



Where Molecular Science Meets Artificial Intelligence

Q2 2026 Earnings Call
August 5, 2026

Important Information and Disclaimer



Forward-Looking Statements

This presentation contains forward-looking statements, within the meaning of the Private Securities Litigation Reform Act of 1995 and other federal securities laws. All statements other than statements of historical facts contained in this presentation are forward-looking statements, including statements regarding our business, solutions, plans, objectives, goals, industry trends, financial outlook and guidance. In some cases forward-looking statements can be identified by words such as “may,” “will,” “should,” “would,” “expect,” “plan,” “anticipate,” “could,” “intend,” “target,” “project,” “potential,” “contemplate,” “believe,” “estimate,” “predict,” or “continue” or similar expressions. You should not rely upon forward-looking statements as predictions of future events. Although we believe that the expectations reflected in these forward-looking statements are reasonable based on information currently available to us, we cannot guarantee that the future results, discoveries, levels of activity, performance or events and circumstances reflected in forward-looking statements will be achieved or occur. Forward-looking statements involve known and unknown risks and uncertainties, some of which are beyond our control. Risks and uncertainties that could cause our actual results to differ materially from those indicated or implied by the forward-looking statements in this presentation include, among other things: continued development, performance and commercialization of Caris Detect; developments in the precision medicine industry; our future financial performance, results of operations or other operational results or metrics; development, analytical and clinical validation, timing and performance of future solutions by us and our competitors; commercial market acceptance for our solutions, including acceptance of preventive as well as diagnostic testing paradigms, and our ability to meet resulting demand; the rapidly evolving competitive environment in which we operate; third-party payer reimbursement and coverage decisions related to our solutions; the impact on our future volumes of the continued execution of our strategy to re-align and expand our sales organization; risks related to data management, storage, and processing capabilities and our ability to integrate and deploy artificial intelligence and advanced data analytics technologies; our ability to protect and enhance our intellectual property; regulatory requirements, decisions or approvals (including the timing and conditions thereof) related to our solutions; reliance on third-party suppliers; risks related to data security, patient privacy, and compliance with healthcare data protection regulations as well as potential cybersecurity threats to our data platforms; our compliance with laws and regulations; the outcome of government investigations and litigation; risks related to our indebtedness; and our ability to hire and retain key personnel as well as risks, uncertainties, and other factors described in the section titled “Risk Factors” and elsewhere in our Annual Report on Form 10-K filed on March 3, 2026, and in our other filings we make with the SEC from time to time. We undertake no obligation to update any forward-looking statements to reflect changes in events, circumstances or our beliefs after the date of this presentation, except as required by law.

Non-GAAP Financial Measures

We use certain financial measures not calculated in accordance with generally accepted accounting principles in the United States (“GAAP”) to supplement our condensed consolidated financial statements, which are presented in accordance with GAAP. We believe the non-GAAP financial measures we use, Adjusted EBITDA and free cash flow, are useful in evaluating our performance and liquidity. Our non-GAAP financial measures have limitations as analytical tools, however, and you should not consider them in isolation or as substitutes for analysis of our results as reported under GAAP. Other companies, including other companies in our industry, may not use these measures or may calculate these measures differently than as presented herein, limiting their usefulness as comparative measures.

We use Adjusted EBITDA in conjunction with GAAP measures as part of our overall assessment of our performance, including the preparation of our annual operating budget and quarterly forecasts, to evaluate the effectiveness of our business strategies, and to communicate with our board of directors concerning our financial performance. We believe Adjusted EBITDA is also helpful to investors, analysts, and other interested parties because it can assist in providing a more consistent and comparable overview of our operations across our historical financial periods. We believe free cash flow is a useful measure of liquidity that provides an additional basis for assessing our ability to generate cash.

For definitions and a reconciliation of our historical non-GAAP financial measures to the most directly comparable financial measures calculated in accordance with GAAP, see our Press Release for the most recently-completed fiscal quarter available on the “News & Events” section of our website at <https://investor.carislifesciences.com/news-events/events>.

Differentiated technology across the cancer journey




MI PROFILE[®]

WES / WTS / IHC

Tissue-based therapy selection

COMMERCIAL


CARIS ASSURE[®]

WES / WTS / CH

Blood-based therapy selection

COMMERCIAL


CARIS CHROMOSEQ[™]

WGS / WTS

Heme therapy selection

COMMERCIAL


CARIS MI CLARITY[™]

Digital AI

Early-stage breast predictor

COMMERCIAL


CARIS DETECT[™]

WGS/WTS

Multi-cancer early detection

COMMERCIAL


CARIS MRD[™]

WES / WTS / WGS

Minimal residual disease

VALIDATION IN-PROCESS

1.13MM+

PROFILED CASES

845,000+

MATCHED CASES WITH OUTCOMES

783,000+

WHOLE TRANSCRIPTOMES CASES

733,000+

WHOLE EXOMES CASES

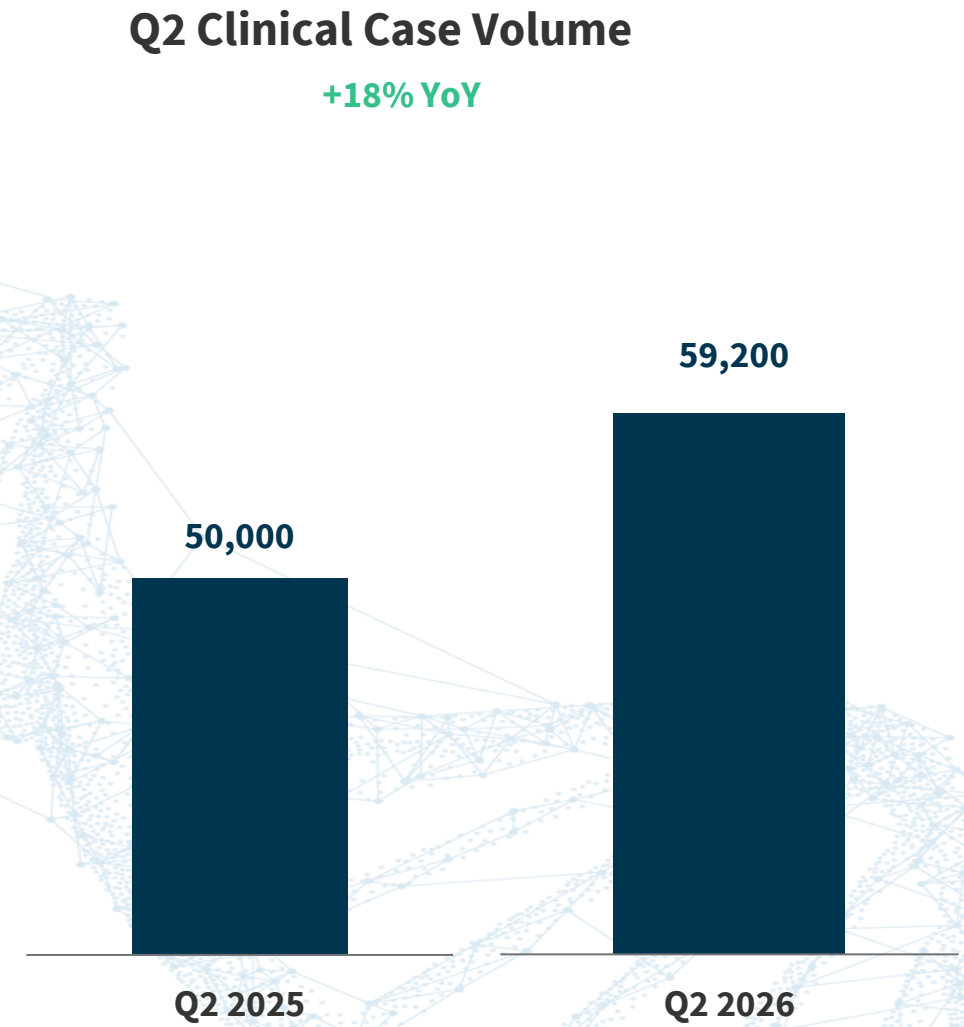
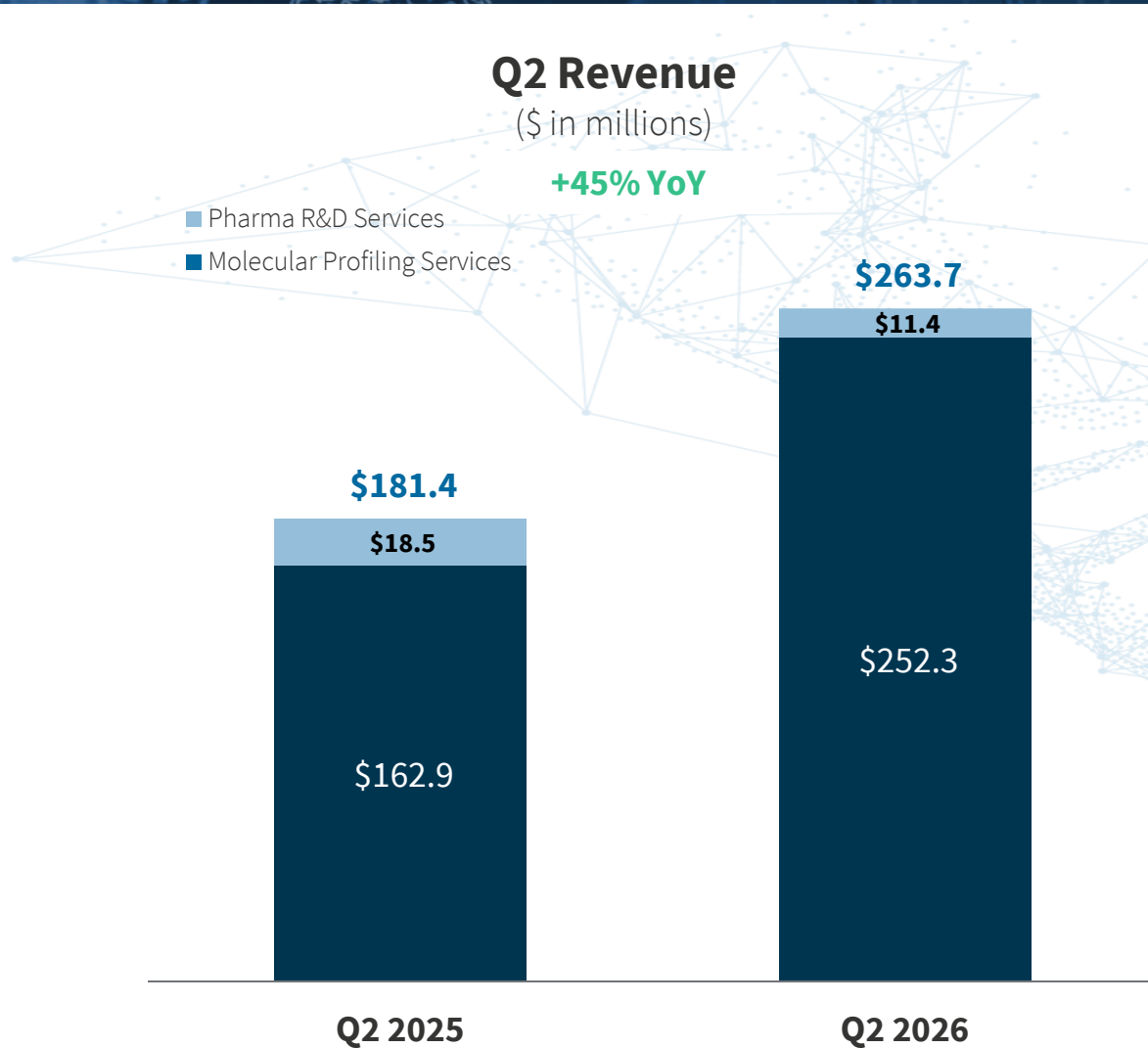
6,200+

ORDERING ONCOLOGISTS

101

POA MEMBERS

Strong Revenue and Clinical Volume Growth



Q2 2026 Performance Highlights



FINANCIAL PERFORMANCE

TOTAL REVENUE

\$263.7M +45% YoY

MOLECULAR PROFILING REV

\$252.3M +55% YoY

CLINICAL CASES

~59,200 +18% YoY

~114k Whole Exome/Whole Transcriptome tests
~345k Clinical Oncology tests

GAAP GROSS MARGIN

68% +500 bps YoY

ADJUSTED EBITDA*

\$55.7M +\$39.0M YoY

NET CASH FROM OPERATIONS

\$28.5M +290% YoY

GAAP net loss \$(0.6)M vs. \$(71.8)M YoY

Free Cash Flow* \$6.4M vs. \$5.9M YoY

\$793.1M cash & securities

ADDITIONAL OPERATING HIGHLIGHTS



Launched Caris Detect

A groundbreaking multi-cancer early detection blood test.



Salesforce Expansion

290 team members as of June 30, 2026.



Volume Growth Acceleration

12% sequential growth from Q126.



Continued Profitability

Positive Adjusted EBITDA for 5th consecutive quarter.



Published Caris Whole-exome TMB study

WES-based TMB linked to improved overall survival.



Published Caris Lookback Program study

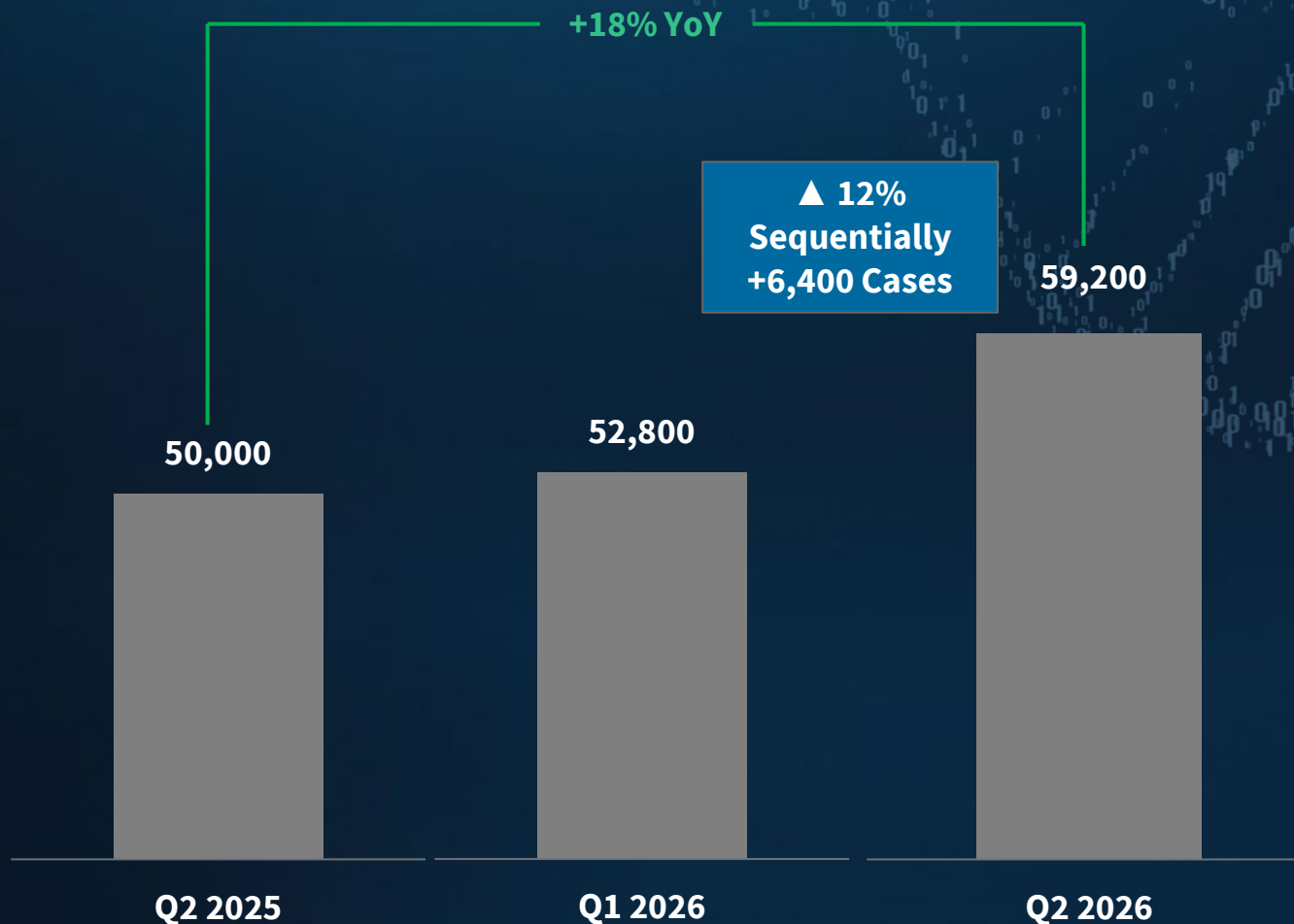
Reinforced the clinical value of MI Cancer Seek testing.

Note: Volume rounded to nearest 100.

* See earnings release for reconciliation.

Commercial Execution Driving Sequential Acceleration from Q126

CLINICAL CASE VOLUME



ADDITIONAL COMMERCIAL HIGHLIGHTS



48,300 MI Profile cases

13% YoY · 11% sequential growth



10,700 Caris Assure cases

50% YoY · 17% sequential growth



290 commercial team members

Continued expansion of commercial sales force



74%+ of orders placed via EMR/Portal

Streamlined digital ordering



3,300+ physicians using EMR

Direct, in-workflow ordering



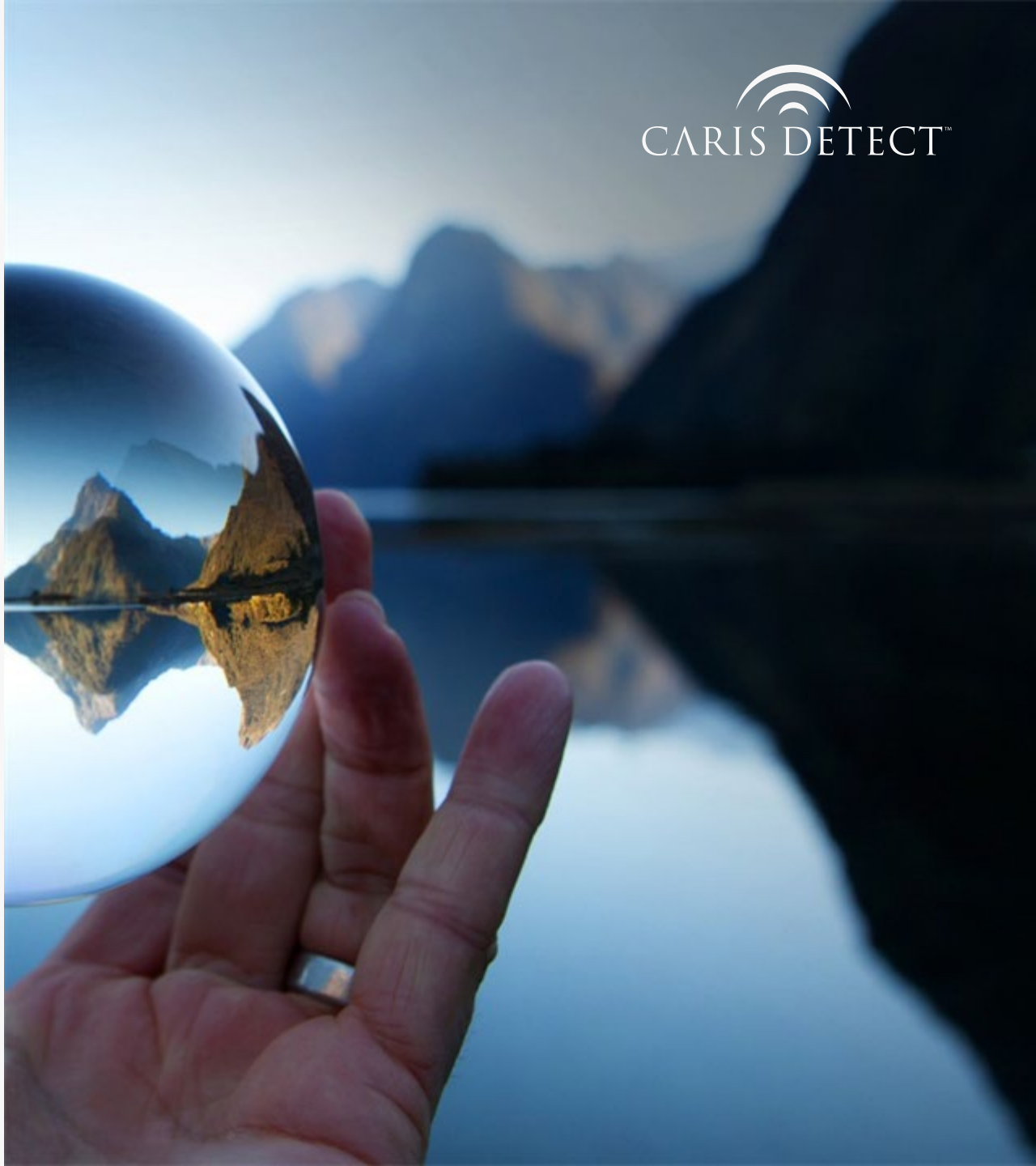
1,300+ Publications/Manuscripts

Evidence generation driving adoption

Note: Volume rounded to nearest 100.

Redefining Early-Stage Cancer Detection

CARIS DETECT™



Caris Detect: Tumor Type Update

58 Cancer Types Detected



Coverage spans solid tumors, hematologic malignancies, and unresolved cancer-signal localization from a routine blood draw.

LUNG / THORACIC

- Non-small cell lung cancer
- Small-cell lung cancer
- Pulmonary neuroendocrine carcinoma
- Thymic carcinoma

GI / CRC

- Colorectal adenocarcinoma
- Appendiceal neoplasms
- Distal small-bowel / terminal-ileum lesions
- Anal squamous cell carcinoma

BREAST

- Invasive ductal carcinoma
- Invasive lobular carcinoma
- Inflammatory
- Metaplastic and other primary breast carcinomas

PROSTATE / TESTIS

- Prostatic adenocarcinomas and other primary prostate malignancies
- Testicular carcinoma

PANCREAS / AMPULLA

- Pancreatic adenocarcinomas
- Ampullary carcinoma

UPPER GI

- Esophageal carcinoma
- Gastroesophageal-junction carcinoma
- Gastric adenocarcinoma
- Proximal-duodenal malignancy
- Gastrointestinal stromal tumor (GIST)

SKIN

- Cutaneous melanoma
- Cutaneous squamous cell carcinoma
- Merkel-cell carcinoma
- Other visible skin malignancies

BILIARY TRACT

- Intrahepatic and extrahepatic cholangiocarcinoma
- Gallbladder carcinoma

HEAD AND NECK

- Oral cavity carcinoma
- Oropharyngeal carcinoma
- Laryngeal carcinoma
- Nasopharyngeal carcinoma
- Salivary-gland cancers
- Thyroid carcinoma

PRIMARY BRAIN TUMOR

- Glioblastoma

UROTHELIAL TRACT

- Urothelial bladder carcinoma
- Upper-tract urothelial carcinoma (UTUC)
- Urethral carcinoma

RENAL

- Renal-cell carcinoma
- Urothelial carcinoma of the renal pelvis

GYNECOLOGIC

- Ovarian epithelial carcinoma
- Fallopian-tube carcinoma
- Primary peritoneal carcinoma
- Germ-cell / sex-cord tumors
- Endometrial carcinoma
- Cervical carcinoma
- Vulvar carcinoma

HEPATIC (HCC)

- Hepatocellular carcinoma

SOFT TISSUE TUMORS

- Angiosarcoma
- Liposarcoma
- Leiomyosarcoma
- Malignant peripheral nerve sheath tumor
- Rhabdomyosarcoma

BONE TUMORS

- Osteosarcoma

HEMATOLOGIC MALIGNANCIES

- Acute myeloid leukemia (AML)
- Myelodysplastic syndromes (MDS)
- Myeloproliferative neoplasms (MPN)
- Large B-cell lymphoma

OTHER / UNCERTAIN PRIMARY

- Tumor type not confidently localized

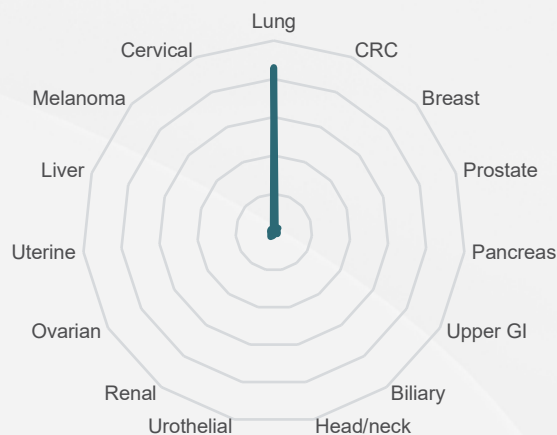
Caris Detect: Tissue-of-Origin Classifier



Pinpointing the tumor's origin, early — with fewer procedures

The classifier concentrates probability on the true site, turning an open-ended cancer hunt into a short, prioritized workup.

NARROWING THE SEARCH



True Positives
(n=2,507)

Potential False
Positives
(n=30)

BY THE NUMBERS

■ **83.9%**

resolved in one work-up

■ **1.19**

procedures per patient (avg.)

■ **99.8%**

localized within two work-ups

■ **30.6%**

resolved in one work-up

■ **1.89**

procedures per patient (avg.)

■ **~100%**

resolved within two work-ups

Representative validation patient — probability collapses onto one origin.

WHY IT MATTERS

● Fewer procedures

Most patients resolve in one or two studies instead of a scattershot battery.

● Less radiation

Narrow-first imaging spares anatomy a whole-body scan would expose by default.

● Faster to answer

A prioritized shortlist compresses time from signal to confirmed primary site.

● Guides unknown primaries

Directly supports metastasis-of-unknown-primary workups where origin is uncertain.

● Transparent & tunable

Every routing step is explainable; the coverage balance is a dial, not a black box.

● Scalable triage

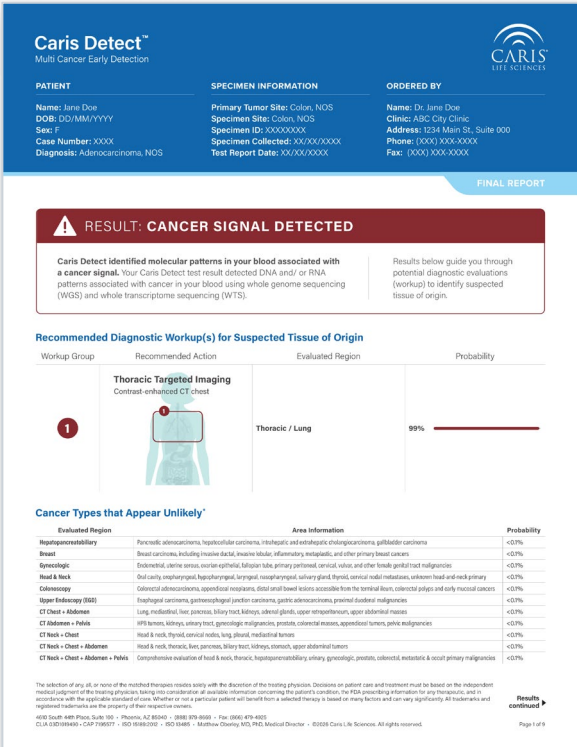
Runs from a routine blood draw — deployable ahead of specialist imaging capacity.

Caris Detect: Tissue-of-Origin Classifier

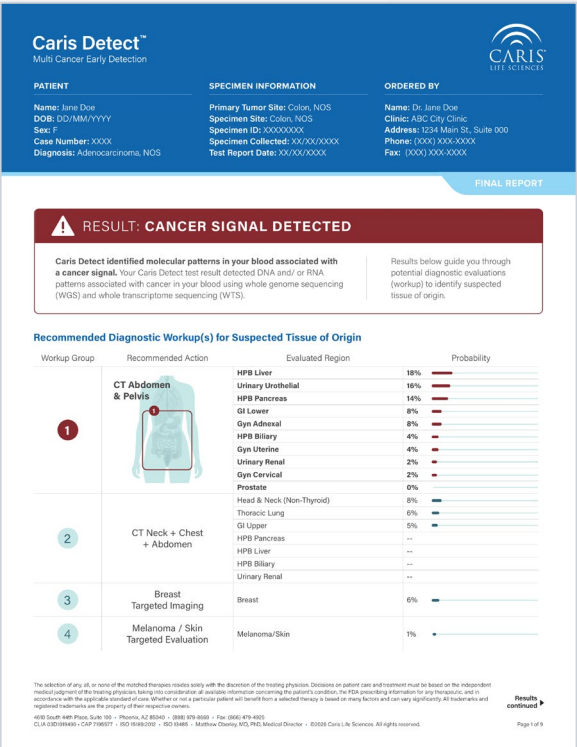


Every result is delivered as a prioritized workup — per-site probabilities paired with the exact screening study to order.

THE DELIVERABLE



Example 1



Example 2

Caris Detect™ Report (sample deliverable)

WHAT THE REPORT COMMUNICATES

1

A clear, actionable result

“Cancer Signal Detected” is stated plainly, backed by whole-genome (WGS) and whole-transcriptome (WTS) sequencing.

2

A ranked, prioritized workup

Suspected tissues of origin are ordered by probability — examples shown.

3

A specific next step per site

Each workup group names the exact study to order and maps it on the body — e.g. contrast-enhanced CT chest, or CT abdomen & pelvis.

4

What the data deprioritizes

Tissue types the signal makes unlikely are listed separately at <0.1%, narrowing the diagnostic search from the outset.

Mutational Cleanse

**From Personalized Medicine
To Personalized Prevention**



Early detection to personalized immune interception

- First Caris Detect identifies disease early
- Caris MAX 10,000x whole-exome sequencing to identify somatic neoepitopes
- Use our proprietary AI to identify immunogenic mutations for personalized immunotherapy.

1**CARIS DETECT**

Blood-based early signal and tumor-of-origin routing while disease burden is low.

60.3% Stage I/II sensitivity

2**CARIS MAX**

Ultra-deep Mutational Analysis eXome plus HLA and germline logic separates real somatic targets from noise.
rare variant discovery

3**AI TARGET RANKING**

Scores pathogenicity, clonality, expression, processing, HLA fit and blood-on-target risk.
immunogenicity triage

4**PERSONALIZED THERAPY**

Top neoepitopes become patient-specific immune targets, with VAF/ctDNA and T-cell monitoring.
closed-loop readout

**Find the dangerous clone early. Make its mutation visible to the immune system.
Remove it before clinically overt disease emerges.**

Driven by Biology: low-burden, blood-accessible clones where the selected mutation is reproducible, pathogenic, HLA-presented and quantitatively measurable over time.

Caris AI Performance identifying immunogenic mutations

83.8%

PPV
top five variants

86.5%

Sensitivity
top variant / patient

81.9%

Sensitivity
top five variants

62.3%

Sensitivity
top 100 variants

Operational implication: focus manufacturing on mutations most likely to prime a mutant-specific T-cell response.

Pipeline Update

**MI Clarity • Caris MRD
Clinical Evidence**



Caris MI Clarity: Upgraded

Upcoming Version 2 brings treatment decision support to patients who were not tested at diagnosis

V1 Capabilities



Both early & late recurrence prognosis

Distant-recurrence risk across Years 0–5 and Years 5–15



Ordering at diagnosis

Actionable recurrence-risk insight right from diagnosis



Fast turnaround time

Rapid results to support timely treatment decisions



Cost

Accessible, cost-effective testing

NEW V2



Chemotherapy decision support

Identifies eligible premenopausal patients more or less likely to benefit from chemotherapy



Extended endocrine therapy decision support

Informs treatment decisions beyond the first five years



Order years later after diagnosis

Expands patient access beyond diagnosis when the extended endocrine therapy decision is due



Integrated early and late treatment decision support

One report combining risk recurrence prognosis with chemotherapy and extended endocrine therapy decision support.

Caris MRD Update

Two complementary approaches to minimal residual disease



WHOLE EXOME

WHOLE TRANSCRIPTOME

Initial intended use: CRC

A minimal residual disease diagnostic for patients with stage II and III solid tumors following curative-intent treatment, prior to adjuvant chemotherapy. Simultaneously profiles cancer-associated ctDNA / RNA alterations from a whole-blood sample.

- ✓ Uses Caris Assure platform
- ✓ Tumor naive
- ✓ Additional indications to follow

Compiling additional data for MolDX TA

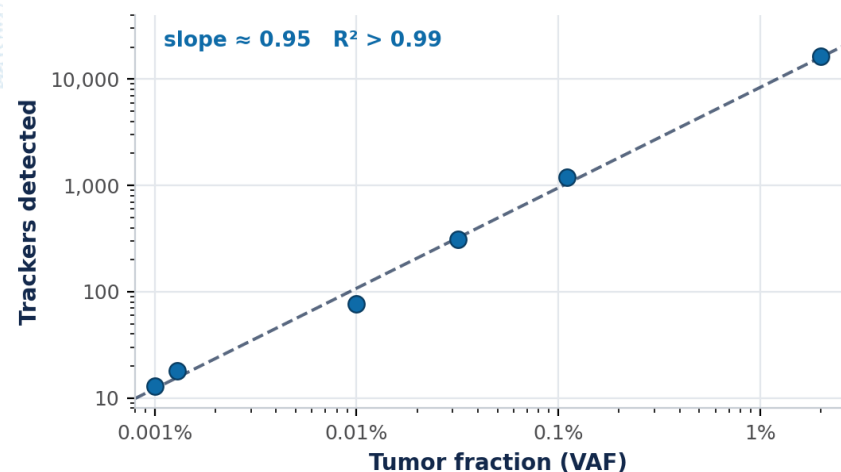


WHOLE GENOME

Intended use: Pan-tumor, Stage I-III. Tumor / normal WGS identifies trackers; a comprehensive approach minimizes false negatives and maximizes tracker count to achieve ultra-low PPM.

- ✓ Uses Caris Precision WG platform
- ✓ Tumor informed
- ✓ Ultra-low sensitivity

TUMOR-INFORMED ANALYTICAL PERFORMANCE



>5 logs

linear dynamic range

~15,000 trackers

median value

$R^2 > 0.99$

log-linear · slope ≈ 0.95

Validation in process, launch planning initiated

The Comprehensive Advantage

Caris testing reveals what **targeted gene panels miss** — in both therapy eligibility and treatment outcomes.

STUDY 1



Caris Lookback Program: comprehensive testing keeps delivering value

13,293

patients newly eligible for FDA-approved targeted therapies — identified from prior results, with no additional testing

483,000+

molecular profiles

87

FDA approvals reviewed

10

tumor types

✓ **More patients matched to therapy — no re-biopsy, no new test**

STUDY 2

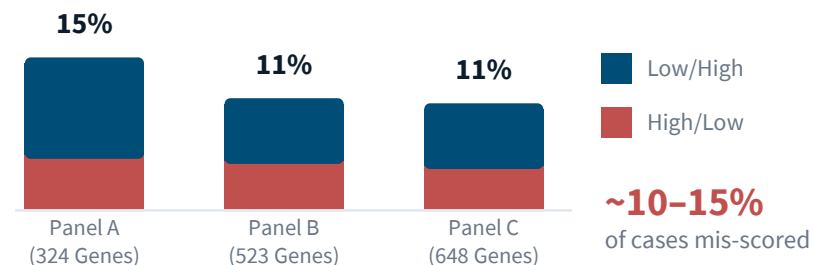


Whole-exome TMB drives better outcomes than targeted gene panels

Longer overall survival

with WES-based tumor mutational burden vs. targeted panels for pembrolizumab selection

PANEL DISCORDANCE WITH WES



✓ **Whole-exome provides more accurate TMB scoring than targeted panels**

The Oncologist, Volume 31, Issue 6, June 2026, oyag169; <https://doi.org/10.1093/oncolo/oyag169>;
Published online 29 April 2026

Cancer Immunology, Immunotherapy (2026) 75:165; <https://doi.org/10.1007/s00262-026-04365-4>;
Published online 11 May 2026

Financial Update

**Financial Overview
2026 Guidance Update**



Q2 2026 Financial Overview

TOTAL REVENUE

\$263.7M

+45% vs. Q2 2025

GROSS MARGIN⁽¹⁾

68%

+500 bps vs. Q2 2025

ADJUSTED EBITDA⁽²⁾

\$55.7M

+39.0MM improvement
vs. Q2 2025

FREE CASH FLOW⁽²⁾

\$6.4M

+\$0.5MM improvement
vs. Q2 2025

(1) Gross margin = total revenue less cost of services, divided by total revenue

(2) Adjusted EBITDA and free cash flow are non-GAAP; see earnings release for reconciliation.

Selected financial metrics

All amounts millions except gross margin

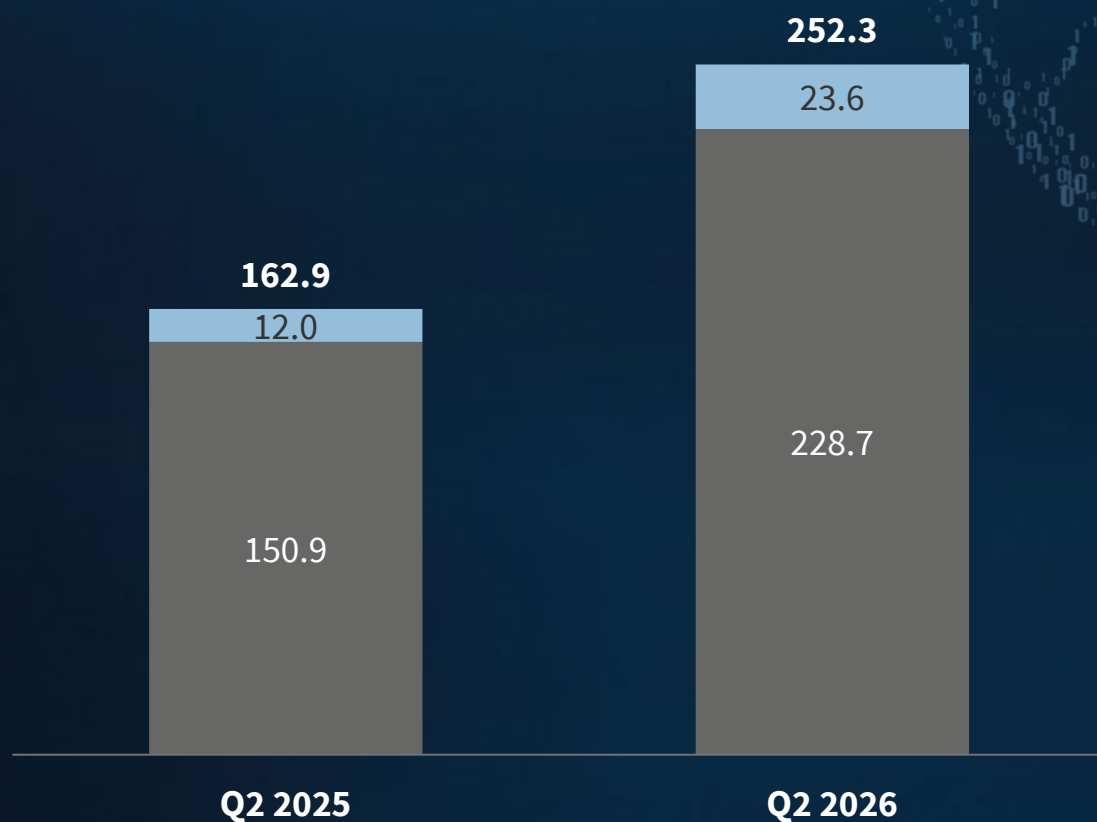
	Q2 2025	Q2 2026	Change
GAAP			
Total Revenue	\$181.4M	\$263.7M	+45%
Molecular Profiling Revenue	\$162.9M	\$252.3M	+55%
Pharma R&D Services Revenue	\$18.5M	\$11.4M	-38%
Gross Margin ⁽¹⁾	63%	68%	+500 bps
Operating Expenses	\$131.7M	\$152.7M	+\$21.0M
Net Income / (Loss)	(\$71.8M)	(\$0.6M)	+\$71.2M
NON-GAAP / CASH FLOW⁽²⁾			
Adjusted EBITDA	\$16.7M	\$55.7M	+\$39.0M
Free Cash Flow	\$5.9M	\$6.4M	+\$0.5M

Molecular Profiling Growth Driven by Volume and ASP

Molecular Profiling Services Revenue
(\$ in millions)

**+55% YoY
Total**

■ Revenue Base Accrual ■ Revenue from prior period performance obligations



ADDITIONAL HIGHLIGHTS



Blended base ASP

Surpassed \$3,850



75% MI Cancer Seek

Stable Tissue volume mix



Clinical volume +18% YoY

59,200 cases consists of ~ 114k Exome/Transcriptome tests



ChromoSeq reimbursement

\$3,228 Medicare reimbursed rate



239.5M covered lives

MI Cancer Seek



131.9M covered lives

Caris Assure

Raising full year guidance on strong Q2 performance



Total Revenue

\$1.03BN – \$1.04BN

27% – 28% YoY

Up from \$1.00–\$1.02BN on profiling strength.



Clinical Therapy Selection Volume

~20% YoY

Broad-based demand across the portfolio and continued commercial expansion.



GAAP Operating Expenses

\$595M – \$600M

20% – 21% YoY

Continued commercial expansion and further increased marketing. Up from \$590-\$595M.



Free Cash Flow / Adjusted EBITDA

Positive

While investing in expansion and catalyst pipeline during FY26, total for full year expected to be positive.

