



SCALING Spatial Biology to
Improve **PATIENT CARE**

Disclaimers

Cautionary Note Regarding Forward-Looking Statements

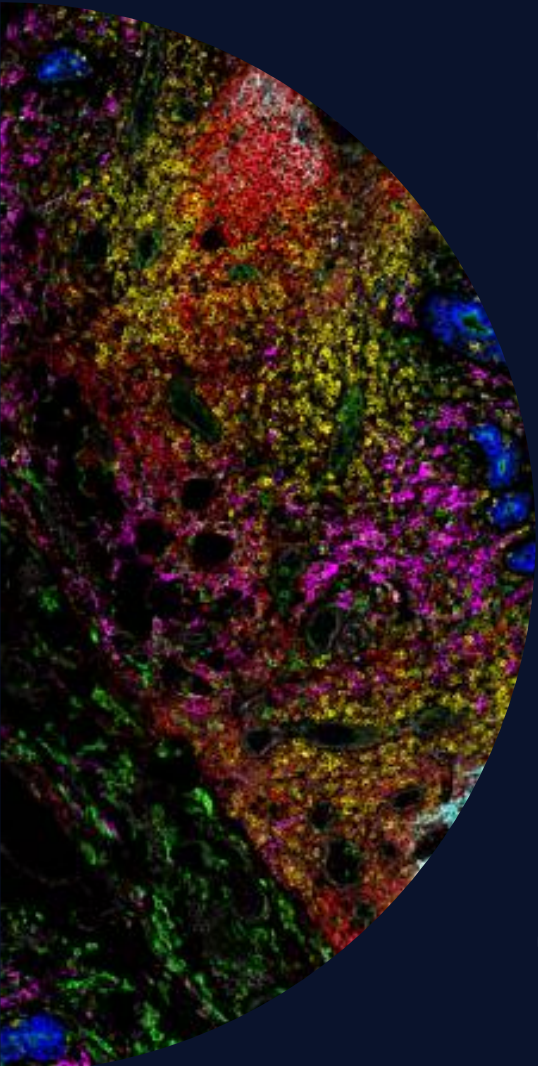
This presentation includes express and implied “forward-looking statements” that are based on management’s beliefs and assumptions and on information currently available to management. All statements contained in this presentation other than statements of historical fact are forward-looking statements, including statements regarding our ability to develop, commercialize and achieve market acceptance of our current and planned products and services, the potential of our current and planned products and services, projected spatial biology market growth, our research and development efforts, our expectations regarding our current and potential partnerships and collaborations, revenues and earnings projections, growth prospects,, and other matters regarding our future performance, market opportunities, business strategies, use of capital, results of operations and financial position, and plans and objectives for future operations. In some cases, you can identify forward-looking statements by terms such as “anticipate,” “estimate,” “expect,” “intend,” “may,” “might,” “plan,” “project,” “will,” “would,” “should,” “could,” “can,” “believe,” “predict,” “potential,” “continue,” “ongoing” or the negative of these terms, and similar expressions intended to identify forward-looking statements. However, not all forward-looking statements contain these identifying words. By their nature, these statements are subject to numerous risks and uncertainties, including factors beyond our control, that could cause actual results, performance or achievement to differ materially and adversely from those anticipated or implied in the statements. For further information regarding these risks, uncertainties and other factors, you should read the “Risk Factors” section of our Quarterly Report on Form 10-Q filed for the period ended September 30, 2024 and our Annual Report on Form 10-K filed for the period ended December 31, 2023 and other documents we file with the Securities and Exchange Commission from time to time. You should not rely upon forward-looking statements as predictions of future events. Although our management believes that the expectations reflected in our statements are reasonable, we cannot guarantee that the future results, levels of activity, performance or events and circumstances described in the forward-looking statements will be achieved or occur. Moreover, neither we, nor any other person, assumes responsibility for the accuracy and completeness of these statements. Recipients are cautioned not to place undue reliance on these forward-looking statements, which speak only as of the date such statements are made and should not be construed as statements of fact. We undertake no obligation to update these forward-looking statements to reflect events or circumstances after the date hereof, to reflect the occurrence of unanticipated events or for any other reason, except as required by law.

Market Industry Data

Projections, estimates, industry data and information contained in this presentation, including the Company’s general expectations and market position and market opportunity, are based on information from third-party sources and management estimates. Although the Company believes that its third-party sources are reliable, the Company cannot guarantee the accuracy or completeness of its sources. The Company’s estimates are derived from third-party sources, publicly available information, the Company’s knowledge of its industry and assumptions based on such information and knowledge. The Company’s estimates have not been verified by any independent source. All of the projections, estimates, market data and industry information used in this presentation involve a number of assumptions and limitations, and you are cautioned not to give undue weight to such information. In addition, projections, estimates and assumptions relating to the Company’s and its industry’s future performance are necessarily subject to a high degree of uncertainty and risk due to a variety of factors, including, but not limited to, those described above, that could cause future performance to differ materially from the Company’s expressed projections, estimates and assumptions or those provided by third parties.

Akoya is Leading the Spatial Biology Revolution

Transforming Discovery to Diagnostics



Best-in-class platforms

Fastest and most robust spatial biology platforms with whole-slide and single-cell imaging



Complete end-to-end solutions

Instruments, reagents, software and services



Emerging clinical platform for next generation patient care

Expanding clinical partnerships to drive precision medicine and companion diagnostics



Established market leader with largest installed base

1,299 instruments installed worldwide*

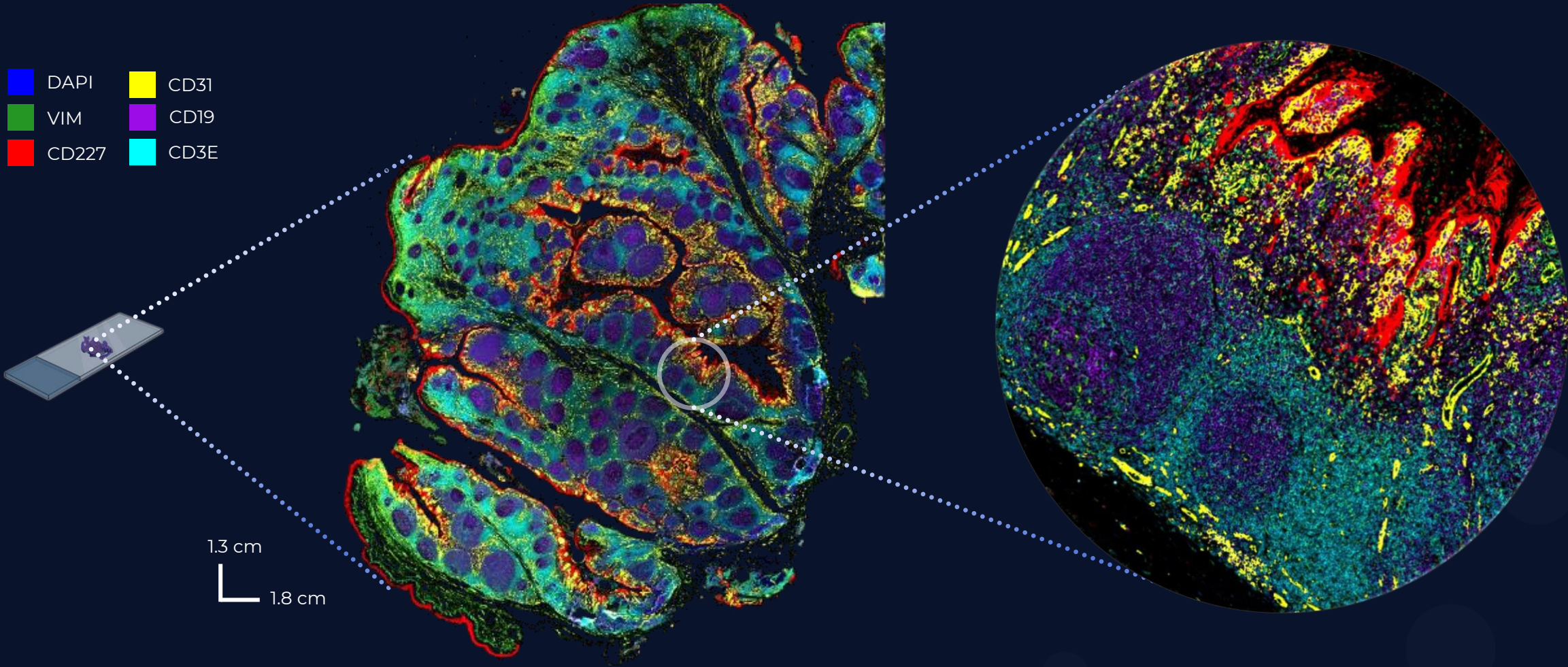


Greatest number of high-impact publications

1,578 total publications citing Akoya's technologies*

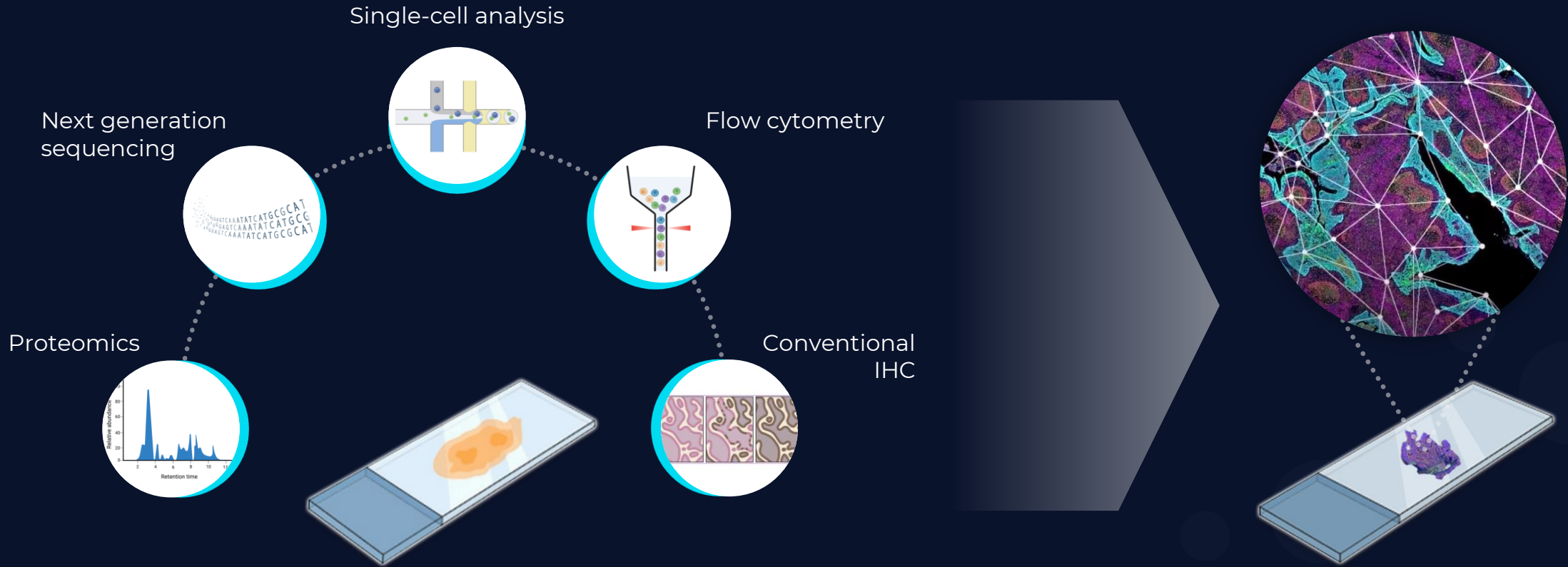
Akoya's Spatial Biology Platforms - Transforming Tissue Analysis

Rapidly Mapping Whole Tissue at Single-cell and Subcellular Resolution



Identifying the **spatial patterns and relationships** that drive disease biology and response to therapy

Current Tissue Analysis Methods Migrating to Spatial



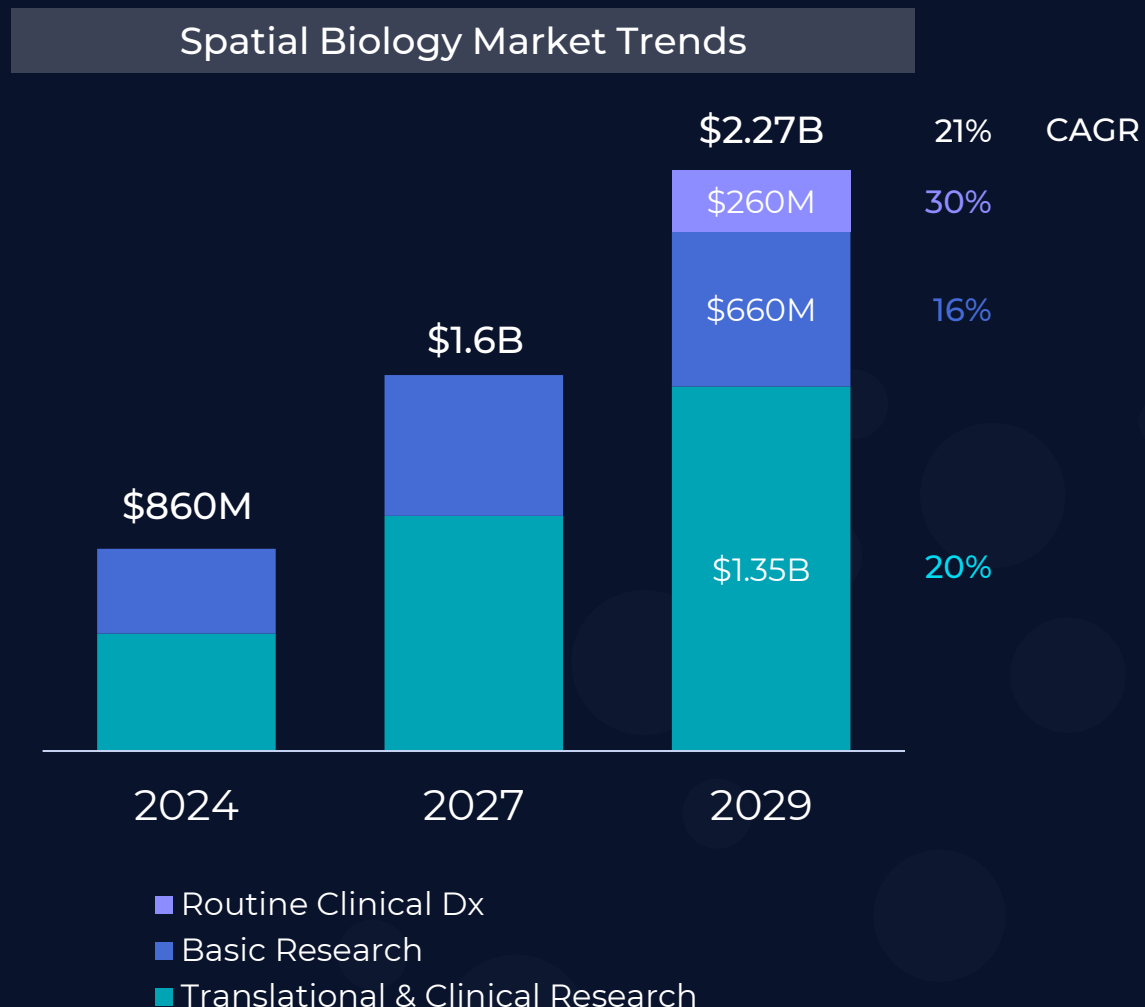
Current tissue analysis methods deliver no or very limited spatial information while destroying the tissue

Spatial Phenotyping: Understanding biomarkers in tissue context while preserving the tissue

Drivers of Spatial Biology Market Growth

Translational, Clinical Research and Routine Dx Estimated to be ~70% of Market in 5 Years

- Spatial biology market expected to grow 21% annually over the next 5 years with spatial proteomics projected to surpass spatial transcriptomics in size
- Translational & clinical research expected to make up the largest market segment as spatial moves into later stage development
- Routine clinical Dx expected to be the fastest growing market segment
- Multi-plex immunofluorescence (mIF) proteomics approaches are expected to accelerate growth more than any other spatial technology



Akoya's Complete End-to-End Spatial Solutions

Discovery

Biomarker discovery with high-plex panels

Translational

Biomarker validation with high-throughput targeted panels

Clinical

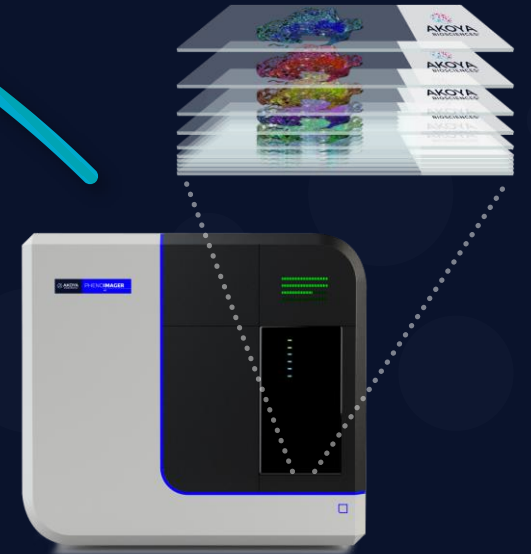
Large-scale studies and diagnostics



PhenoCycler-Fusion

Unique 2-in-1 spatial platform enabling **HIGH-TO-LOW PLEX** AND **HIGH-THROUGHPUT** whole-slide imaging

Only **CLINICAL-GRADE** spatial platform with proven **ROBUSTNESS, REPRODUCIBILITY** and **SENSITIVITY**

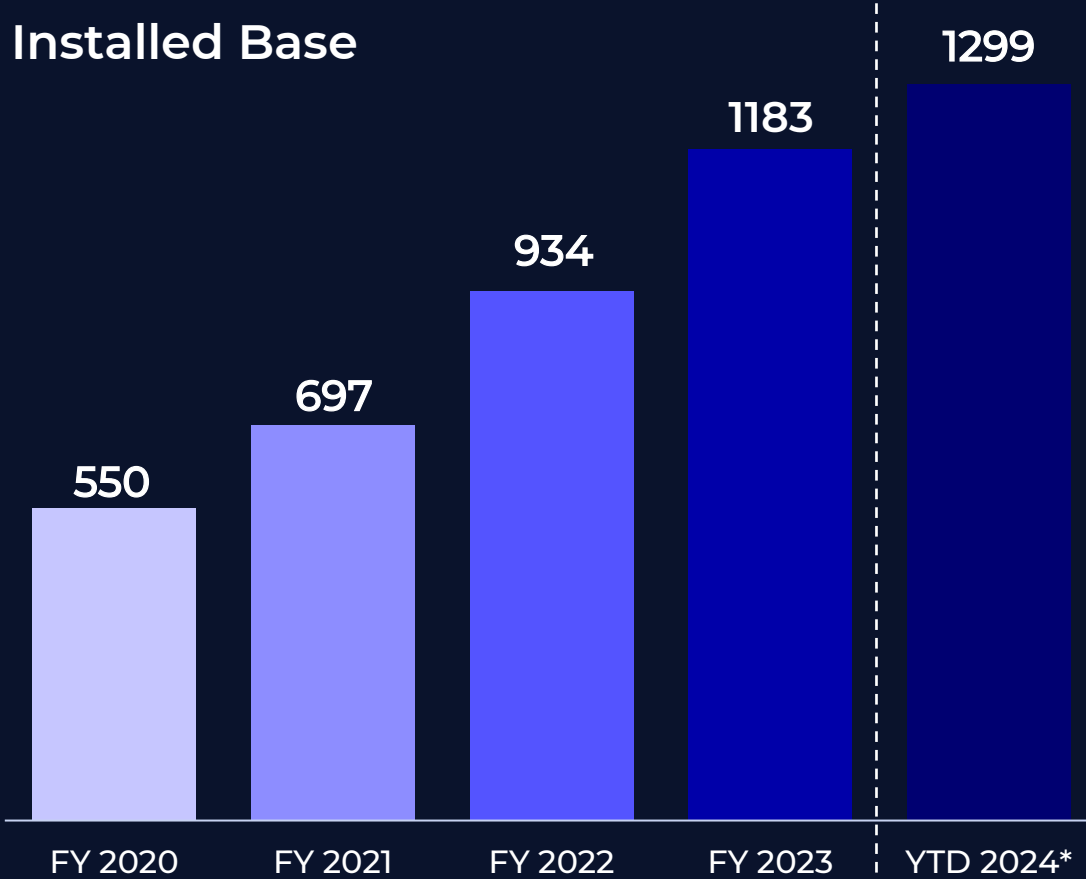


PhenoImager HT

Largest and Rapidly Growing Installed Base in the Industry

Products Across Discovery, Translational, and Clinical Markets

Installed Base



Installed base of
1299
Owning the biomarker journey...



PhenoCycler-Fusion



Phenolmager HT

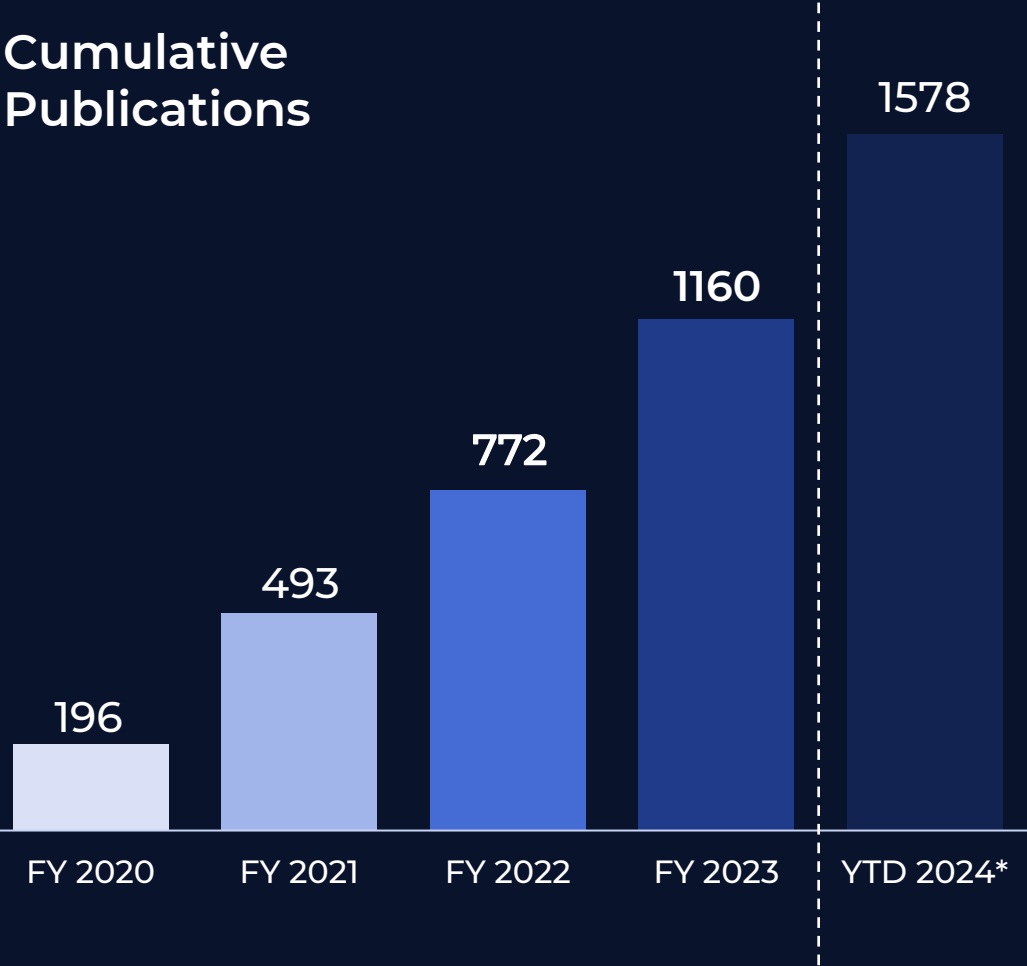


* As of September 30, 2024

Accelerating and Market Leading Publication Volume

Akoya's Technology Consistently Featured in Leading Journals for Groundbreaking Findings

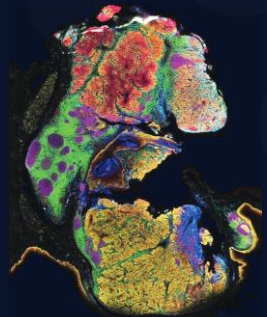
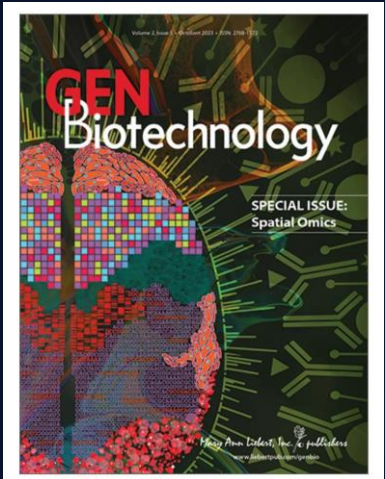
Cumulative Publications



1578
total publications featuring Akoya's technology



Featured Publication

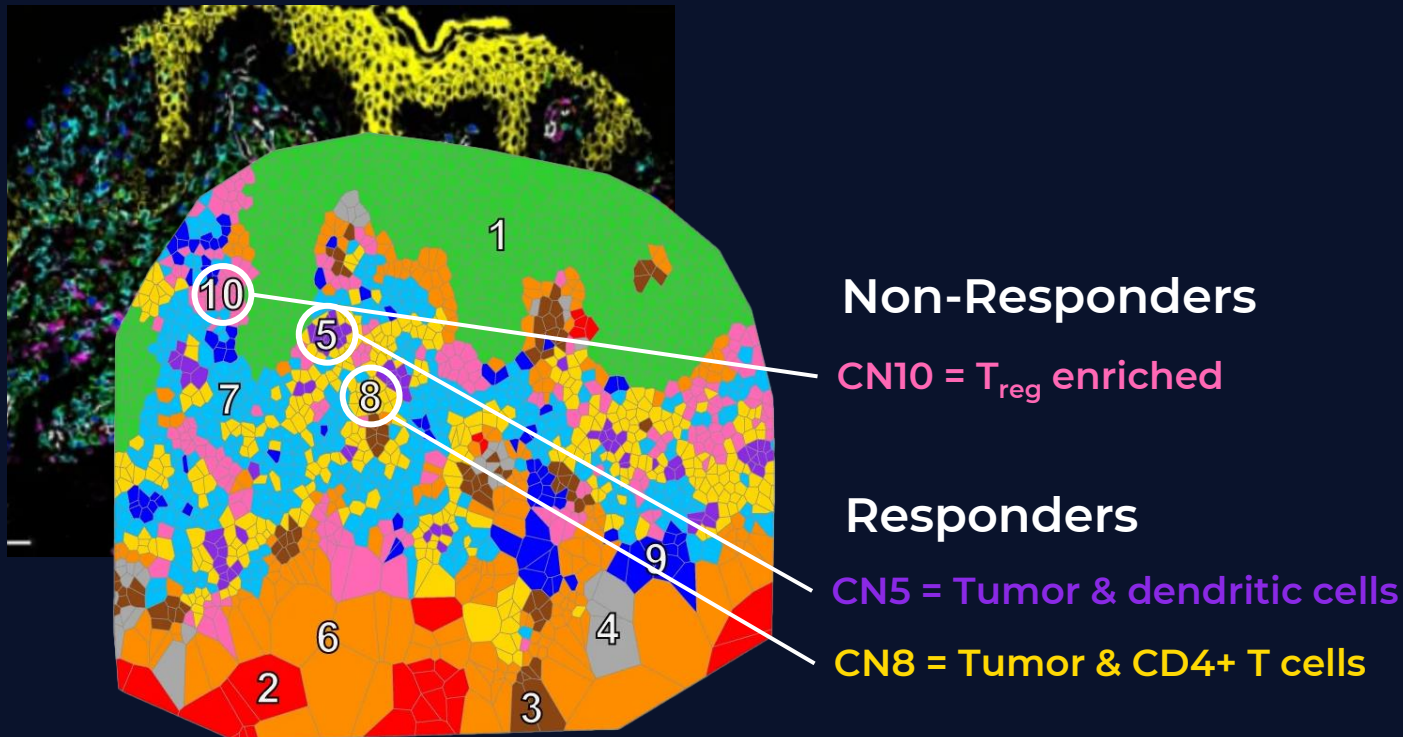


First published 100+ protein plex whole-slide image, comprehensively mapping the spatial proteome of head and neck squamous cell carcinoma, on PhenoCyler-Fusion¹

*As of September 30, 2024
¹Published October 17, 2023

Spatial Biology Markers Predicting Response to Therapy

Predicts Response to PD-1 Blockade in Cutaneous T Cell Lymphoma (CTCL)



Cellular Neighborhood (CN)
Analysis Identified Localized
Enrichment

- *SpatialScore* derived from spatial relationship b/w PD-1+CD4+ T cells, tumor cells and immunosuppressive Tregs.
- *SpatialScore* demonstrates high correlation with response to pembrolizumab in CTCL
- PhenoCycler-Fusion high-plex data used to develop a targeted panel for larger cohort studies on the Phenolmager HT

ACR-368 OncoSignature Assay – a New Era of Precision Medicine

First-of-its-kind Spatial Signature CDx Assay to Identify Patients for a Targeted Oncology Agent



OncoSignature®



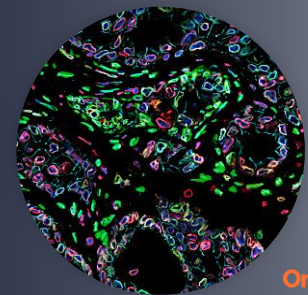
Acrivon and Akoya partnering on ongoing clinical development and future commercial use of the ACR-368 OncoSignature Assay

Acrivon granted Breakthrough Device Designation: ACR-368 OncoSignature Assay + PhenolMager HT + Akoya Software for the identification of ovarian cancer patients who may benefit from ACR-368

Acrivon granted Fast Track Designation: Investigation of ACR-368 as monotherapy based on ACR-368 OncoSignature-predicted sensitivity in patients with platinum-resistant ovarian and endometrial cancer



Multiplex immunofluorescence CDx assay developed on PhenolMager HT



Acrivon's ACR-368 OncoSignature test, a drug-tailored spatial signature assay



Patient screened using ACR-368 OncoSignature test to determine clinical treatment in Acrivon's registrational intent Phase 2 trial of ACR-368



Pending FDA approval, results of ACR-368 OncoSignature test used to assign therapy

Akoya and Acrivon will co-develop, validate and **EXCLUSIVELY** commercialize the ACR-368 OncoSignature test



Exclusive Partnership with NeraCare in Early-stage Melanoma

Immunoprint® Assay and Phenolmager HT Platform to Enable Personalized Therapy Selection



Stage	5-Y survival	Patients		Adjuvant therapy
	RFS	# ⁽⁴⁾	% of total ⁽⁵⁾	
IA	95% ⁽¹⁾	133,573	57%	X
IB	88% ⁽¹⁾	30,740	13%	X
IIA	73% ⁽¹⁾	18,817	8%	X
IIB	62% ⁽¹⁾	11,559	5%	✓
IIC	44% ⁽¹⁾	4,903	2%	✓
III	25% ^(2,3)	23,596	10%	✓
IV	n/a	12,057	5%	✓
Total		235,479	100%	

Identified using Immunoprint®

Over 235,000 new cases of melanoma diagnosed globally every year

- Currently, adjuvant therapy is only approved for stages IIB-IV resectable cutaneous melanoma (c.15-20% of patients), however, earlier-stage patients contribute a significant share of overall melanoma mortality but cannot be identified with AJCC staging
- Immunoprint® is a **multiplex assay** which has demonstrated robust clinical performance in identifying **early-stage melanoma patients at high risk of relapse and death** that could potentially benefit from on-market therapies
- Akoya and NeraCare will develop market opportunities to combine Akoya's Phenolmager HT platform and NeraCare's Immunoprint assay for therapy selection in early-stage melanoma patients

Note: Survival data in stage III refers to an untreated population / placebo arms of phase III adjuvant therapy trials

(1) Garbe et al.: Prognosis of Pts With Melanoma Stage I and II According to AJCC v8 [...]: Implications for Adjuvant Treatment, J Clin Oncol (2022), [LINK](#)

(2) Weber et al: Nivolumab versus placebo as adjuvant therapy for resected stage III melanoma, Cancer Immunology, Immunotherapy (2023), [LINK](#)

(3) Garbe, [...] Eggermont: Prognosis of Patients With Stage III Melanoma According to AJCC v8, J Clin Oncol (2020), [LINK](#)

(4) Annually diagnosed in Europe (106,310), USA & Canada (106,369), Australia & New Zealand (22,800)

(5) Patient distribution (i) according to SEER database [LINK](#) (Adjusted for unstaged pts: 5% stage IV, 11% stage III; 85% stage I&II) and (ii) further stage I&II substage distribution according to Garbe et al., J Clin Oncol (2022)

Akoya and KR Pharmtech Announce NMPA Approval for KR-HT5

Based on Phenolmager HT to Drive Next Generation Pathology Clinical Solutions in China

- KR-HT5, co-developed with Shanghai KR Pharmtech utilizing Phenolmager HT technology as its foundation, has secured **premarket approval** from China's National Medical Products Administration (NMPA)
- NMPA clearance allows **clinicians** to use the instrument



KR-HT5 high-throughput
mIF scanning system

Akoya's Workflow – Owning the Biomarker Journey

Consistency and Flexibility Drive Platform Utilization and Pull Through Across a Continuum

PROBE & STAIN



Rapid Menu Expansion

- Consistent chemistries
- Ready-to-use panels and off-the-shelf antibody compatibility
- Flexible plex options

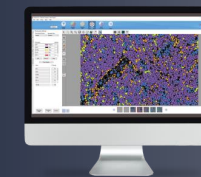
IMAGE



Accelerating Workflows

- Whole-slide imaging area
- Market-leading throughput
- Workflow simplification

ANALYZE



Flexible Data Analysis

- Consistent image analysis methods
- Proprietary data compression
- Solutions serve every user need

PhenoCycler-Fusion 2.0 Platform

More Discoveries, Faster Than Ever — High-plex Panels for Comprehensive Coverage



Fastest Imaging Technology

Scalable Chemistry

Standardized Gigabyte-sized **QTIFF** Files

PhenoCode
Discovery Panels

TISSUE ARCHITECTURE MODULE

IMMUNE PROFILING CORE

LYMPHOCYTE PROFILING MODULE

IMMUNE ACTIVATION & PROLIFERATION MODULE



350+ Antibodies



100+ Phenotypes



30+ Tissue Types



Multiple Species

PhenoImager HT 2.0 Platform

The Fastest End-to-end Solution for Immuno-Oncology Spatial Signature Development

Immuno-Contexture
CD8
CD68
PD-L1
FoxP3
PanCK

Immune Profile
CD8
CD68
CD3
CD20
PanCK

Activated TIL Status
CD8
CD3
Ki67
GrzB
PanCK

M1/M2 Polarization
CD8
CD68
CD163
PD-1
PD-L1

T Cell Status
CD8
CD4
FoxP3
CD20
PD-1

PhenoCode Signature Panels

Fastest Multispectral Imaging Technology

Onboard Spectral Unmixing

Standardized Gigabyte Sized 16-bit **QPTIFF files**

Labels in diagram: CAMERA, FILTER CUBE, LIGHT SOURCE FOR HEIGHT SENSING, OBJECTIVE, SAMPLE, Wavelength (nm)

Data Analysis Ecosystem Across Akoya's Workflows

Powerful Ultrahigh-Plex
Analysis in the Cloud

ENABLE MEDICINE

Flexible Open Source



Machine Learning and AI

VISIOPHARM®

indica labs HALO
QUANTITATIVE PATHOLOGY

PathAI

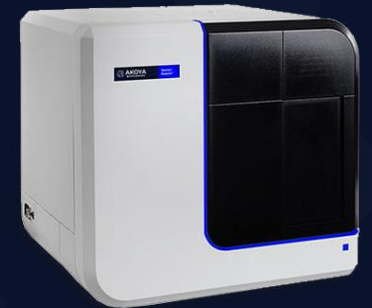
Leading Analysis
Service Providers

OracleBio



Large installed based and QTIFF enables a growing ecosystem

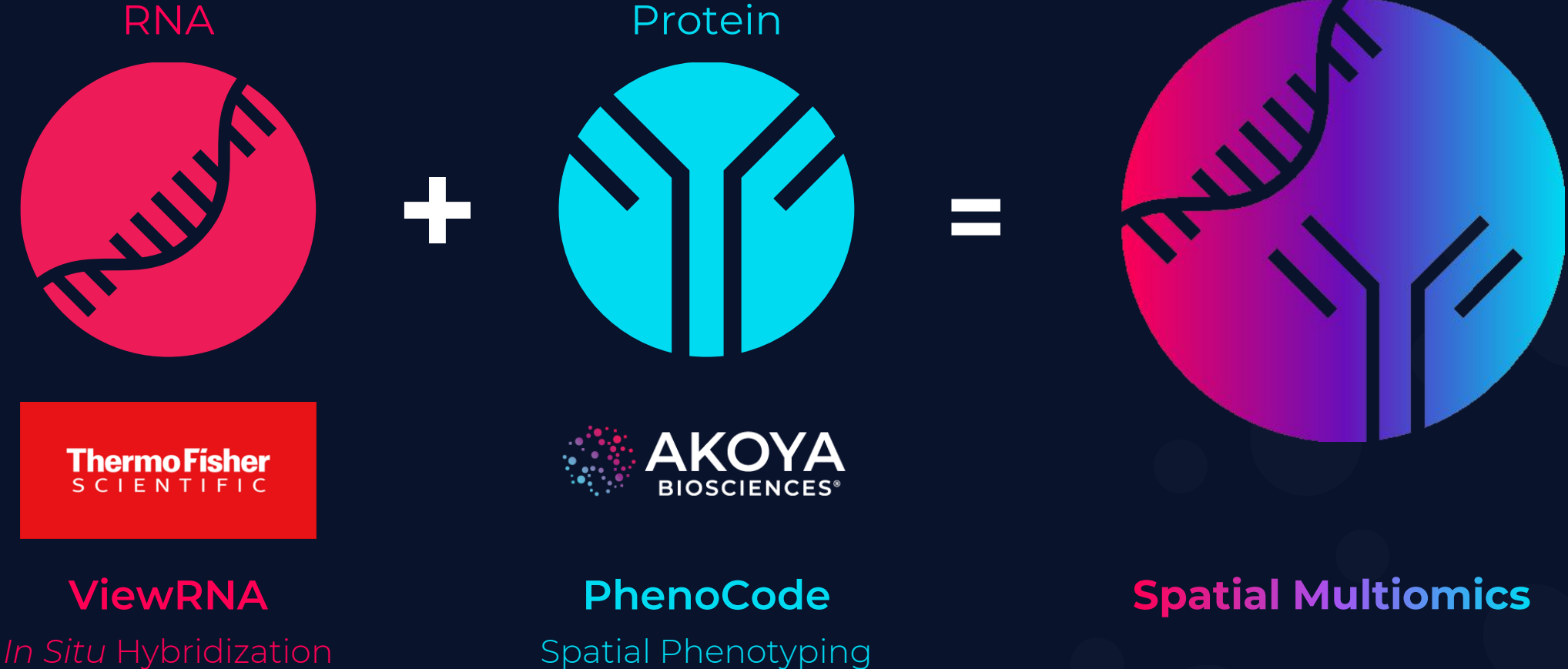
Accessibility to cutting-edge analysis



Software partnerships offer **powerful data analysis solutions** to meet the **varying requirements** of our customers

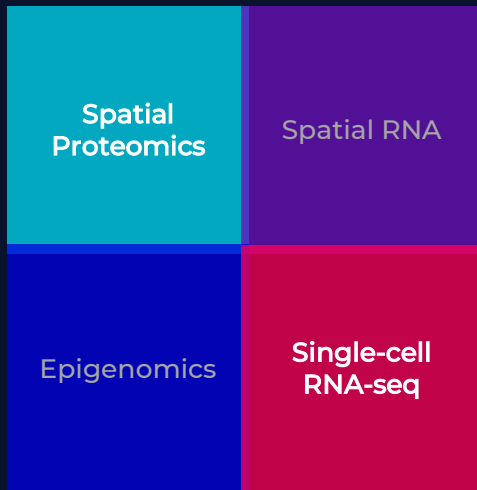
Akoya and Thermo Fisher to Deliver Leading Spatial Multiomics

Streamlined Workflow for Rapid and Whole-slide Imaging of RNA and Protein Biomarkers



MaxFuse – Multiomic Integration of Spatial and Single-cell Data

AI Driven Digital Integration of Proteomic, Transcriptomic and Epigenomic Data on Same Tissue Types



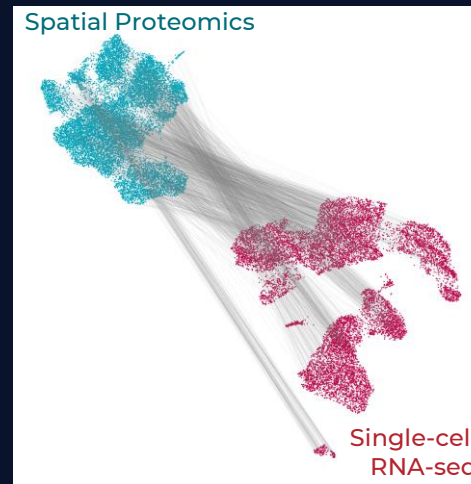
MaxFuse



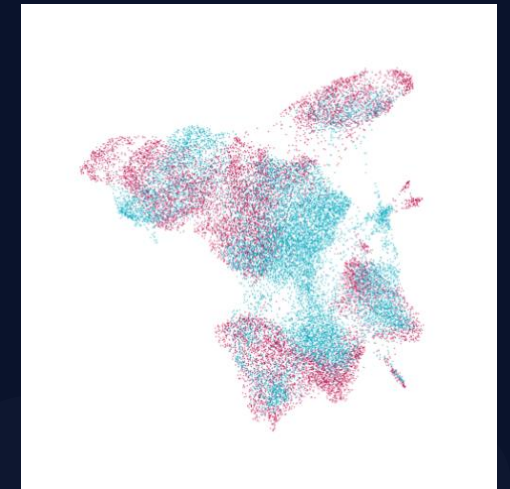
Iterative co-embedding

Data smoothing

Cell matching



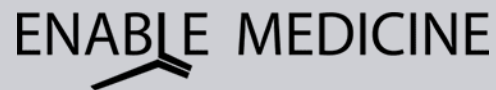
MaxFuse algorithm uses spatial proteomic data to **infer spatial information** on single-cell RNA-seq data sets



Using spatial proteomic data to **maximize the value of new and historical** scRNA and epigenomic data

Rapidly Expanding Qualified CRO Service Provider Network

- Partnership with best-in-class CROs **amplify** the use of Akoya's platforms
- Qualification process ensures **consistent** and **best practices** across the network.



Akoya's New Scientific Advisory Board

Leading Experts in Innovation, Immunobiology and Immunotherapy



James Allison, Ph.D.

Chair of the Department of Immunology,
MD Anderson Cancer Center

2018 Nobel Prize Winner in Physiology or Medicine



Garry Nolan, Ph.D. (Chair)

Professor in the Department of Pathology,
Stanford University School of Medicine



Padmanee Sharma, M.D., Ph.D.

Professor in the Departments of Genitourinary
Medical Oncology and Immunology,
MD Anderson Cancer Center

Akoya's 2024 Strategic Priorities

Driving Operational Leverage and Gross Margin Improvements to Meet our Profitability Goals



Accelerate Pull Through

- Expand menu of applications
- Continuous platform improvements drive throughput
- Streamline data analysis and time to answer



Build Clinical IVD Menu

- Expand clinical trial participation – leveraging our CLIA services lab and CRO partner network
- Rapidly grow CDx pipeline
- Advance clinical workflow and regulatory capabilities / readiness



Financial Overview



Recurring revenue model

- Recurring reagent revenue from global installed base driving projected gross margin increase



Favorable growth profile

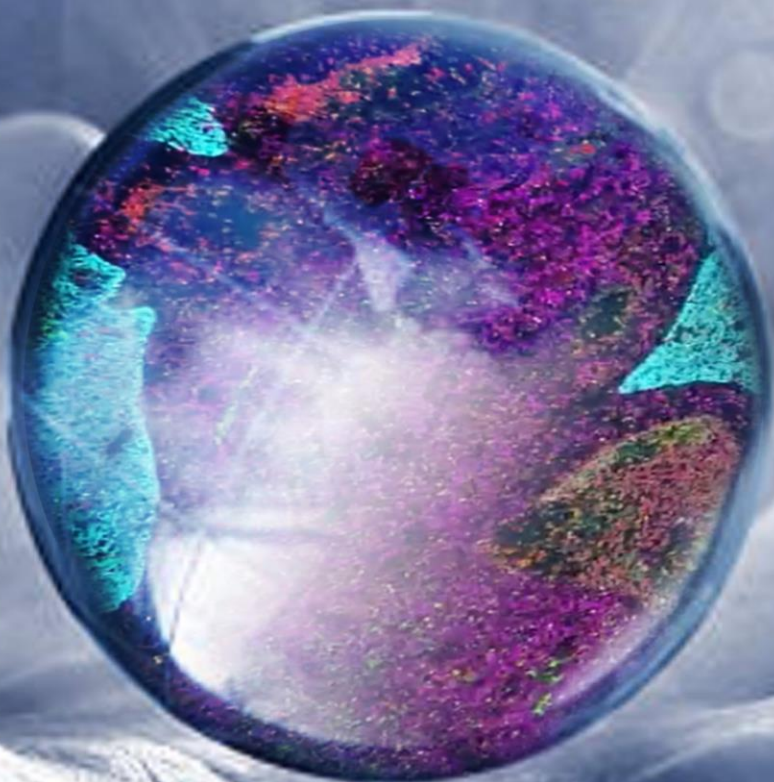
- Expanding installed base, menu, pull through and clinical lab services



Path to profitability

- \$39.3 million of cash, cash equivalents and marketable securities as of September 30, 2024

2024 Revenue Guide:
\$80 – 85 Million



Catalyzing **Discovery** and Improving **Patient Care**