	NCT	National clinical trial	TROP2	Trophoblast cell-surface antigen 2
EC	Endometrial cancer	NEN	Neuroendocrine neoplasm	UK	United Kingdom
ECOG	Eastern Cooperative Oncology Group	NIH	National Institutes of Health	U.S.	United States
EGFR	Epidermal growth factor receptor	NSCLC	Non-small cell lung cancer	VEGF-A	Vascular endothelial growth factor A
ELCC	European Lung Cancer Congress	ORR	Objective response rate	VHH	Heavy chain variable
EpCAM	Epithelial cell adhesion molecule	OS	Overall survival	WT	Wild type