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 2026

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 15, 2026, BioNTech SE held the Annual General Meeting ("AGM") 2026. The press release and the AGM presentation are attached as Exhibits 99.1 and 99.2, respectively. The voting results are attached as Exhibit 99.3.

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/ Ramón Zapata Gomez
Name: Ramón Zapata Gomez
Title: Chief Financial Officer

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

Date: May 15, 2026

EXHIBIT INDEX

Exhibit	Description of Exhibit
99.1	Press Release: BioNTech SE Shareholders Approve All Agenda Items at the Annual General Meeting 2026
99.2	Annual General Meeting 2026 Presentation
99.3	Annual General Meeting 2026 Voting Results

BioNTech SE Shareholders Approve All Agenda Items at the Annual General Meeting 2026

MAINZ, Germany, May 15, 2026 (GLOBE NEWSWIRE) -- BioNTech SE (Nasdaq: BNTX, “BioNTech” or “the Company”) held its Annual General Meeting (“AGM”) today. A total of 92 per cent of the share capital was represented at the virtual assembly. All resolutions proposed on the agenda items put to the vote at today’s AGM were approved by a majority of the shareholders.

BioNTech is strengthening its focus on the Company’s growing late-stage oncology pipeline, while continuing its discovery and early research aimed at long-term innovation. Consequently, the Company's shareholders approved expanding the Supervisory Board from six to eight members and adding additional expertise: Prof. Iris Löw-Friedrich, M.D., Ph.D., and Susanne Schaffert, Ph.D., were elected as new members of the Supervisory Board.

Iris Löw-Friedrich has many years of expertise in the field of clinical development and broad experience in the scientific and medical fields. She also possesses knowledge in the areas of sales and commercialization, management, innovation, and international markets relevant to the Company.
She is an experienced Supervisory Board member and adjunct professor of internal medicine at the faculty of medicine at Goethe University in Frankfurt am Main, Germany.

Susanne Schaffert is a member of supervisory boards in the healthcare sector, including that of Merck KGaA. She possesses particular expertise in the field of oncology as well as in sales and commercialization with a focus on product launches. Her knowledge spans innovation, research and development, and organizational leadership and management.

Additionally, shareholders approved the extension of the mandates of BioNTech’s Supervisory Board members Helmut Jeggle, Prof. Anja Morawietz, Ph.D., and Prof. Rudolf Staudigl, Ph.D.

At a meeting held following the AGM, the Supervisory Board elected Helmut Jeggle as its Chairman.

The voting results for all agenda items can be viewed on the Annual General Meeting 2026 website under the section ‘Voting Results’. The speeches by Chief Executive Officer Prof. Ugur Sahin, M.D., Chief Financial Officer Ramón Zapata and the slides presented at the AGM 2026 can be found in section ‘Speeches and Presentations’ under the same link.

About BioNTech
BioNTech is a global next generation biopharmaceutical company pioneering novel investigative therapies for cancer and other serious diseases. In oncology, BioNTech is committed to transforming how cancer is treated. Its ambition is to develop innovative medicines with pan-tumor or synergistic potential to address cancer from multiple angles and across the full continuum of the disease from early- to late-stage. Its growing late-stage oncology pipeline comprises complementary treatment approaches spanning immunomodulators, antibody drug conjugates, and mRNA cancer immunotherapies. BioNTech has partnered with multiple global and specialized pharmaceutical collaborators leveraging complementary expertise and resources to accelerate innovation and drive progress, including Bristol Myers Squibb, Duality Biologics, Genentech, a member of the Roche Group, Genmab, MediLink, OncoC4, and Pfizer.

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

BioNTech 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 the potential benefits of appointed Supervisory Board members. 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.

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, 2026, 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

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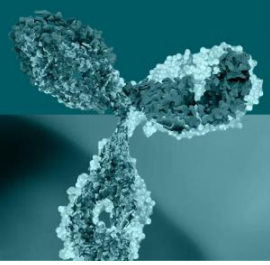
Media@biontech.de


Exhibit 99.2
Annual General Meeting
15 May 2026, 14:00 CEST
BIONTECH

Management Report



1 Operations Development 2025 & Q1 2026 and Outlook 2026
Prof. Dr. Uğur Şahin, Co-Founder & Chief Executive Officer

2 Financial Development 2025 & Q1 2026 and Financial Outlook 2026
Ramón Zapata, Chief Financial Officer

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1

Operations Development 2025 & Q1 2026 and Outlook 2026

Prof. Dr. Uğur Şahin, Co-Founder & Chief Executive Officer

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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; 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; BioNTech's expectations regarding the impact of changes to its manufacturing operations; discussions with regulatory agencies; BioNTech's expectations with respect to intellectual property; the impact of BioNTech's collaboration and licensing agreements, including BioNTech's partnership with BMS; BioNTech's expectations with respect to developments in law, public policy, and international trade; BioNTech's estimates of revenues, research and development expenses, selling, general and administrative expenses, and capital expenditures for operating activities; 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 presentation, 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 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; 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 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; 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 its product candidates, if approved; 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, 2026, 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.

Continuing to Execute on BioNTech's Strategy

COVID-19 Vaccine Global Impact

5 Billion vaccine doses distributed[1]

Pfizer

Addressing Oncology Unmet Medical Need

>25 Ongoing Phase 2 & Phase 3 trials[2]

17 Clinical programs[3]

Bristol Myers Squibb · Genentech (A Member of the Roche Group) · Genmab · OncoC4 · MediLink Therapeutics · DualityBio

Focused Infectious Disease Innovation

7 High unmet need clinical programs[4]

Gates Foundation · CEPI

In-House GMP Manufacturing Platforms

Capabilities and facilities for key platforms: mRNA therapeutics, including individualized mRNA and bispecific antibodies

Fully Integrated AI-Driven Innovation

Tech-bio company with AI-infused target and drug discovery and development capabilities

InstaDeep™ a BioNTech Company

1. Includes globally distributed doses from 2020 to-date; 2. Includes Phase 2 or 3 trials for BNT113, autogene cevumeran, gotistobart, trastuzumab pamirtecan and pumitamig; 3. Includes BNT113, BNT116, autogene cevumeran, BNT211, BNT314/GEN1059, gotistobart, BNT317, trastuzumab pamirtecan, BNT324/DB-1311, BNT325/DB-1305, BNT326/YL202, pumitamig, BNT329, BNT3212, BNT3213; BNT3214; 4. Includes BNT162, BNT161, BNT163, BNT164, BNT165, BNT166, BNT351; Numbers on the slide are as of 5 May 2026.

2025 Achievements: Strong Performance and Pipeline Momentum

COVID-19 Market Leadership[1]

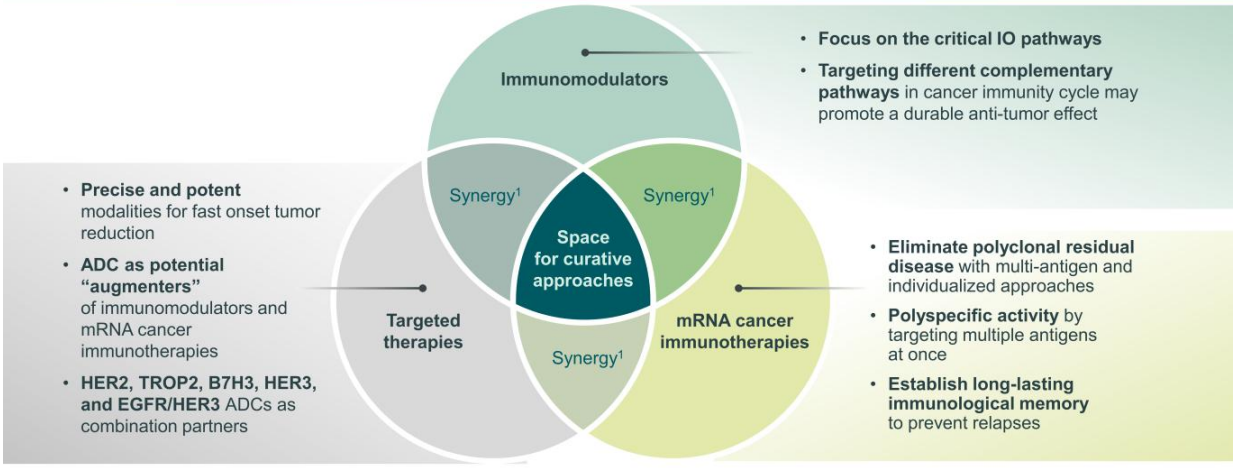
Advanced Key Oncology Programs

Executed Key Strategic Deals

Strengthened Financial Position

1. Over 50%, including Italy, Spain, France, Germany, USA, Japan, Australia.

Strategic Oncology Multi-Modal Immunotherapy Approach



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

- **Precise and potent** modalities for fast onset tumor reduction
- **ADC as potential "augmenters"** of immunomodulators and mRNA cancer immunotherapies
- **HER2, TROP2, B7H3, HER3, and EGFR/HER3** ADCs as combination partners

- **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

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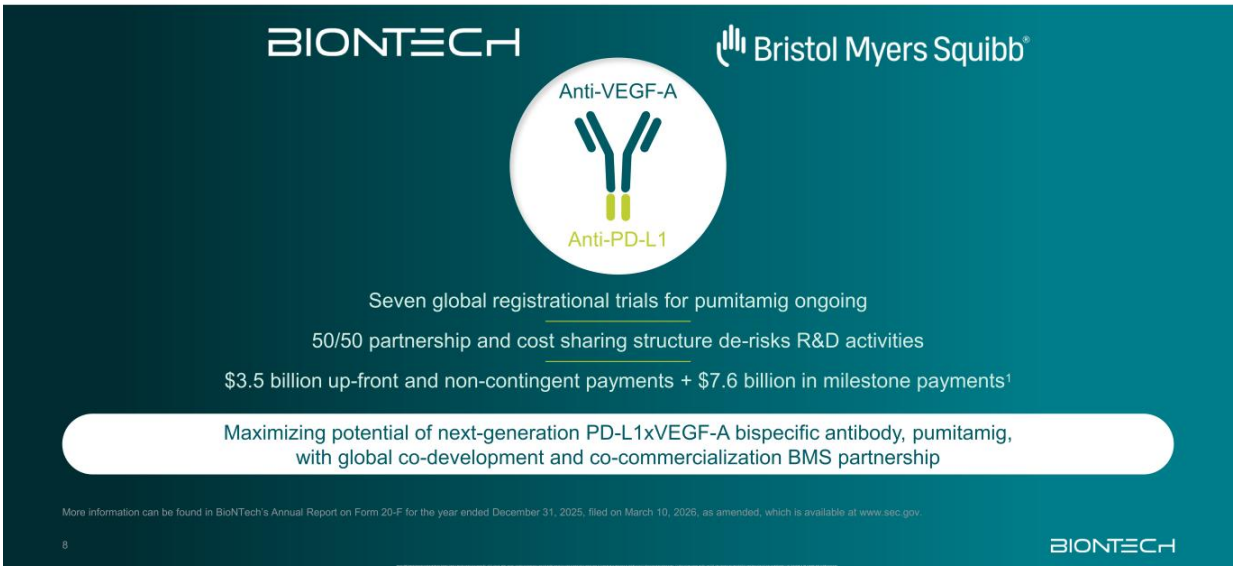
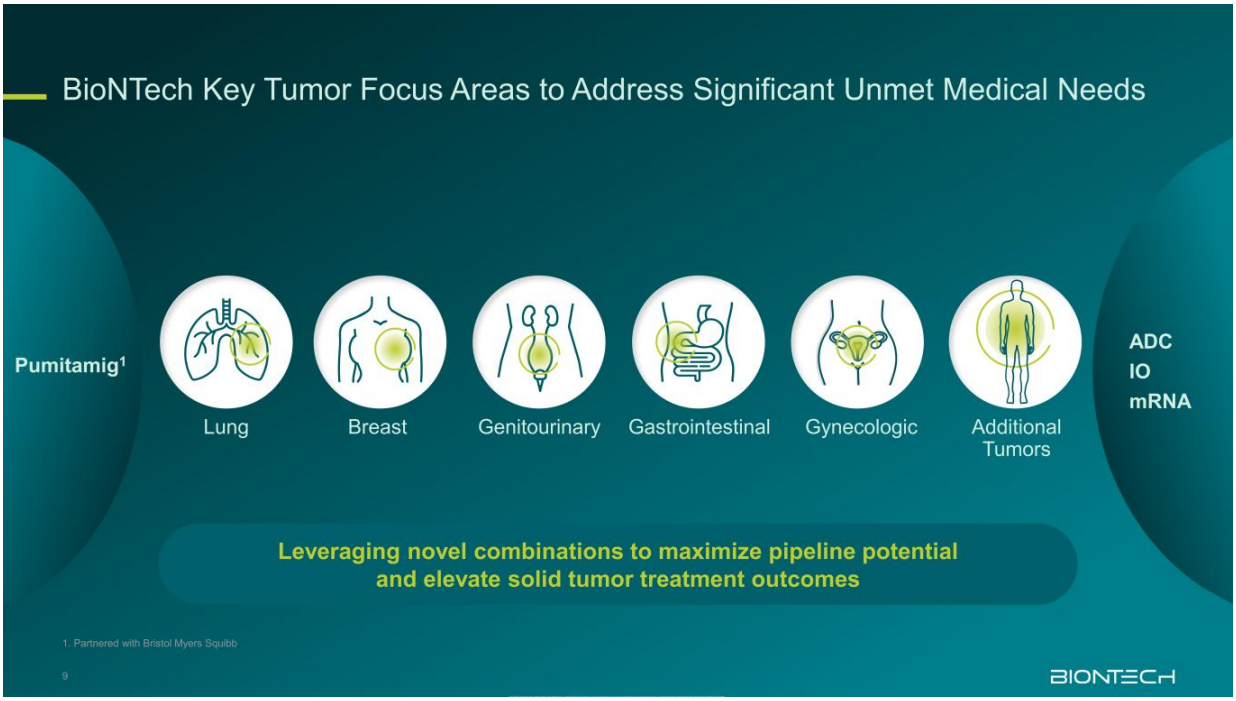
Innovative BMS Partnership Structured to Accelerate and Maximize Pumitamig's Potential




BioNTech Key Tumor Focus Areas to Address Significant Unmet Medical Needs
Pumitamig[1]
Lung
Breast
Genitourinary
Gastrointestinal
Gynecologic
Additional Tumors
ADC
IO
mRNA
Leveraging novel combinations to maximize pipeline potential and elevate solid tumor treatment outcomes
1. Partnered with Bristol Myers Squibb
9
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Building a Multi-Product Company by 2030

Targeting 17+ Late-Stage/Pivotal Trial Readouts Through 2030+ Informing Multiple Launch Opportunities

1. Expected data readouts may be from interim or final analyses and are event-driven, and in some cases may not translate into commercial launches; Partnered with 2. Bristol Myers Squibb; 3. OncoC4; 4. DualityBio; 5. Genentech, a member of the Roche group; 6. These are Phase 1/2 trials. The anticipated pivotal trials evaluating pumitamig in these tumor types are expected to readout after 2030.

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Our vision and mission have not changed since the founding of BioNTech.
Our Vision
We aspire to harness the power of the immune system to develop novel therapies against cancer, infectious diseases and other severe diseases. We believe in scientific rigor, innovation and passion as driving forces.
Our Mission
We are committed to improving the health of people worldwide with our fundamental research and work in the development of immunotherapies.

2

Financial Development 2025 & Q1 2026 and Financial Outlook 2026

Ramón Zapata,
Chief Financial Officer

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Full Year 2025 Financial Results Compared to Guidance

In € millions	FY 2025 IFRS Results[1]	FY 2025 IFRS Guidance
Total Revenues	2,870	2,600 – 2,800
R&D Expenses	2,105	2,000 – 2,200
SG&A Expenses	624	550 – 650
Capital Expenditures for Operating Activities	198	200 – 250

1. All numbers have been rounded. More information can be found in BioNTech's Report on Form 20-F for the year ended December 31, 2025, filed on March 10, 2026, as amended, which is available at www.sec.gov.

First Quarter 2026 Unaudited Financial Results Highlights[1]

118 M€

Revenues

527 M€

Adjusted Research & Development Expenses[2]

(IFRS: 557 M€)

151 M€

Sales, Marketing, General & Administrative Expenses

16.8 B€

Cash and Cash Equivalents plus Security Investments[3]

1. All numbers have been rounded. 2. In addition to BioNTech's results determined in accordance with International Financial Reporting Standards ("IFRS"), or IFRS Accounting Standards, or IFRS results, BioNTech reports certain adjusted, non-IFRS measures used internally as a supplemental measure of our business performance (each referred to with the prefix "Adjusted" or, as a whole, "Adjusted Results"). The calculation of these measures and the adjusted results as a whole is based on the concepts of the applicable IFRS Accounting Standards, but includes certain adjustments. Reconciliation of the adjusted results to BioNTech's measures based on IFRS Accounting Standards and more information can be found in the appendix and in BioNTech's Report on Form 6-K for the period ended March 31, 2026, filed on May 5, 2026, which is available at www.sec.gov. While non-IFRS measures may offer additional insights, BioNTech's non-IFRS measures are not, and should not be viewed as a substitute for their most directly comparable IFRS Accounting Standards measures, and should always be considered alongside our financial statements prepared in accordance with IFRS Accounting Standards. 3. Cash and cash equivalents plus security investments as of March 31, 2026, reached €16,763.3 million, comprising €9,939.4 million in cash and cash equivalents, €4,696.9 million in current security investments disclosed as financial assets and €2,127.0 million in non-current security investments disclosed as financial assets.

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Reaffirming Full Year 2026 Financial Guidance[1]

In € millions	FY 2026 non-IFRS Guidance
Total Revenues	2,000 – 2,300
Adjusted R&D Expenses	2,200 – 2,500
Adjusted SG&A Expenses	700 – 800

Revenue Guidance Considerations

- Competitive market dynamics in the United States
- Begin managing transition away from multi-year contracts in Europe, and specifically in Germany where BioNTech recognizes direct sales for its COVID-19 vaccine
- Stable revenues from the collaboration with BMS, from a pandemic preparedness contract with the German government, and from the BioNTech Group service businesses
- No operationally-driven one-time revenue effect, such as from Pfizer opt-out from further development of shingles program

1. Excludes risks that are not yet known and/or quantifiable and related activities. Includes effects identified from licensing arrangements, collaborations and Merger & Acquisitions ("M&A") transactions to the extent disclosed. The guidance is based on non-IFRS measures and excludes certain effects compared to measures based on IFRS Accounting Standards. More information can be found in BioNTech's Report on Form 6-K for the period ended March 31, 2026, filed on May 5, 2026, which is available at www.sec.gov.

Capital Transactions During Full Year 2025 and First Quarter 2026

Acquisition of CureVac

Issuance of new ADSs or use of ADSs[1] held in treasury	Period	Number of issued ADSs	Percentage of share capital[2]	Issue price or fair value	Volume[3]
First and Second Offer Period (issuance)	December 2025	10,475,287	4.04%	€82.90[5]	€868.4 million
Post-Offer Reorganization (held in treasury)	January 2026	1,552,300	0.60%	€82.90	€128.7 million[4]
Total number of ADSs		**12,027,587**	**4.64%**	**Ø €82.90**	**€997.1 million**

Use of ADSs[1] held in treasury	Period	Number of issued ADSs	Percentage of share capital[2]	Issue price	Volume[3]
CureVac Transaction Award	December 2025	57,211	0.02%	€78.75	€4.5 million
Total number of used ADSs previously held in treasury		**57,211**	**0.02%**	**€78.75**	**€4.5 million**

Employee Participation Programs

Use of ADSs[1] held in treasury	Period	Number of issued ADSs	Percentage of share capital[2]	Issue price	Volume[3]
Biotheus Founder SBP Program	January 2025	421,818	0.16%	€116.58	€49.2 million
ESOP 2018 Settlement	March to December 2025	24,147	0.01%	€91.68	€2.2 million
LTI 2020 Settlement	April to June 2025	636	< 0.01%	€112.10	€0.1 million
InstaDeep Tranche 2 Settlement	July 2025	49,143	0.02%	€93.92	€4.6 million
LTI 2024 Tranche 1 Settlement	December 2025	229,870	0.09%	€81.90	€18.8 million
LTI 2021 Settlement	December 2025	96,424	0.04%	€81.10	€7.8 million
Total number of used ADSs previously held in treasury		**822,038**	**0.32%**	**Ø €100.63**	**€82.7 million**

1. American Depositary Share (ADS), each representing one ordinary share. 2. The "percentage of share capital" ratio is calculated based on the shares issued as of December 31, 2025 and as of March 31, 2026 (both 259,027,487), respectively. 3. Numbers have been rounded. 4. Disclosed as an obligation to issue share capital in BioNTech's Annual Report on Form 20-F as of and for the year ended December 31, 2025. Post-offer reorganization took place in Q1 2026. Withholding tax of €4.0 million was paid during the post-offer reorganization in Q1 2026 in cash ("withhold to cover" for approx. 48,000 ADSs). 5. Fair value of $96.73 per ADS, calculated using the exchange rate of 0.86.

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Capital Transactions During Full Year 2025 and First Quarter 2026

Employee Participation Programs

Use of ADSs[1] held in treasury	Period	Number of issued ADSs	Percentage of share capital[2]	Issue price	Volume[3]
LTI 2021 Settlement	February 2026	169	< 0.01%	€81.10	€0.01 million
ESOP 2018 Settlement	March 2026	6,452	< 0.01%	€78.07	€0.5 million
Total number of used ADSs previously held in treasury		**6,621**	**< 0.01%**	**Ø €78.15**	**€0.51 million**

1. American Depositary Share (ADS), each representing one ordinary share. 2. The "percentage of share capital" ratio is calculated based on the shares issued as of March 31, 2026 (259,027,487). 3. Numbers have been rounded.

Focused Capital Allocation Strategy for Sustainable Value Creation

Focusing R&D Investments

Concentrate investments on advancing BioNTech's growing oncology pipeline toward commercialization, including pumitamig and ADC candidates

Announced Share Repurchase Program

Initiating share repurchase program of up to $1.0 billion over the next twelve months

Manufacturing Footprint Consolidation

Enhance operational efficiency with expected cost savings to ramp up over time, reaching approximately €500 million in recurring annual savings upon full implementation of the measures in 2029[1]

1. Expected savings relative to BioNTech's 2025 cost base and CureVac's 2026 budget; do not reflect partially offsetting costs for CDMO use or transfer to other sites; and exclude exit costs, which will be recorded as incurred.

BioNTech Oncology Vision: Translating Science into Survival

Today

Advanced Strategy, Matured Pipeline and De-risked Development

Progress key programs into pivotal stage, leverage partnership with BMS, fortified balance sheet to fund our pipeline

2026-2029

Drive Oncology Execution at Scale and Speed

Advance combination therapy studies, accelerate pivotal trial execution, build indication-specific oncology portfolios and execute oncology launches

2030

Diversified Multi-Product Company

Build a diversified, multi-product global immunotherapy powerhouse addressing high unmet medical need of cancer patients worldwide

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Thank you
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Reconciliation of IFRS to Adjusted Results – Q1 2026 & 2025 Financial Results

In € millions except per share data[1]	Q1 2026 (unaudited) IFRS Results	Non-IFRS Adjustments	Adjusted Results[2]	Q1 2025 (unaudited) IFRS Results	Non-IFRS Adjustments	Adjusted Results[2]
Revenues	118	-	118	183	-	183
Cost of sales	(71)	-	(71)	(84)	-	(84)
Research and development expenses	(557)	30	(527)	(526)	-	(526)
Sales, marketing, general and administrative expenses	(151)	-	(151)	(121)	-	(121)
Other operating result	(16)	7	(9)	13	(15)	(2)
Operating loss	**(677)**	**37**	**(640)**	**(534)**	**(15)**	**(549)**
Net loss[3]	**(532)**	**37**	**(495)**	**(416)**	**(15)**	**(431)**
Basic and diluted loss per share	(2.10)		(1.95)	(1.73)		(1.79)

1. All Numbers have been rounded and may not add up to the totals. Presentation of the consolidated statements of profit or loss has been condensed. 2. In addition to BioNTech's results determined in accordance with International Financial Reporting Standards ("IFRS"), or IFRS Accounting Standards, or IFRS results, BioNTech reports certain adjusted, non-IFRS measures used internally as a supplemental measure of our business performance (each referred to with the prefix "Adjusted" or, as a whole, "Adjusted Results"). The calculation of these measures and the adjusted results as a whole is based on the concepts of the applicable IFRS Accounting Standards, but includes certain adjustments. Reconciliation of the adjusted results to BioNTech's measures based on IFRS Accounting Standards and more information can be found in BioNTech's Report on Form 6-K for the period ended March 31, 2026, filed on May 5, 2026, which is available at www.sec.gov. While non-IFRS measures may offer additional insights, BioNTech's non-IFRS measures are not, and should not be viewed as a substitute for their most directly comparable IFRS Accounting Standards measures, and should always be considered alongside our financial statements prepared in accordance with IFRS Accounting Standards. 3. Tax effects are not considered as part of BioNTech's non-IFRS adjustments.

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Abbreviation Directory

n L	*nth* line	DNA	Desoxyribonucleic acid	(sq) NSCLC	(squamous) Non-small cell lung cancer
ADC	Antibody-drug conjugate	EGFR	Epidermal growth factor receptor	P&L	Profit and loss statement
adj.	Adjuvant	ESOP	Employee stock ownership plan	PD-L1	Programmed cell death protein ligand 1
ADS	American Depositary Shares	FY	Fiscal year	PDAC	Pancreatic ductal adenocarcinoma
AI	Artificial intelligence	GMP	Good Manufacturing Practice	Q1	First quarter
B7H3	B7 Homolog 3	HCC	Hepatocellular carcinoma	R&D	Research and development
BC	Breast cancer	HER2 (3)	Human epidermal growth factor receptor 2 (3)	RCC	Renal cell carcinoma
BMS	Bristol Myers Squibb			SBP	Share-based payment
CDMO	Contract Development and Manufacturing Organization	HNSCC	Head and neck squamous cell carcinoma	(ES)SCLC	(Extensive stage) small cell lung cancer
		HPV 16	Human papilloma virus 16	SEC	Securities and Exchange Commission
CEPI	Coalition for Epidemic Preparedness Innovations	HR	Hormone receptor	SG&A	Selling, general and administrative expenses
		IFRS	International financial reporting standards		
CPS	Combined positive score	IO	Immuno-oncology	TNBC	Triple-negative breast cancer
CRC	Colorectal cancer	LTI	Long-term incentive	TROP2	Trophoblast cell-surface antigen 2
CRPC	Castration resistant prostate cancer	mRNA	Messenger ribonucleic acid	VEGF-A	Vascular endothelial growth factor A
ctDNA	Circulating tumor DNA	MSS	Microsatellite stability		

BioNTech SE, Mainz
ordinary Annual General Meeting on 15/05/2026

Voting results - Overview

		Valid votes cast		Yes votes		No votes	
		Shares	% of capital stock	Shares	%	Shares	%
Item 2	Resolution on the appropriation of the balance sheet profit for the 2025 financial year	234,799,767	90.65 %	234,734,083	99.97 %	65,684	0.03 %
Item 3	Resolution on the approval of the actions of the Management Board	198,284,335	76.55 %	197,873,174	99.79 %	411,161	0.21 %
Item 4	Resolution on the approval of the actions of the Supervisory Board	236,818,981	91.43 %	236,071,180	99.68 %	747,801	0.32 %
Item 5	Resolution on the appointment of the auditor and the Group auditor for the 2026 financial year and the auditor for any audit or review of interim reports	238,289,587	91.99 %	237,926,738	99.85 %	362,849	0.15 %
Item 6	Resolution on the approval of the Compensation Report	238,280,961	91.99 %	230,176,677	96.60 %	8,104,284	3.40 %
Item 7	Resolution on the amendment of Section 9 para. 1 of the Articles of Association to increase the size of the Supervisory Board	238,286,403	91.99 %	238,238,932	99.98 %	47,471	0.02 %
Item 8.1	Resolution on Elections to the Supervisory Board - Helmut Jeggle	238,286,432	91.99 %	236,945,474	99.44 %	1,340,958	0.56 %
Item 8.2	Resolution on Elections to the Supervisory Board - Prof. Dr. Anja Morawietz	238,288,562	91.99 %	238,000,533	99.88 %	288,029	0.12 %
Item 8.3	Resolution on Elections to the Supervisory Board - Prof. Dr. Rudolf Staudigl	238,289,153	91.99 %	235,551,089	98.85 %	2,738,064	1.15 %
Item 8.4	Resolution on Elections to the Supervisory Board - Dr. Susanne Schaffert	238,288,876	91.99 %	238,242,633	99.98 %	46,243	0.02 %
Item 8.5	Resolution on Elections to the Supervisory Board - Prof. Dr. Iris Loew-Friedrich	238,288,266	91.99 %	216,215,291	90.74 %	22,072,975	9.26 %

Note: Percentages rounded to 2 decimal places

Voting results - Overview

		Valid votes cast		Yes votes		No votes	
		Shares	% of capital stock	Shares	%	Shares	%
Item 9	Resolution on the amendment of Section 16 para. 5 of the Articles of Association to reauthorize the Management Board to provide for the holding of a virtual Annual General Meeting	238,288,290	91.99 %	217,903,070	91.45 %	20,385,220	8.55 %
Item 10	Resolution on the cancellation of the existing Authorized Capital 2025 and the creation of new Authorized Capital 2026 with the possibility of excluding subscription rights and corresponding amendment to the Articles of Association	238,290,388	91.99 %	236,478,038	99.24 %	1,812,350	0.76 %
Item 11	Resolution on the approval of the conclusion of a domination and profit and loss transfer agreement between BioNTech SE and BioNTech Discovery GmbH	238,284,305	91.99 %	237,836,243	99.81 %	448,062	0.19 %

Note: Percentages rounded to 2 decimal places