



Unleashing Tools to Accelerate Breakthroughs in Human Health™

INVESTOR PRESENTATION
September 2024



Legal Information

Forward-looking statements

This presentation contains forward-looking statements that are subject to risks, uncertainties and other factors that could cause actual results to differ materially from those referred to in the forward-looking statements. All statements other than statements of historical fact (including statements containing the words “believes,” “plans,” “anticipates,” “expects,” “estimates,” “targets” and similar expressions) are statements that could be deemed forward-looking statements, although not all forward-looking statements contain these identifying words. Readers should not place undue reliance on these forward-looking statements. Forward-looking statements may include statements regarding financial outlook and business performance, including related to revenues, growth, margin, and operating expenses; statements regarding future financial performance and expectations, operational and strategic plans, deployment of capital, cash runway and sufficiency of cash resources, market and growth opportunity and potential, potential M&A activity, potential and ongoing restructuring plans; the potential to realize the expected benefits following the merger with SomaLogic, Inc. (“SomaLogic”), our revenue outlook for the full year 2024, and our 2026 financial targets, including with respect to revenue, non-GAAP gross margin, non-GAAP gross profit, non-GAAP operating expenses, adjusted EBITDA, cash, and free cash flow the competitive ability and position of the combined company, the success, cost and timing of the combined company’s product development, sales and marketing, and research and development activities, the combined company’s ability to obtain and maintain regulatory approval for its products, the sufficiency of the combined company’s cash, cash equivalents and short-term investments to fund operations, and any assumptions underlying any of the foregoing. Statements regarding future events are based on the parties’ current expectations and are necessarily subject to associated risks and uncertainties related to, among other things, the outcome of any legal proceedings related to the merger; risks that the anticipated benefits of the merger or other commercial opportunities may otherwise not be fully realized or may take longer to realize than expected; risks that we may not realize expected cost savings from our restructuring plans, including the anticipated decrease in operational expenses, at the levels we expect; possible restructuring and transition-related disruption, including through the loss of customers, suppliers, and employees and adverse impacts on our development activities and results of operation; restructuring activities, including our subleasing plans, customer and employee relations, management distraction, and reduced operating performance; risks that internal and external costs required for ongoing and planned activities may be higher than expected, which may cause us to use cash more quickly than we expect or change or curtail some of our plans, or both; risks that our expectations as to expenses, cash usage, and cash needs may prove not to be correct for other reasons such as changes in plans or actual events being different than our assumptions; our ability to achieve future financial targets; changes in our business or external market conditions; challenges inherent in developing, manufacturing, launching, marketing, and selling new products; interruptions or delays in the supply of components or materials for, or manufacturing of, our products; reliance on sales of capital equipment for a significant proportion of revenues in each quarter; seasonal variations in customer operations; unanticipated increases in costs or expenses; continued or sustained budgetary, inflationary, or recessionary pressures; uncertainties in contractual relationships; reductions in research and development spending or changes in budget priorities by customers; uncertainties relating to our research and development activities, and distribution plans and capabilities; potential product performance and quality issues; risks associated with international operations; intellectual property risks; and competition. Therefore, actual results may differ materially and adversely from those expressed in any forward-looking statements. For information regarding other related risks, see the “Risk Factors” section of our most recent annual report on Form 10-K filed with the SEC on March 1, 2024. We undertake no obligation to revise or update any forward-looking statements for any reason.

Non-GAAP financial information

Standard BioTools has presented certain financial information in accordance with U.S. GAAP and also on a non-GAAP basis. The non-GAAP financial measures included in this presentation are non-GAAP gross margin, non-GAAP gross profit, non-GAAP operating expenses, and adjusted EBITDA. Management uses these non-GAAP financial measures, in addition to GAAP financial measures, as a measure of operating performance because the non-GAAP financial measures do not include the impact of items that management does not consider indicative of the Company’s core operating performance. Management believes that non-GAAP financial measures, taken in conjunction with GAAP financial measures, provide useful information for both management and investors by excluding certain non-cash and other expenses that are not indicative of the Company’s core operating results. Management uses non-GAAP measures to compare the Company’s performance relative to forecasts and strategic plans and to benchmark the company’s performance externally against competitors. Non-GAAP information is not prepared under a comprehensive set of accounting rules and should only be used to supplement an understanding of the company’s operating results as reported under U.S. GAAP. Standard BioTools encourages investors to carefully consider its results under GAAP, as well as its supplemental non-GAAP information and the reconciliations between these presentations, to more fully understand its business. Reconciliations between GAAP and non-GAAP operating results are presented in the accompanying tables of this release.

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Standard BioTools products are provided for Research Use Only. Not for use in diagnostic procedures.

Executing on our Vision

Building a Diversified Life Science Tools Industry Leader



LEADING PLATFORMS AND TEAM

Leading Proteomics and Multi-omics Platforms

High content solutions in serum proteomics
spatial proteomics and single cell proteomics

Performance via SBS

Leveraging Standard BioTools Business
Systems (SBS) driving execution and profitability

Strategic M&A and Partnerships

Leveraging M&A and strategic partnerships (i.e. Illumina/
OEM partnerships) to expand sources of revenue

OPERATING AT SCALE

\$170-175MM

2024 Revenue Forecast

\$396MM

Cash, cash equivalents, and investments
(at 6/30/2024)

2026

Break-even adjusted EBITDA

Proven Leadership Team with Operational and Scientific Prowess



Michael Egholm, PhD
Chief Executive Officer
Experience:

- Danaher
- Pall
- Roche



Alex Kim
Chief Operating Officer &
Interim Chief Financial Officer
Experience:

- Danaher
- Pall
- Milliken



Sean Mackay
Chief Business Officer
Experience:

- IsoPlexis
- PhenomeX
- Lazard



Agnieszka Gallagher
SVP, Chief Legal Officer
Experience:

- Sandoz
- Alnylam
- GSK



Jeremy Davis
Chief Commercial Officer
Experience:

- Danaher
- Culligan
- CG



Stephen Williams, MD PhD
Chief Medical Officer
Experience:

- Somalogic
- Pfizer
- NIH



Betsy Jensen
Chief H.R. Officer
Experience:

- Danaher
- Gibraltar
- ITW



David King, PhD
SVP, Global R&D
Experience:

- IntegenX
- Affymetrix
- Guava



Mona Abou-Sayed
SVP, SBS
Experience:

- Mitel
- Danaher
- Tektronix



Anders Davas
SVP, Global Operations
Experience:

- Danaher
- Mölnlycke
- Telair

- Deep experience executing M&A and integration to drive profitability and growth
- Diversified life sciences tools and biopharma experience to drive progress and growth in critical end markets
- Deeply experienced multi-omic development team to provide innovation across all platform areas



Agilent Technologies



Standard BioTools™

Our Four Proteomic & Multi-Omic Platforms Uniquely Scale

Our Solutions Cover the Four Sources of Critical Biomarker Information

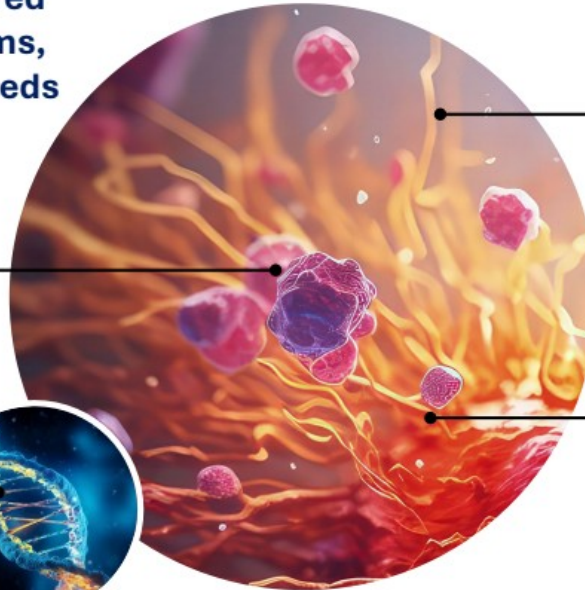
Our M&A Strategy has delivered four complementary platforms, indexed to key biopharma needs



SINGLE CELL
PROTEOMICS



GENOMICS/
MULTIOMICS



PLASMA
PROTEOMICS



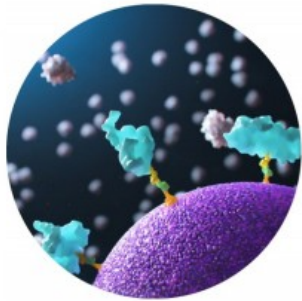
SPATIAL
PROTEOMICS

Standard BioTools Product Leadership and Growth Drivers

PROTEOMICS PLATFORMS

MULTIOMICS PLATFORM

SOMASCAN
PLASMA PROTEOME



800+

Clinical Publications

illumina[®]
Proteomics Partnership
Standard BioTools™

CYTOF
SINGLE CELL PROTEOMICS



2800+

Publications Utilizing Mass Cytometry

2600+ Mass Cytometry Active User Base

HYPERION
SPATIAL PROTEOMICS



BIOMARK
GENOMICS/ MULTIOMICS

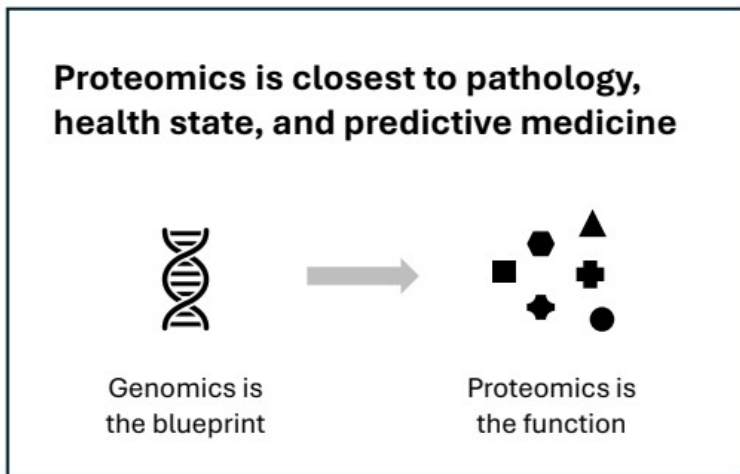


9200+

Publications in Multiomics
and Genomics

Olink
OEM Proteomics Partnership

Our Proteomic Solutions are Key to the Highest Growth Therapeutic Areas



12,800+ proteomic and multiomic studies

SOMASCAN
PLASMA PROTEOME

HYPERION
SPATIAL PROTEOMICS

CYTOF
SINGLE CELL PROTEOMICS

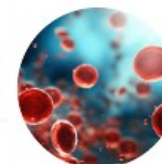
BIOMARK
GENOMICS/ MULTIOMICS



Leading published clinical trial studies in below growth areas

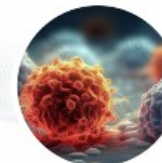
3,937 Trials¹

Cardiometabolic, GLP-1s



15,755 Trials¹

Oncology



8,655 Trials¹

Immunology



¹ clinical trials database
Standard BioTools™

Our High Content Proteomics Uniquely Scale to Meet Customer Needs

Widest coverage and consistency is driving our collective leadership in translational medicine

SOMASCAN PLASMA PROTEOME

Widest Coverage & Highest Precision

3x more proteins
vs ANTIBODY BASED APPROACH

<0.5x coefficient of variations (CVs)

CYTOF SINGLE CELL PROTEOMICS

Widest Coverage & Highest Multiplex

5-10x more functional proteins
vs FLOW

2x more markers

HYPERION SPATIAL PROTEOMICS

Widest Dynamic Range & Highest Throughput

10-15x more simultaneous markers
vs CYCLIC IMMUNOFLUORESCENCE

35-100x more throughput



High Content
Cardiometabolic
Biomarkers



High Content
Immuno-oncology
Biomarkers

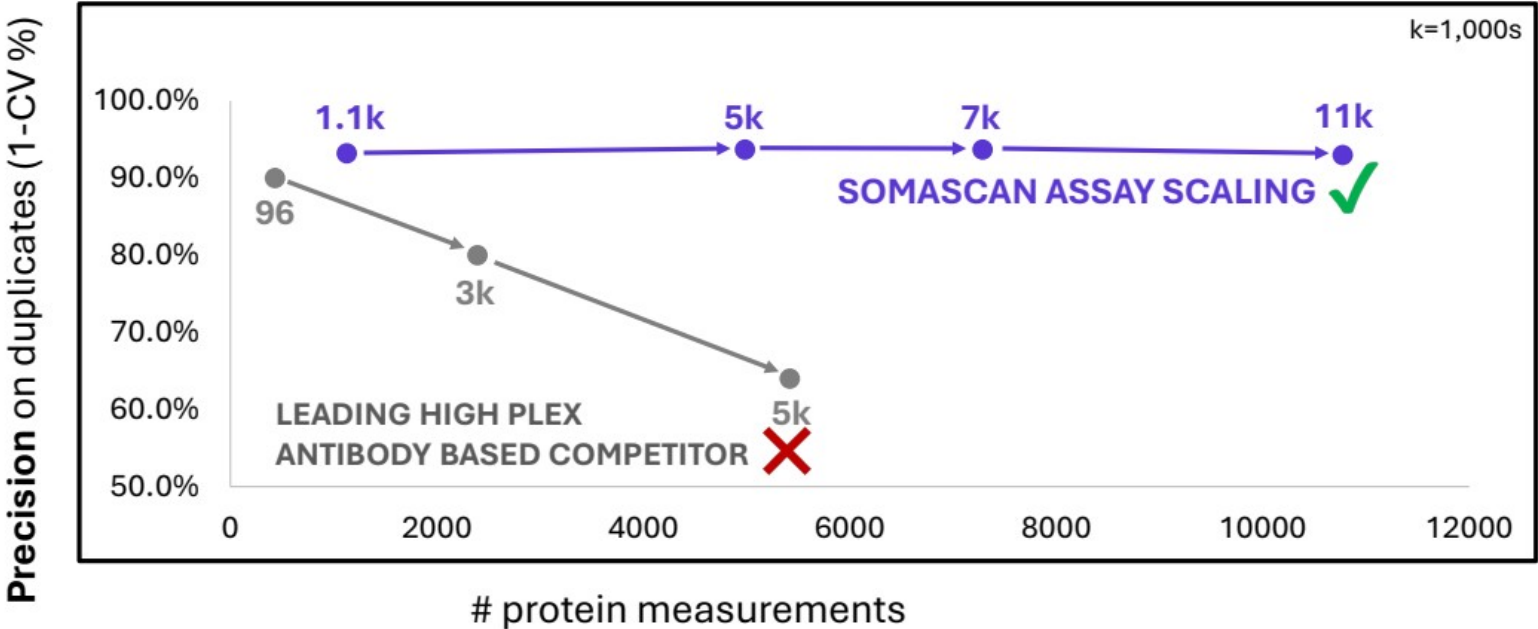


High Content
Immuno-oncology
Biomarkers

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2024 Data: SomaScan is the Only Proteomic Approach that Scales

Greatest Precision Moat: SomaScan is the only plasma proteomics technology that scales



Rooney et al. Plasma proteomic comparisons change as coverage expands for SomaLogic and Olink. MedRxiv Preprint. 2024
Standard BioTools™

SomaScan Leadership in GLP-1 / Cardiometabolic and Key Growth Areas

Large studies, qualifying biomarkers, over 10 years create key moats

Novo Nordisk¹

Prevalence of, and effect of semaglutide on, features of non-alcoholic steatohepatitis using SomaSignal tests in patients with obesity with/without type 2 diabetes, and correlation of SomaSignal tests with histology in patients with non-alcoholic steatohepatitis: analysis of data from three randomised trials

Novo M. Schattenberg¹, Henning Grundbois², Eric Kliewt³, Steen Lakkundi⁴, Michelle T. Longhi⁵, Sara Doric Negishi⁶, Jean S. Capparelli⁷, Andrew Dawson⁸

Semaglutide reduced the odds of having NASH components in populations with overweight/obesity, as measured by SomaSignal models

Table 1. Trial design

Parameter	Novo Nordisk	Novo Nordisk	Novo Nordisk
Population	Obese with/without T2D	Obese with/without T2D	Obese with/without T2D
Intervention	Semaglutide	Placebo	Placebo
Duration	12 weeks	12 weeks	12 weeks
Primary endpoint	Change in liver enzymes	Change in liver enzymes	Change in liver enzymes

Table 2. Baseline prevalence of NASH and NASH components based on SomaSignal models

Parameter	Novo Nordisk	Novo Nordisk	Novo Nordisk
NASH	11%	11%	11%
NAFLD	36%	36%	36%
Steatosis	12%	12%	12%
Steatosis without NASH	25%	25%	25%
Steatosis with NASH	11%	11%	11%
NASH without steatosis	0%	0%	0%
NASH with steatosis	11%	11%	11%

Table 3. Odds ratios for changes in NASH components

Parameter	Novo Nordisk	Novo Nordisk	Novo Nordisk
NASH	0.48 (0.18-1.28)	0.48 (0.18-1.28)	0.48 (0.18-1.28)
NAFLD	0.85 (0.45-1.65)	0.85 (0.45-1.65)	0.85 (0.45-1.65)
Steatosis	0.65 (0.35-1.25)	0.65 (0.35-1.25)	0.65 (0.35-1.25)
Steatosis without NASH	0.75 (0.40-1.45)	0.75 (0.40-1.45)	0.75 (0.40-1.45)
Steatosis with NASH	0.55 (0.25-1.25)	0.55 (0.25-1.25)	0.55 (0.25-1.25)
NASH without steatosis	0.45 (0.15-1.25)	0.45 (0.15-1.25)	0.45 (0.15-1.25)
NASH with steatosis	0.55 (0.25-1.25)	0.55 (0.25-1.25)	0.55 (0.25-1.25)

Figure 1. Odds ratios for changes in NASH components

Table 4. Odds ratios for changes in liver enzymes

Parameter	Novo Nordisk	Novo Nordisk	Novo Nordisk
ALT	0.95 (0.85-1.05)	0.95 (0.85-1.05)	0.95 (0.85-1.05)
AST	0.95 (0.85-1.05)	0.95 (0.85-1.05)	0.95 (0.85-1.05)
ALP	0.95 (0.85-1.05)	0.95 (0.85-1.05)	0.95 (0.85-1.05)
Gamma-GT	0.95 (0.85-1.05)	0.95 (0.85-1.05)	0.95 (0.85-1.05)

Conclusion

Semaglutide significantly reduced the odds of having NASH components in populations with overweight/obesity, as measured by SomaSignal models. This effect was observed across all NASH components and was independent of changes in liver enzymes.

Schattenberg et al. Journal of Hepatology 78:S811-S812. June 2023

Gilead¹

Utility of SomaSignal[®] Panels for Drug Response and Monitoring Disease Progression in Patients with Advanced Fibrosis Due to Nonalcoholic Steatohepatitis

Raja S. Kanbay¹, Raji Zhai², Jun Xu², Jason H. Meekins³, Lisa Boyette⁴, Timothy R. Williams⁵, Shariene Lev⁶, Vlad Rabkin⁷, Andrew R. Brown⁸, Mazen Hourani⁹, Ronit Liorat¹⁰

Key Findings

- SomaSignal panels for serum biomarkers for nonalcoholic steatohepatitis (NASH) are significantly associated with histologically confirmed NASH.
- SomaSignal panels for serum biomarkers for nonalcoholic steatohepatitis (NASH) are significantly associated with histologically confirmed NASH.

Conclusions

The data demonstrate the potential for using personalized NASH SomaSignal[®] panels for assessing longitudinal disease response in NASH patients.

Introduction

Nonalcoholic steatohepatitis (NASH) is a chronic liver disease characterized by hepatic steatosis, hepatocellular injury, inflammation, and fibrosis. The pathogenesis of NASH is complex and involves genetic, metabolic, and environmental factors. Current treatment options are limited, and there is a need for novel therapies and biomarkers for NASH diagnosis, prognosis, and monitoring of treatment response.

Objective

To evaluate the utility of SomaSignal panels for drug response and monitoring disease progression in patients with advanced fibrosis due to NASH.

Methods

A total of 100 patients with advanced fibrosis due to NASH were enrolled in a phase 2 clinical trial. The trial was designed to evaluate the efficacy of a novel drug in treating NASH. SomaSignal panels were used to monitor disease progression and treatment response in these patients.

Results

SomaSignal panels were significantly associated with histologically confirmed NASH. The panels were also associated with changes in liver enzymes and other biomarkers of NASH. These findings suggest that SomaSignal panels may be useful for monitoring disease progression and treatment response in patients with advanced fibrosis due to NASH.

Kowdley et al. EASL June 2023

Greatest Study Moat: 10 years of major cardiometabolic and oncology studies sponsored by pharma companies such as Novo Nordisk, BMS, Gilead, etc. have driven efforts to significantly accelerate cardiometabolic trials

¹Listed firms are authors on Somascan studies
Standard BioTools™

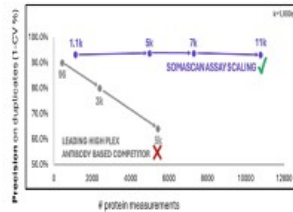
Data Moat + Our Illumina Partnership Create Growth Opportunity

Integrating SomaLogic's products with Illumina's sequencing technologies, informatics toolsets and software

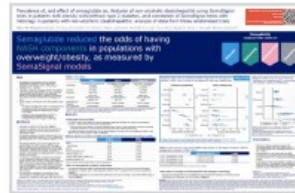
Illumina Partnership



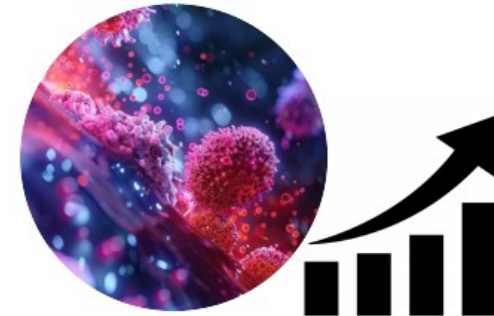
+



+



Growth Opportunity



2000+
Installed base
of NovaSeq
Instruments

**Only Scalable
Solution on the
Market Today**

**Leading
Biomarker
Studies**

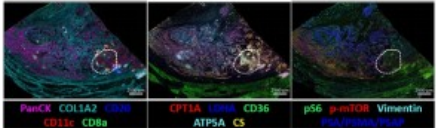
**Large proteomic translational
opportunity over the next decade**

Standard BioTools™

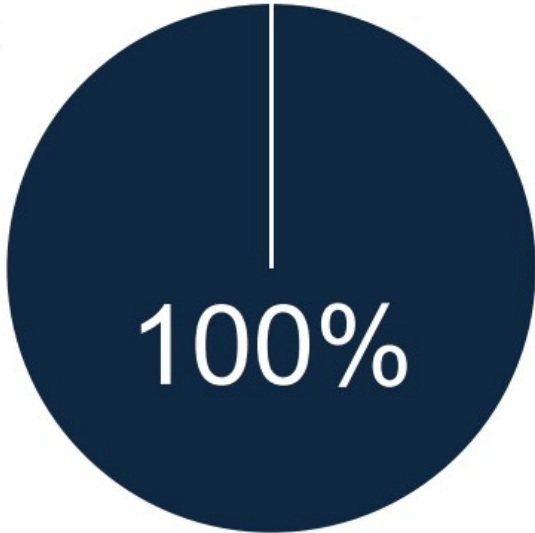
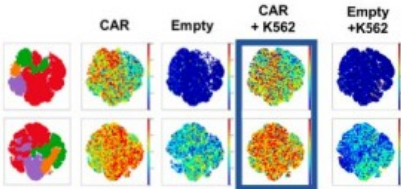
2024 Single Cell Proteomics & Spatial Proteomics Innovation



'24: Hyperion launch – with up to 100x throughput over other technologies



'22 CyTOF XT launch: Approaching walkaway operations



- ✓ KEYTRUDA[†] (pembrolizumab)
- ✓ OPDIVO[†] (nivolumab)
- ✓ YERVOY[†] (ipilimumab)
- ✓ TECENTRIQ[†] (atezolizumab)
- ✓ DARZALEX[†] (daratumumab)

100% of the top 10 cancer immunotherapies¹ have leveraged unique biomarkers in trials, found by mass cytometry

1. Top cancer immunotherapies by revenue in 2023. [†]Therapies listed in peer reviewed publications Standard BioTools™

CyTOF Uniquely Scales with Critical Biomarkers for Oncology

Studies using conventional flow panels would have missed the critical differentiating markers

Conventional Panel in Spectral Fluorescence*

25 surface markers*

Additional Simultaneous Coverage only on CyTOF

+25 intracellular/functional markers



Biomarker for Identification of Drug Targets in MM Trial

BMS† | nivolumab

Predictive Biomarker of Response in Combination Nivo Phase 2 Trial

MERCK† | pembrolizumab

Predictive Biomarker of Survival in Pembro Phase 1 Trial

NOVARTIS† | CAR-T

Predictive Biomarker of Response in CAR-T

Unique Predictive Biomarker

*Competitor 25-color immunoprofiling assay Standard BioTools™ †Pharma listed are the firms that manufacture and/ or market the therapies in the published studies

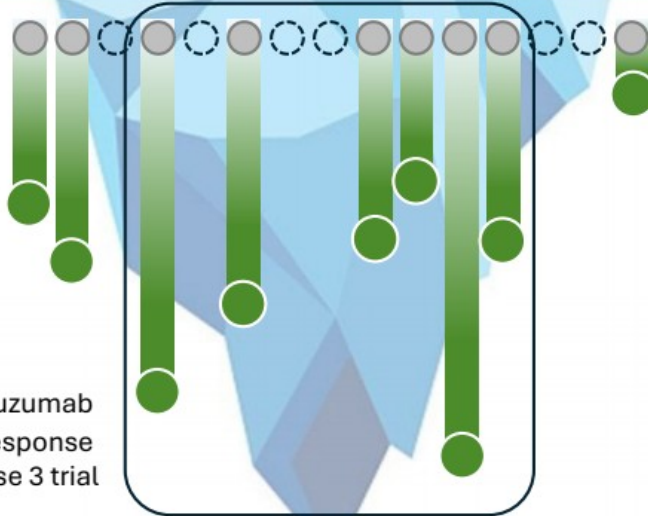
Spatial Proteomics Scales in Dynamic Range, Multiplex, and Throughput

Cyclic Fluorescence is missing dynamic range

Highest Dynamic Range & Highest Multiplex



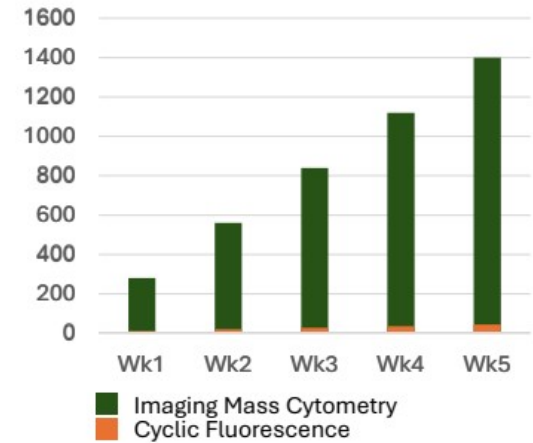
Imaging Mass Cytometry captures necessary dynamic range: i.e. Highest data quality



+

Highest Throughput

Cumulative Samples Per Week



GENENTECH† | trastuzumab
 Predictive biomarker of patient response stratification in trastuzumab phase 3 trial

†Pharma listed are the firms that manufacture and/or market the therapies in the published studies
 Standard BioTools™

Standard BioTools Business System: Creates Flywheel Through M&A

SBS Practice	Impact
LEAN TRANSFORMATION Build awareness of opportunities to eliminate excess/ waste and achieve the best possible efficiency	GAIN OPERATING EFFICIENCY Reduce cost per unit, increase gross margins, and reduce operating cash use
CUSTOMER CENTRICITY Ensure delivering value to the customer is at the center of everything we do	CUSTOMER GROWTH Provide valuable proteomic and multi-omic solutions and insights to impact a growing customer base
KAIZEN CULTURE Facilitate continuous improvement (Kaizen) through standard work to ensure maximum quality and efficiency	GROW PROFITABLY Drive daily execution with quality systems, improving operational excellence around a leading suite of products

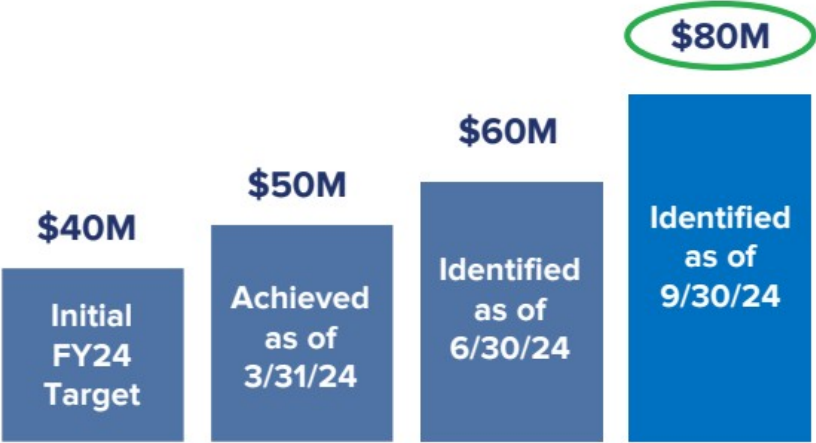


SBS

culture of continuous improvement to achieve world class operational excellence and exceptional customer value

Capturing \$80M Synergy Opportunity

Non-GAAP OpEx Reduction Expected to Reach \$80M in 2024



- Able to operationalize synergies ahead of schedule as initial target goal was to realize by YE 2024
- This preserves investments for future growth opportunities

Revenue

Continued Focus on Diversifying Revenue Mix

Pro Forma Combined	H1 2024	YOY
Instruments	\$12M	(32%)
Consumables	\$19M	(11%)
Service & Other	\$52M	(4%)
TOTAL	\$83M	(11%)

Pro Forma Combined	H1 2024	YOY
Proteomics (Px)	\$65M	(12%)
Genomics (Gx)	\$18M	(6%)
TOTAL	\$83M	(11%)

- YTD revenue declined due to continued pressure from macroeconomic headwinds
- SomaScan Assay services business down over 2023 driven primarily by the timing of large projects
- SomaScan Assay Kits business up over 2023 from continued expansion in authorized sites and related pull-through
- Illumina early access program underway and on track for 2025 full commercial release
- While macroeconomic conditions have continued to be a near-term headwind, our instrument revenue improved sequentially; pipeline remains robust/building
- Consumables and services in both Px and Gx impacted by prior year declines in legacy installed base; new installations expected to expand pull-through in late 2024 and beyond

Reflects combined historical information, assuming the merger closed on January 1, 2023. | Numbers may not add, and percentages may not foot due to rounding. All amounts presented are non-GAAP

Gross Margin (Non-GAAP)

Executing Roadmap to Expanded Gross Margin Profile

Pro Forma Combined	Q2 2024	YoY	H1 2024	YoY
Non-GAAP Gross Profit \$	\$17M	(\$9M)	\$43M	(\$6M)
Non-GAAP Gross Margin %	45%	(840 bps)	51%	(124 bps)

- ~250 bps impact in Q2 from lower capacity utilization related to decrease in SomaScan assay services volumes
- ~300 bps impact in Q2 from strategic decisions to replace or upgrade instruments in the field

NON-GAAP GROSS MARGIN EXPANSION OPPORTUNITY

2026 Target GM Profile:

Mid 60%s

+ Continued deployment of SBS / Lean principles

+ Sales growth
+ Product mix shift

+ Overhead absorption
+ Reduced replacement and upgrade costs

Reflects combined historical information with certain adjustments, assuming the merger closed on January 1, 2023. | Non-GAAP gross margin excludes amortization of developed technology, non-cash stock-based compensation, and depreciation and amortization. Refer to Appendix for a reconciliation between GAAP and non-GAAP gross margin. | Numbers may not add, and percentages may not foot due to rounding. All amounts presented are non-GAAP

Standard BioTools™



Appendix

2024 Nature correction: SomaScan leads in precision/# of proteins

Section 1 and Figure 1 in BioRx preprint and Nature paper were both corrected

Article | [Open access](#) | Published: 04 October 2023

Large-scale plasma proteomics comparisons through genetics and disease associations

[Grimur Hjorleifsson Eldjarn](#), [Egil Ferkingstad](#), [Sigrun H. Lund](#), [Hannes Helgason](#), [Olafur Th. Magnusson](#), [Kristbjorg Gunnarsdottir](#), [Thorunn A. Olafsdottir](#), [Bjarni V. Halldorsson](#), [Pall I. Olason](#), [Florian Zink](#), [Sigurjon A. Gudjonsson](#), [Gardar Sveinbjornsson](#), [Magnus I. Magnusson](#), [Agnar Helgason](#), [Asmundur Oddsson](#), [Gisli H. Halldorsson](#), [Magnus K. Magnusson](#), [Saedis Saevarsdottir](#), [Thjodbjorg Eiriksdottir](#), [Gisli Masson](#), [Hreinn Stefansson](#), [Ingilei](#)

Nature 622, 348–357 (2023) | [View article](#) | [Download PDF](#) | [Full text access](#)

56k Accesses | 3 Citations

An Author Correction | [View](#)

This article | [View](#)

Count

Assay CV relative to population

— Olink
— SomaScan

“Count” label misleading: not # of proteins measured

Author Correction | [Open access](#) | Published: 22 May 2024

Author Correction: Large-scale plasma proteomics comparisons through genetics and disease associations

[Grimur Hjorleifsson Eldjarn](#), [Egil Ferkingstad](#), [Sigrun H. Lund](#), [Hannes Helgason](#), [Olafur Th. Magnusson](#), [Kristbjorg Gunnarsdottir](#), [Thorunn A. Olafsdottir](#), [Bjarni V. Halldorsson](#), [Pall I. Olason](#), [Florian Zink](#), [Sigurjon A. Gudjonsson](#), [Gardar Sveinbjornsson](#), [Magnus I. Magnusson](#), [Agnar Helgason](#), [Asmundur Oddsson](#), [Gisli H. Halldorsson](#), [Magnus K. Magnusson](#), [Saedis Saevarsdottir](#), [Thjodbjorg Eiriksdottir](#), [Gisli Masson](#), [Hreinn Stefansson](#), [Ingilei](#)

Nature 622, 348–357 (2023) | [View article](#) | [Download PDF](#) | [Full text access](#)

3388 Accesses | 1 Citation

This article | [View](#)

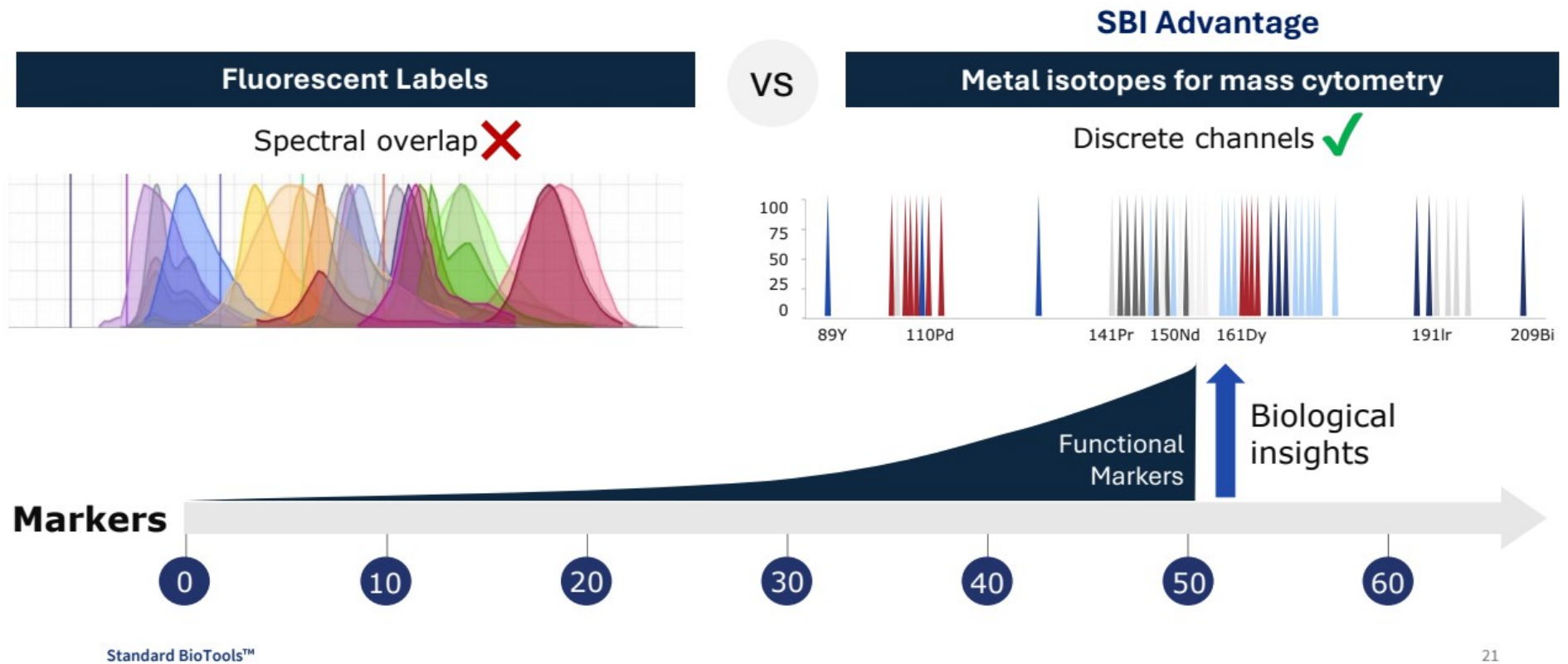
Fraction of assays

Assay CV

— Olink
— SomaScan

High-Parameter Cell Analysis is a Challenge with Proteins

Mass cytometry solves fundamental limitation of fluorescence-based cell analysis

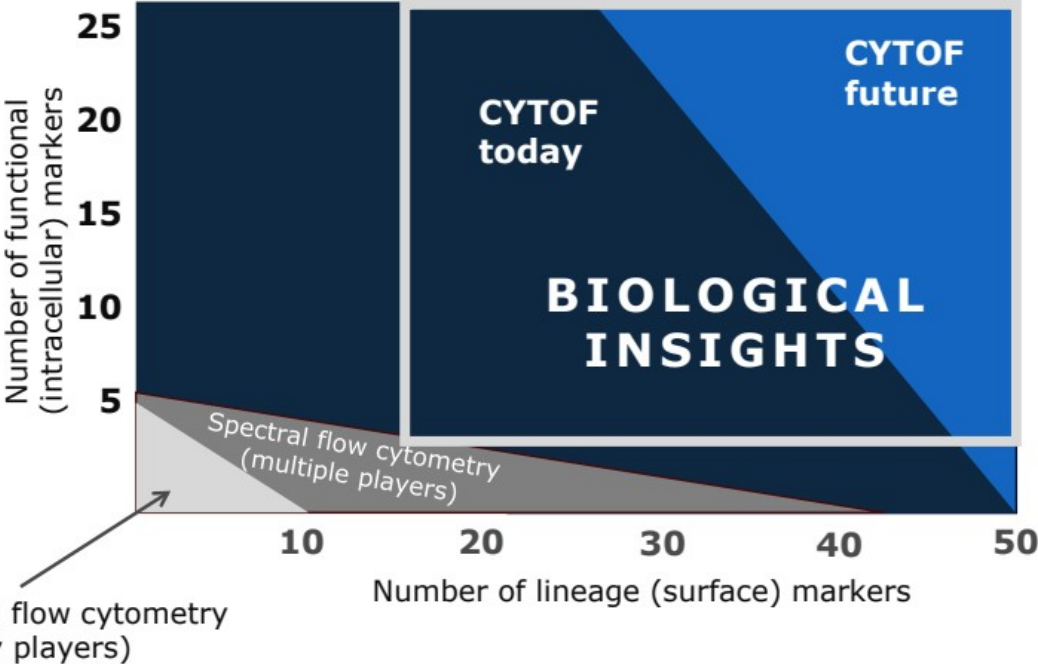


CyTOF Uniquely Scales for Translational Research

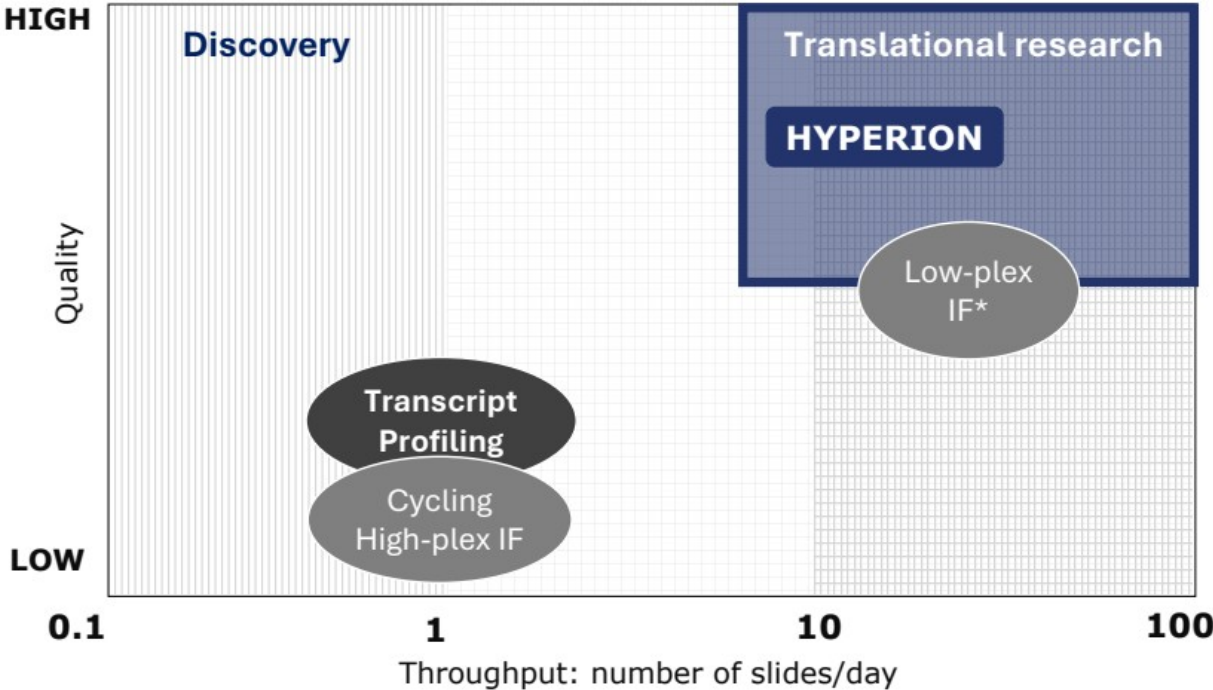
The most robust solution in high-parameter flow cytometry market segment



SBI Advantage
No limit to how many different markers can be detected at once



Spatial Biology: Hyperion System is a Game-Changer



SBI Advantage
 40 Slides | 40 Markers
 24 Hours



Standard BioTools™

*IF: immunofluorescence

Non-GAAP Reconciliation

Gross Margin

	As Reported		Pro Forma	
	Q2 2024	Q2 2023	Q2 2024	Q2 2023
GAAP Gross Profit (\$M)	\$14.9	\$13.6	\$14.9	\$21.5
Add: Amortization of Acquired Intangible Assets	\$0.6	\$2.8	\$0.6	\$3.4
Add: Depreciation and Amortization in COGS	\$1.0	\$0.3	\$1.0	\$0.7
Add: Stock-Based Comp in COGS	\$0.3	\$0.1	\$0.3	\$0.2
Add: Restructuring in COGS	\$0.0	\$0.0	\$0.0	\$0.0
Add: Cost of Sales Adjustment	\$0.0	\$0.0	\$0.0	\$0.0
Non-GAAP Gross Profit	\$16.7	\$16.8	\$16.7	\$25.7
GAAP Gross Margin	40.1%	49.2%	40.1%	44.6%
Add: Amortization of Acquired Intangible Assets	1.5%	10.1%	1.5%	7.0%
Add: Depreciation and Amortization in COGS	2.6%	1.2%	2.6%	1.5%
Add: Stock-Based Comp in COGS	0.8%	0.4%	0.8%	0.3%
Add: Restructuring in COGS	0.0%	0.0%	0.0%	0.0%
Add: Cost of Sales Adjustment	0.0%	0.0%	0.0%	0.0%
Non-GAAP Gross Margin	45.0%	60.9%	45.0%	53.4%

Figures are derived from Condensed Consolidated Statements of Operations as reported in the Company's Reports on Form 10-Q for the relevant periods. | Numbers may not add, and percentages may not foot due to rounding.

Non-GAAP Reconciliation - YTD

Gross Margin

	As Reported		Pro Forma	
	Q2 YTD 2024	Q2 YTD 2023	Q2 YTD 2024	Q2 YTD 2023
GAAP Gross Profit (\$M)	\$39.1	\$25.9	\$37.7	\$41.8
Add: Amortization of Acquired Intangible Assets	\$2.5	\$5.6	\$2.5	\$6.7
Add: Depreciation and Amortization in COGS	\$2.0	\$0.7	\$2.0	\$1.4
Add: Stock-Based Comp in COGS	\$0.5	\$0.5	\$0.5	\$0.6
Add: Restructuring in COGS	\$0.0	\$0.0	\$0.0	\$0.0
Add: Cost of Sales Adjustment	(\$1.8)	\$0.0	\$0.0	(\$1.3)
Non-GAAP Gross Profit	\$42.3	\$32.6	\$42.7	\$49.1
GAAP Gross Margin	47.2%	49.0%	45.2%	44.6%
Add: Amortization of Acquired Intangible Assets	3.0%	10.6%	3.0%	7.2%
Add: Depreciation and Amortization in COGS	2.4%	1.2%	2.4%	1.5%
Add: Stock-Based Comp in COGS	0.6%	0.9%	0.6%	0.6%
Add: Restructuring in COGS	0.0%	0.0%	0.0%	0.0%
Add: Cost of Sales Adjustment	(2.2%)	0.0%	0.0%	(-1.4%)
Non-GAAP Gross Margin	51.1%	61.7%	51.2%	52.5%

Figures are derived from Condensed Consolidated Statements of Operations as reported in the Company's Reports on Form 10-Q for the relevant periods. | Numbers may not add, and percentages may not foot due to rounding.