



Conquering Cancer with Data

Q1 2026

May 7, 2026

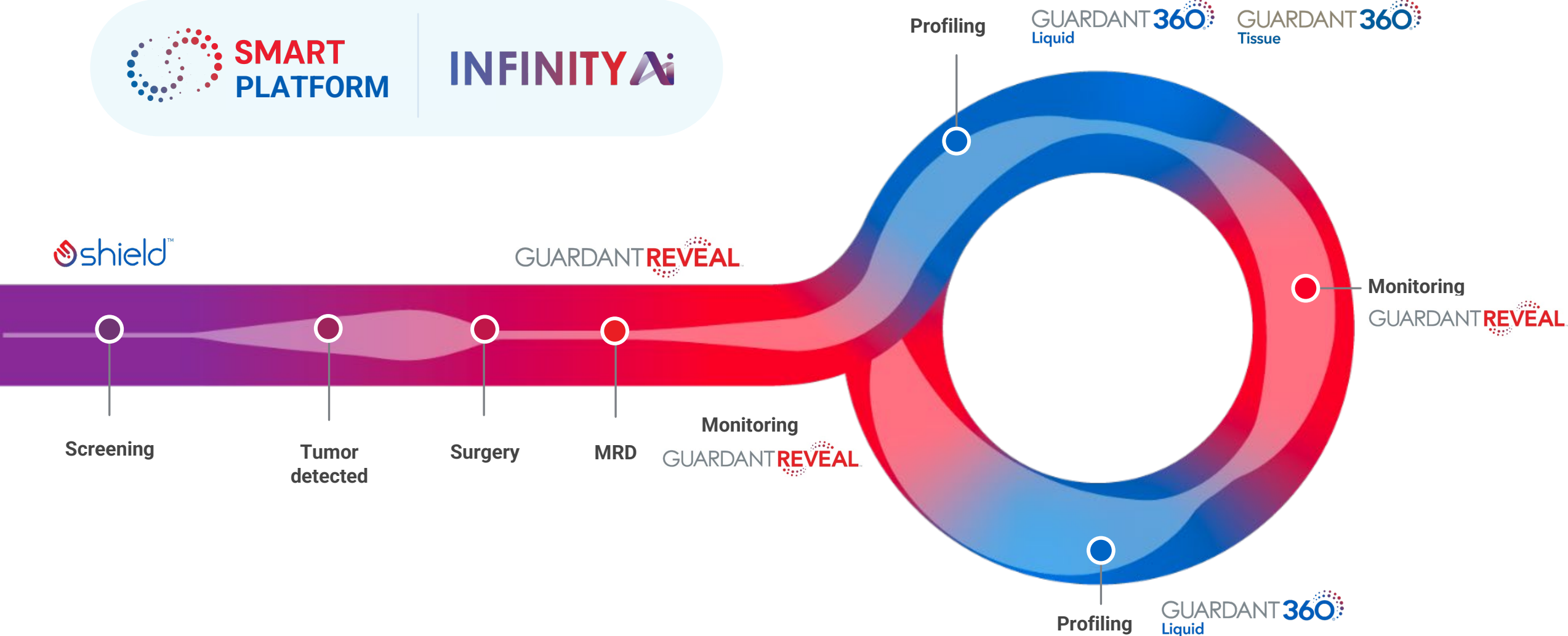
Safe harbor and non-GAAP disclosures

Certain statements in this presentation and the accompanying oral commentary are forward-looking statements within the meaning of federal securities laws. These statements relate to future events or Guardant Health, Inc. (the “Company”)’s future results and involve known and unknown risks, uncertainties and other factors that may cause the actual results, levels of activity, performance or achievements of the Company or its industry to be materially different from those expressed or implied by any forward-looking statements. In some cases, forward-looking statements can be identified by terminology such as “may,” “will,” “could,” “would,” “should,” “to,” “target,” “expect,” “plan,” “anticipate,” “intend,” “believe,” “estimate,” “predict,” “potential” or other comparable terminology. All statements other than statements of historical fact could be deemed forward-looking, including any expectations regarding the Company’s commercial engine as a force multiplier for research and development initiatives; any projections of market opportunities; statements about the Company’s ability to assess potential market opportunities or any statements about the Company’s ability to successfully develop new products and services; any statements regarding expectations for future reimbursement opportunities; any statements regarding the Company’s long-term expectations, including with respect to oncology, liquid biopsy, and other aspects of the Company’s industry; any statements about launching planned new products and additional laboratories, including with respect to Guardant Shield, CGP tissue assay, and laboratories outside the United States; any statements about the Company’s ECLIPSE study; any statements regarding expectations for future regulatory approvals; any statements about historical results that may suggest trends for the Company’s business; any statements of the plans, strategies, and objectives of management for future operations and directions; any statements of expectation or belief regarding future events, opportunities to drive future growth, potential markets or market size, or technology developments; and any statements of assumptions underlying any of the items mentioned. The Company has based these forward-looking statements on its current expectations, assumptions, estimates and projections. While the Company believes these expectations, assumptions, estimates and projections are reasonable, such forward-looking statements are only predictions and involve known and unknown risks and uncertainties, many of which are beyond the Company’s control. These and other important factors may cause actual results, performance or achievements to differ materially from those expressed or implied by these forward-looking statements. The forward-looking statements in this

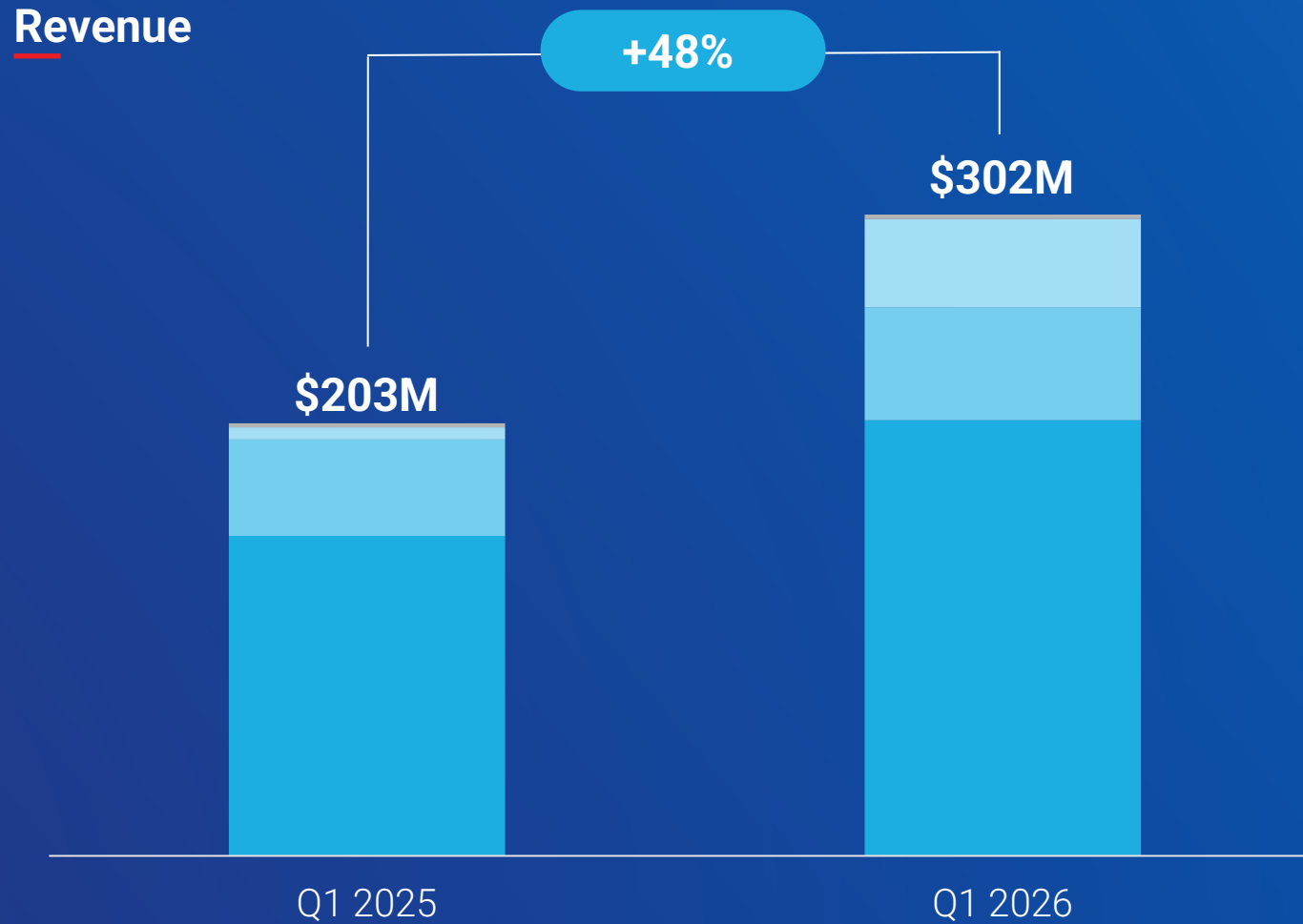
presentation are made only as of the date hereof. For a further description of the risks and uncertainties that could cause actual results to differ from those expressed in these forward-looking statements, as well as risks relating to the business of the Company in general, see the Company’s periodic filings with the Securities and Exchange Commission, including its most recently filed Annual Report on Form 10-K, and in its other reports filed with or furnished to the Securities and Exchange Commission. Except as required by law, the Company assumes no obligation and does not intend to update these forward-looking statements or to conform these statements to actual results or to changes in the Company’s expectations. This presentation also contains estimates and other statistical data made by independent parties and by the Company relating to market size, penetration and growth and other data about the Company’s industry, which involve a number of assumptions and limitations, and you are cautioned not to give undue weight to such estimates. In addition, projections, assumptions, and estimates of the Company’s future performance and the future performance of the markets in which the Company operates are necessarily subject to a high degree of uncertainty and risk. In light of the foregoing, investors are urged not to rely on any forward-looking statement or third-party data in reaching any conclusion or making any investment decision about any securities of the Company.

This presentation includes references to certain financial measures that are not calculated in accordance with GAAP. Reconciliation to the most directly comparable GAAP financial measure may be found in the earnings release furnished to the SEC. We define our non-GAAP measures as the applicable GAAP measure adjusted for the impacts of stock-based compensation and related employer payroll tax payments, contingent consideration, amortization of intangible assets, impairment of non-marketable equity securities, gain on extinguishment of convertible notes, and other non-recurring items. Free cash flow is defined as net cash used in operating activities in the period less purchases of property and equipment in the period.

One platform for the entire patient journey



Q1 revenue accelerates to 48% growth



Oncology drives growth with strong revenue and volume expansion

Oncology Revenue

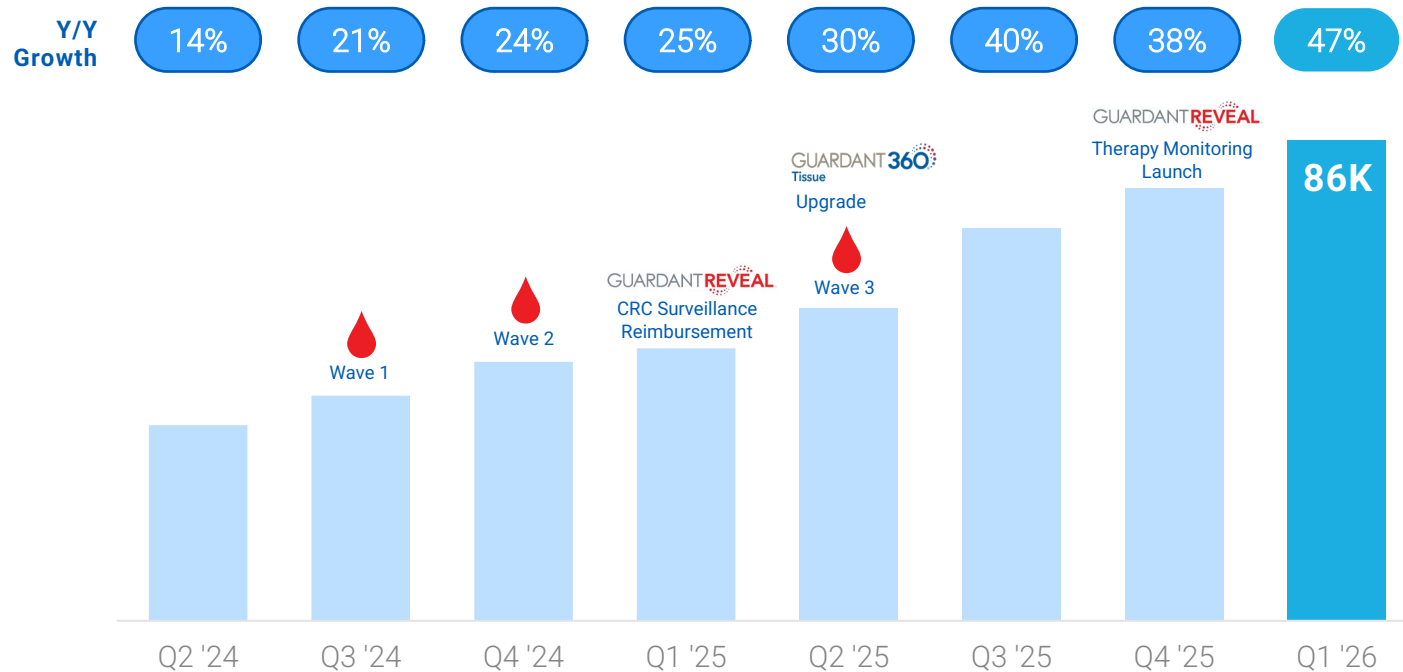


Oncology Volume



Multi-product momentum driving Oncology volume acceleration

Oncology Volume



- ✓ Guardant360 Liquid y/y volume growth of approximately 30% driven by Smart Platform
- ✓ Guardant360 Tissue remains second fastest growing Oncology product with volume growth accelerating in Q1
- ✓ Reveal continues to be the fastest growing Oncology product with volume growth exceeding 100% y/y

Rapidly scaling data + InfinityAI is fueling innovation and widening competitive moat

>1,000,000
PATIENT SAMPLES

Genomics

Multi-modal

Claims

Longitudinal Data

EMR

>500,000
EPIGENETIC PROFILES

>100
TUMOR TYPES

INFINITY *ai*

Smart Apps

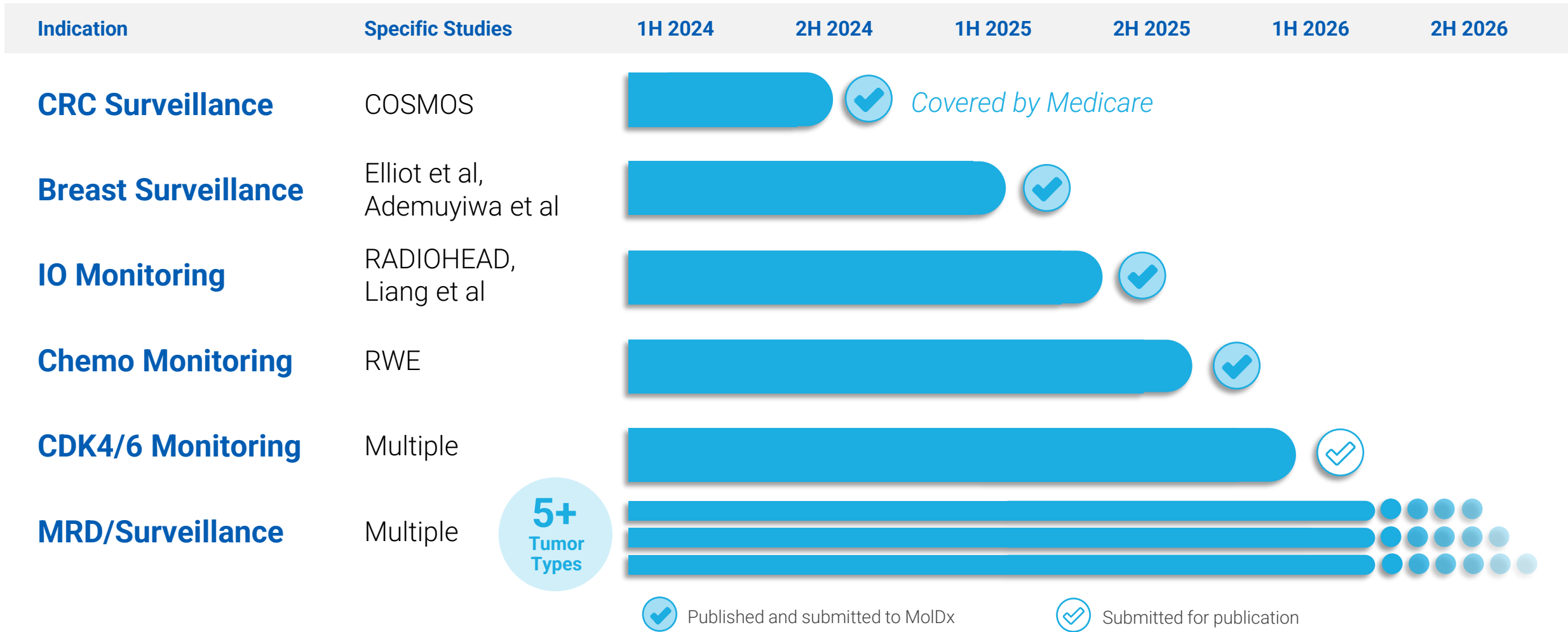
New product development

Novel biological signatures

Real world data generation

Faster regulatory approval

Strong Reveal data pipeline supporting indication expansion



Raising the bar for liquid and tissue-based therapy selection to potentially unlock greater volume

Expected 2026 launch

GUARDANT360
Liquid CDx

Will be the most comprehensive FDA approved liquid biopsy for therapy selection

Streamlines therapy selection portfolio

Whole transcriptome upgrade

GUARDANT360
Tissue

Second major platform upgrade in less than a year for Guardant360 Tissue

Further expands transcriptomic power of the Smart Platform

AACR 2026: Strong presence across the portfolio, including new Smart Apps

INFINITY^{AI} + Smart Apps

AI driven prediction & tracking of therapeutic response:

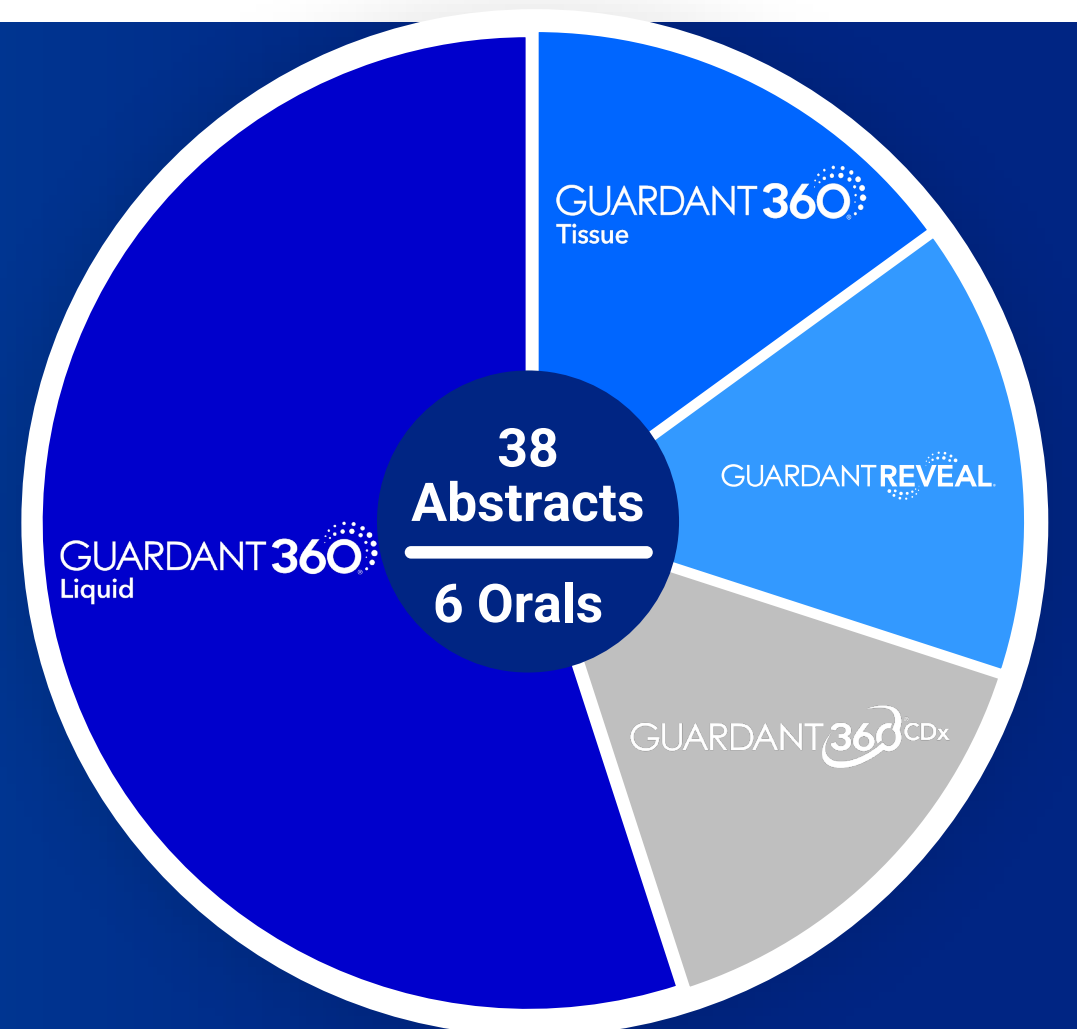
- Therapy response (SERD, anti-EGFR)
- Tumor Fraction as a measure of molecular progression

Multi-modal modeling:

- Integration of clinical + genomic + epigenomic data

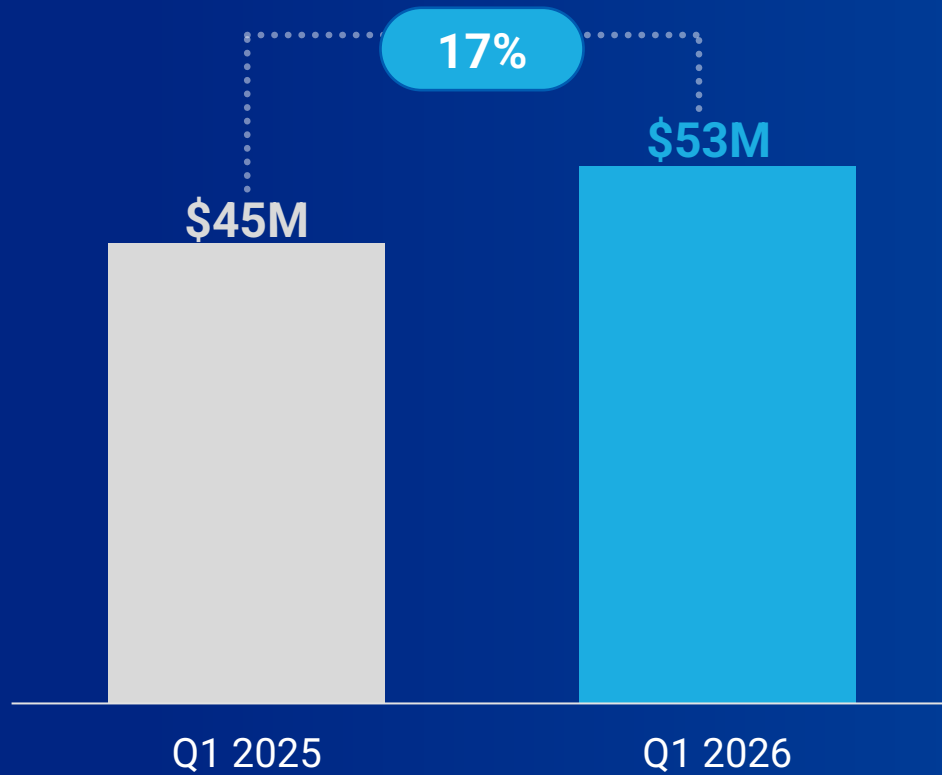
Improved detection of challenging genomic alterations with epigenomics:

- ALK-fusion rescue
- MTAP deletions



InfinityAI and Smart Platform deepening strategic partnerships with leading biopharma companies

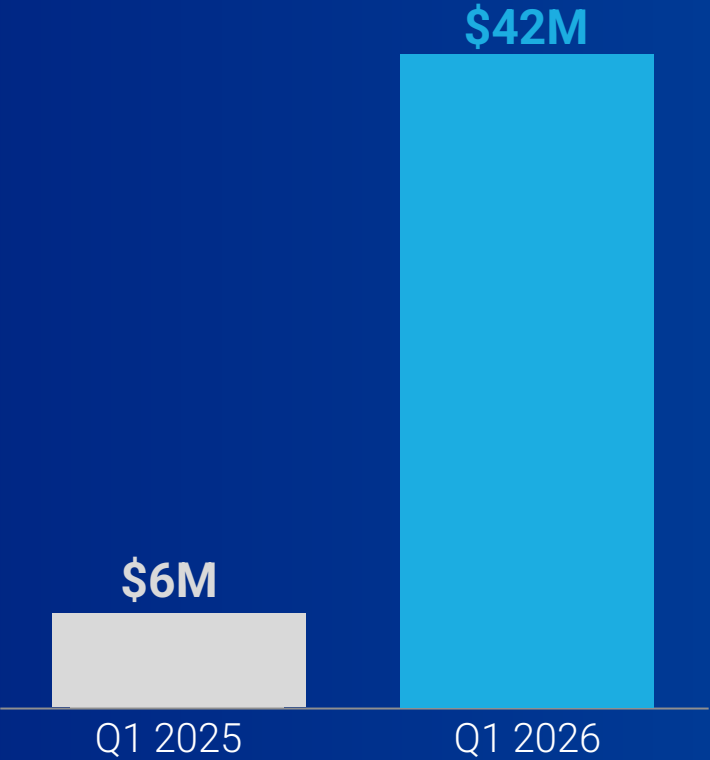
Biopharma & Data Revenue



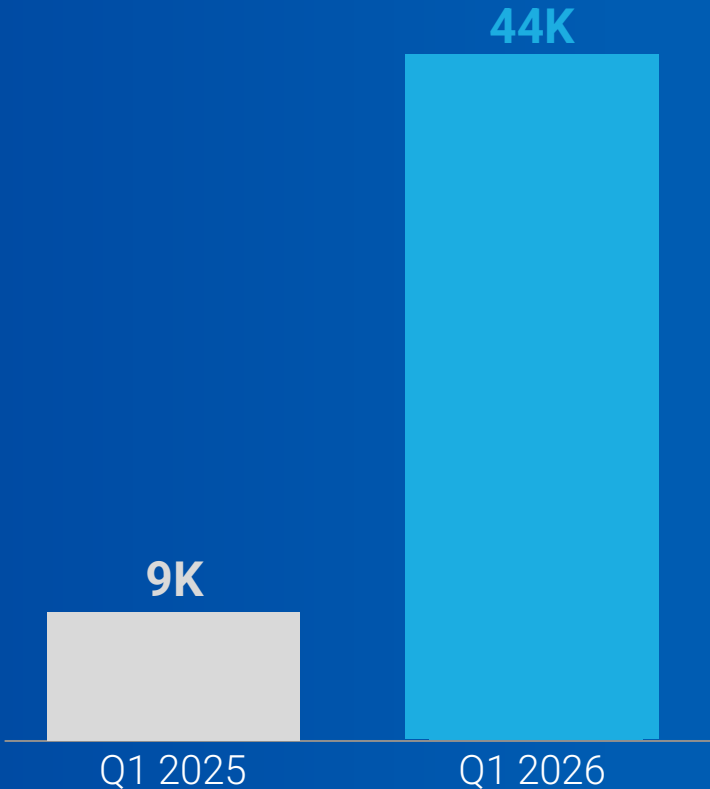
- ✓ FDA approval for Guardant360 CDx as a companion diagnostic for Arvinas and Pfizer's VEPPANU for ER+/HER2- ESR1 mutated advanced breast cancer
- ✓ FDA approval of Guardant360 CDx as a companion diagnostic for Pfizer's Braftovi in BRAF V600E-mutant metastatic colorectal cancer
- ✓ Guardant InfinityAI contributed to the first tumor agnostic approval of Daiichi's ENHERTU® in Japan
- ✓ Collaboration with Merck to develop companion diagnostics and commercialize new therapies
- ✓ Collaboration with Nuvalent to develop companion diagnostics in targeted cancer therapy, with initial emphasis on Guardant360 Tissue

Shield momentum continues to drive rapid revenue and volume growth

Screening Revenue



Screening Volume



Q1 Screening highlights

- ✓ Strong volume throughout the quarter with **accelerated momentum** in March
- ✓ **Direct-to-Consumer and Influencer campaigns** launched in conjunction with Colorectal Cancer Awareness Month
- ✓ **Quest collaboration** commenced nationwide
- ✓ Shield continues to demonstrate **>90% adherence rate**
- ✓ Launched **Shield Multi-Cancer Detection (MCD)** in Asia through Manulife partnership

DTC campaigns across TV and digital driving strong consumer engagement



Based on data from clinical studies, Shield has limited detection (55%-65%) of Stage 1 colorectal cancer and does not detect 87% of precancerous lesions. One out of 10 patients with a negative result may have a precancer that would have been detected by a screening colonoscopy.

PATRICK DEMPSEY
screened for colon cancer with the Shield™ blood test

Patrick Dempsey is a paid partner of Guardant Health

shield by GUARDANT Rx only

Patrick Dempsey screens for colon cancer with the Shield blood test.

Shield Cancer Screen [Subscribe](#)

53,493,299 views 2 months ago



How Shield works

The Shield blood test detects signals for colon cancer from DNA shed into the blood.^{1,2}

You and your healthcare provider will receive your test results in approximately 2 weeks.

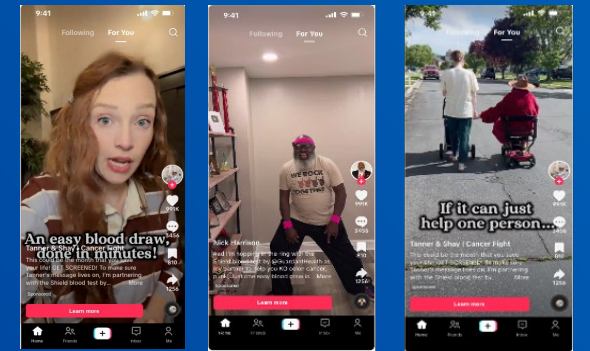
100,000 Shield blood tests have been completed since FDA approval.³

Ask if Shield is right for you

- Discuss the benefits and risks of colon cancer screening with Shield as part of a shared decision made with your healthcare provider
- Provide your medical history to assess your eligibility
- Shield is not a replacement for a follow-up diagnostic colonoscopy or surveillance colonoscopy in high risk individuals. Shield has limited ability to prevent the development of colorectal cancer, given its limited detection of advanced precancerous lesions.

Shield™, screen for colon cancer with just a blood test

From Guardant Health, a company trusted in blood tests for cancer for over a decade.



Expanded marketing, EMR, and phlebotomy initiatives fueling record HCP engagement

Nationwide EMR

Coast to Coast Phlebotomy

Extensive Guardant network across 50 states and over 40,000 phlebotomists

[View in browser](#)

March is Colon Cancer Awareness Month. Don't wait. Get screened with Shield™

Patrick Dempsey for Shield*
Actor, Advocate, Screened with Shield™
*Patrick Dempsey is a paid partner of Guardant Health.

[See Patrick get screened with Shield™](#)

Did you know that ~75% of people who die from colorectal cancer are not up to date with their screenings?

The Shield™ blood test

The only FDA-approved blood test for CRC screening with Medicare coverage

Now recommended by the NCCN Clinical Practice Guidelines in Oncology (NCCN Guidelines®) for Colorectal Cancer Screening
The first and only blood test recommended for first-line CRC screening*

CRC remains the 2nd-deadliest cancer in the US† | Among those who died of CRC, 3 out of 4 people were not up to date with screening**

Shield is a convenient, accurate, and easy solution for low CRC screening adherence*†

- ✓ HIGHLY ACCURATE CRC DETECTION
Validated by the CLIPSE study, published in NEJM*
83% OVERALL CRC SENSITIVITY | 90% SPECIFICITY
- ✓ 90% ADHERENCE

NOW AVAILABLE |

Search "Shield" to order
Quantum® Lab Services Manager
your EHR
Order test code 18441
Test is performed and billed through Guardant Health.



Improving the Shield experience through product enhancements

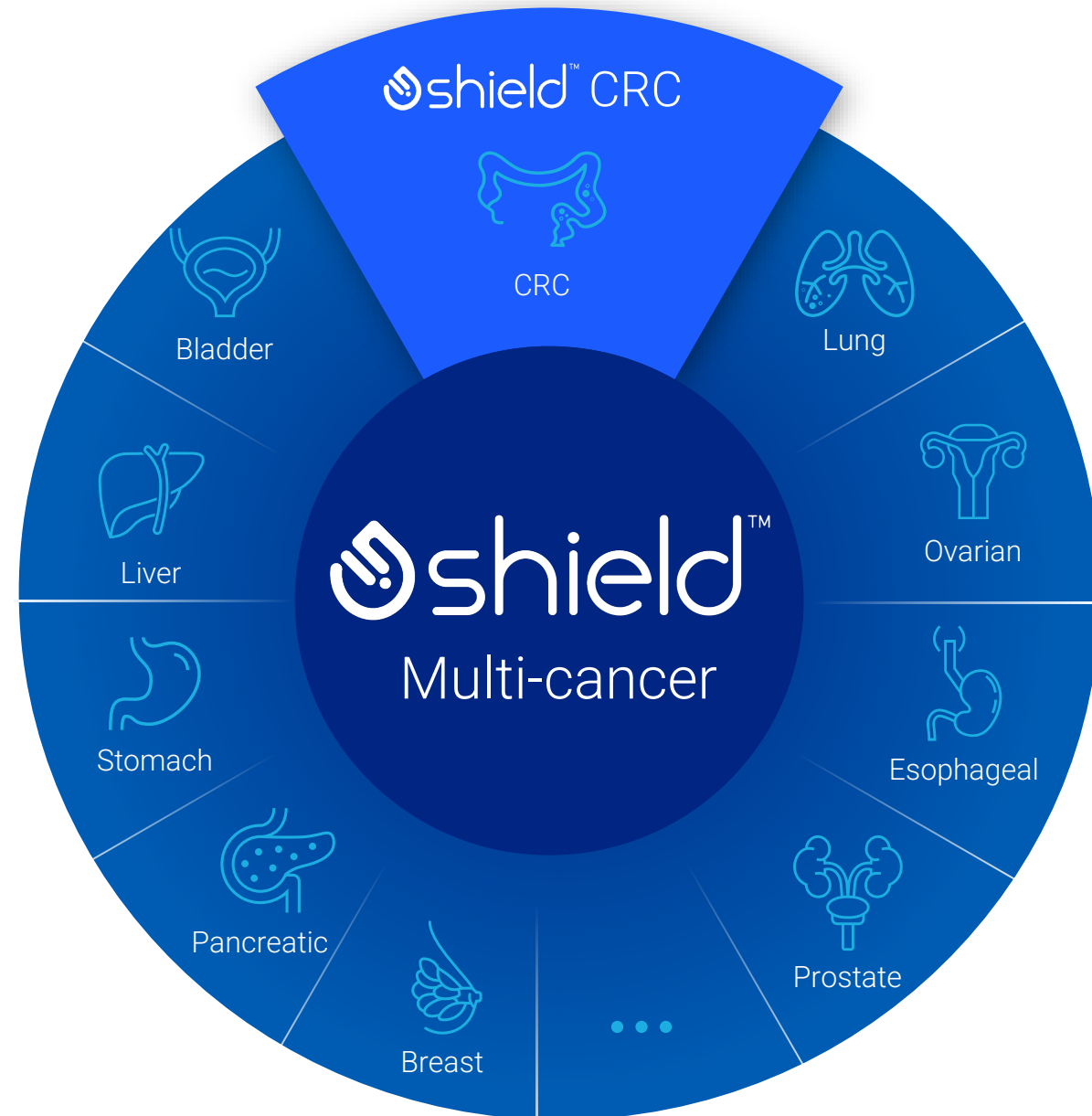
2-tube Shield kit now FDA approved



Shield is a multi-cancer detection platform

When Shield is ordered for CRC screening:

- Physician opts in to receive MCD results report
- Patient authorizes release of medical data



Shield MCD launched in Asia through partnership with Manulife



- ✓ Leading global life and health insurance provider
- ✓ Strong presence in Asia with >13 million customers
- ✓ Shield MCD initially offered to Manulife customers in Hong Kong, Philippines and Singapore



Revenue growth acceleration led by Oncology & Screening

	Q1'26	Q1'25	% Growth
Total Revenue	\$302M	\$203M	48%
Oncology	\$205M	\$151M	36%
Biopharma & Data	\$53M	\$45M	17%
Screening	\$42M	\$6M	>600%
Licensing & Other	\$2M	\$2M	--

Q1 2026 non-GAAP financial highlights

Non-GAAP Measures ²	Q1'26	Q1'25
Gross Margin ¹	66%	65%
Total Operating Expense	\$268M	\$200M
R&D	\$77M	\$75M
S&M	\$154M	\$94M
G&A	\$37M	\$31M
Adjusted EBITDA ³	\$(59M)	\$(59M)
Free Cash Flow ⁴	\$(71M)	\$(67M)
	March 31, 2026	December 31, 2025
Cash & investments ⁵	\$1.2B	\$1.3B

1. Gross margin is defined as gross profit divided by total revenue.
2. Non-GAAP measures are defined as the applicable GAAP measures adjusted for the impacts of stock-based compensation and related employer payroll tax payments, contingent consideration, amortization of intangible assets, impairment of non-marketable equity securities, gain on extinguishment of convertible notes, and other non-recurring items. Please refer to the relevant non-GAAP tables in the associated press release for reference.
3. Adjusted EBITDA is defined as net loss adjusted for interest income; interest expense; other income (expense), net; provision for income taxes; depreciation and amortization expense; stock-based compensation expense and related employer payroll tax payments; contingent consideration; and other non-recurring items.
4. Free cash flow is defined as net cash used in operating activities in the period less purchase of property and equipment in the period.
5. Cash & investments include cash, cash equivalents, restricted cash & marketable securities.

Screening gross margin expansion as Shield volume increases

Screening Non-GAAP
Gross Margin

18%

56%

Shield Non-GAAP
Cost per Test

\$520

\$420

Q1 2025

Q1 2026

Raising full year guidance on strong Q1 performance

Revenue	Current Guidance	Prior Guidance
Total	\$1.30B – \$1.32B 32% - 34% y/y growth	\$1.25B - \$1.28B 27% - 30% y/y growth
Oncology	28% - 29% y/y growth	25% - 27% y/y growth
Biopharma & Data	Low double-digit growth	Low double-digit growth
Screening	\$186M - \$198M 230K - 245K Shield volume	\$162M - \$174M 210K - 225K Shield volume

Non-GAAP Gross Margin

64% – 65%

Non-GAAP Operating Expenses

\$1.05B – \$1.07B

Free Cash Flow

(\$185M) – (\$195M)

Key catalysts across the continuum of cancer care in 2026

ONCOLOGY

Therapy Selection

- Guardant360 Liquid FDA approval and launch
- Transition to NovaSeq X
- ESR1 monitoring launch
- Guardant360 Smart Platform app expansion

MRD

- MoIDx coverage for Reveal Breast MRD
- MoIDx coverage for Reveal therapy monitoring
- MoIDx submissions in additional tumor types and indications
- Reveal Ultra launch

BIOPHARMA & DATA

- CDx approvals, including SERENA-6
- Additional strategic biopharma partnerships
- InfinityAI data partnerships

SCREENING

- ACS guidelines
- Quest collaboration launch
- OUS self-pay expansion

