

Investor Overview

February 2022

Legal Information

Forward-looking statements

This presentation and the accompanying presentation contain forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995, including, among others, statements regarding Fluidigm's vision and strategy, target markets and revenues, anticipated collaborations, growth plans, recurring revenue streams, expense management, product applications and adoption in certain markets, addressable market growth projections, adoption of, benefits of and demand for new and recently introduced products, OEM and collaboration opportunities, product innovation pipeline, and market segment growth and penetration. Forward-looking statements are subject to numerous risks and uncertainties that could cause actual results to differ materially from currently anticipated results, including but not limited to risks relating to the potential adverse effects of the coronavirus pandemic on our business and operating results; any failure to obtain required stockholder approval of a pending financing transaction (the "Transaction"); the possibility that the conditions to the closing of the Transaction are not satisfied; potential litigation relating to the Transaction; uncertainties as to the timing of the consummation of the Transaction; the ability of each party to consummate the Transaction; possible disruption related to the Transaction to Fluidigm's current plans and operations, including through the loss of customers, suppliers and employees; changes in Fluidigm's business or external market conditions; customers and prospective customers continuing to curtail or suspend activities utilizing our products; our ability and/or the ability of the research institutions utilizing our products and technology to obtain and maintain Emergency Use Authorization from the FDA and any other requisite authorizations or approvals to use our products and technology for diagnostic testing purposes; 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, Fluidigm 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; uncertainties in contractual relationships; reductions in research and development spending or changes in budget priorities by customers; Fluidigm research and development and distribution plans and capabilities; interruptions or delays in the supply of components or materials for, or manufacturing of, Fluidigm products; potential product performance and quality issues; risks associated with international operations; intellectual property risks; and competition. Information on these and additional risks and uncertainties and other information affecting Fluidigm's business and operating results is contained in its Annual Report on Form 10-K for the year ended December 31, 2020, and in its other filings with the Securities and Exchange Commission. These forwardlooking statements speak only as of the date hereof. Fluidigm disclaims any obligation to update these forward-looking statements except as may be required by law.

Non-GAAP financial information

This presentation and the accompanying presentation have certain financial information in accordance with U.S. GAAP and also on a non-GAAP basis for the three-month and twelve-month periods ended December 31, 2021. 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. The time and amount of certain material items needed to estimate non-GAAP financial measures are inherently unpredictable or outside of our control. Material changes to any of these items could have a significant effect on guidance and future GAAP results. 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. Fluidigm encourages investors to carefully consider its results under GAAP, as well as its supplemental non-GAAP information and the reconciliation between these presentations, to more fully understand its business. Reconciliations between GAAP and Non-GAAP operating results are presented in the tables of this presentation or in the accompanying "Reconciliations and Financial Package" available at supplemental financials.

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Drive Meaningful Insight in Health and Disease to Improve Life



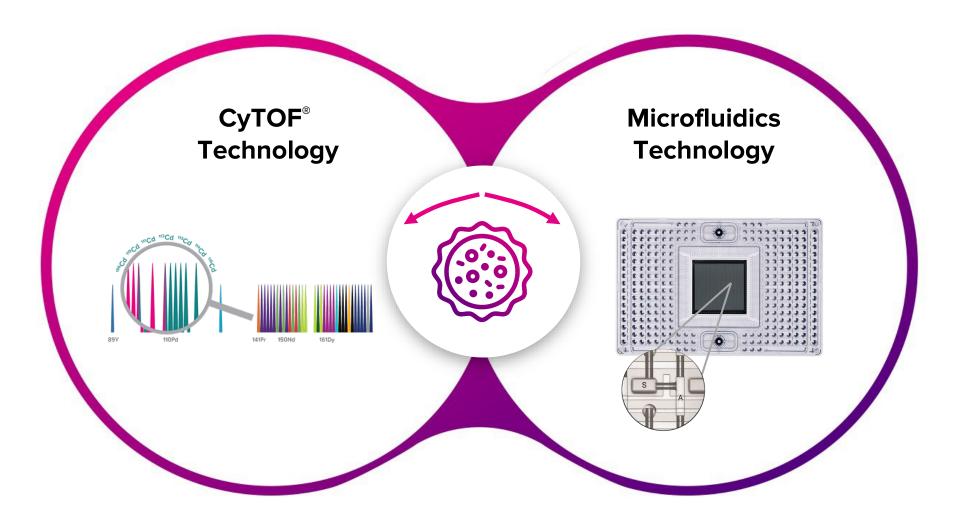


Advance human health by deploying innovative technologies.



Reveal, understand and address the biological complexities of disease.

Harnessing the Power of Two Technologies





Powerful Growth Drivers

Vision 2025

We will span the spectrum from discovery to diagnostics, delivering double-digit revenue growth with sustained profitability.





Innovation

Launching instruments, expanding menu and creating new content and workflows



Partnerships

Building new capabilities, broadening our customer base and penetrating applied markets



Beachhead Expansion

Building a transformative diagnostics base and moving closer to health care decision making



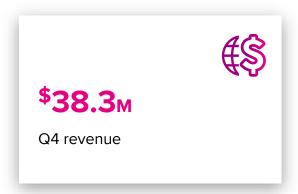
Key Investment Highlights

- 1. Addressing large market opportunities
- 2. Offering proprietary platform technologies with demonstrated clinical research and real-world utility
- **3.** Driving recurring revenue streams
- 4. Targeting long-term double-digit revenue growth and sustained profitability

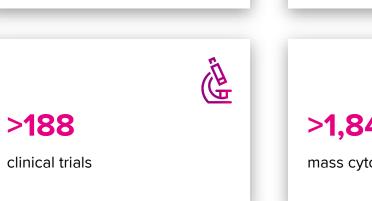


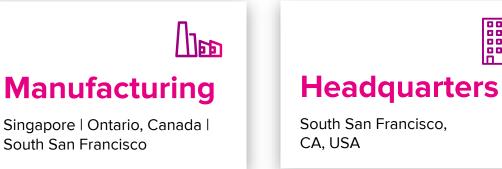


Leading Provider of Indispensable Tools and Consumables











575 issued or pending patents (worldwide)

For the quarter ended December 31, 2021. For reconciliations of the Non-GAAP financial measures to the GAAP measures, please refer to: supplemental financials.



Operational Efficiencies Driving Productivity

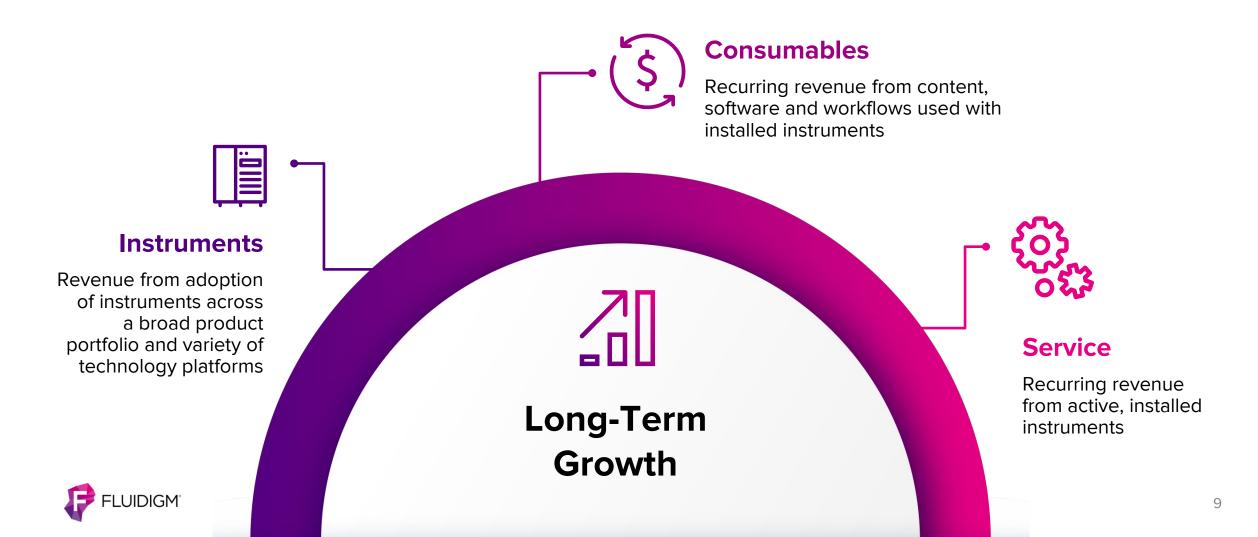
Disciplined operating expense management

Instrument placements, recurring revenue, partnerships

Manufacturing productivity

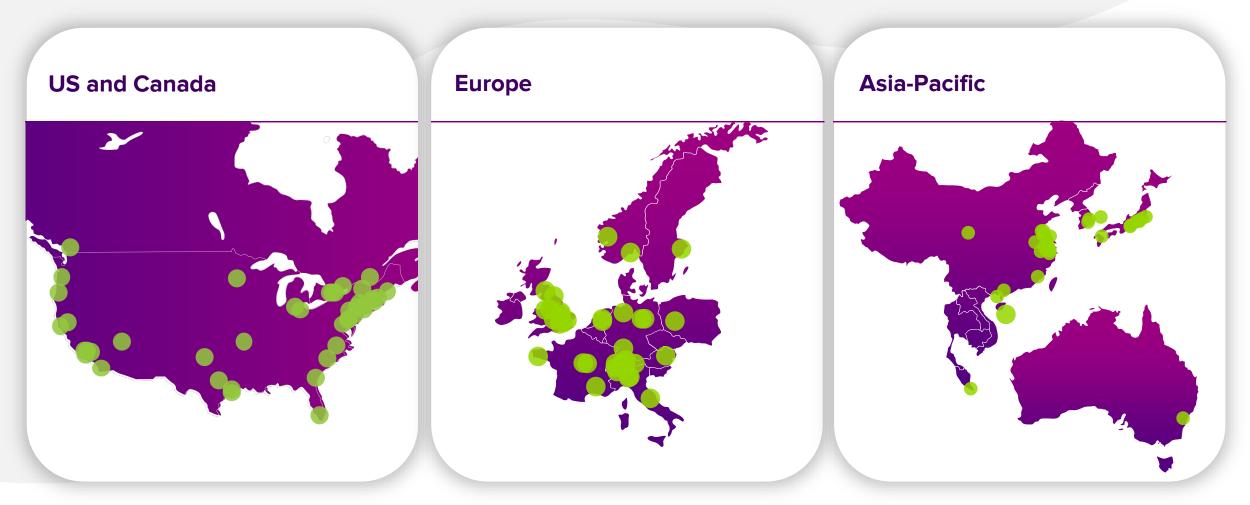


Long-Term Recurring Revenue Growth



Global Presence

Presence in 9 of Top 10 Pharma (WW) and 61% of Comprehensive Cancer Centers (US)





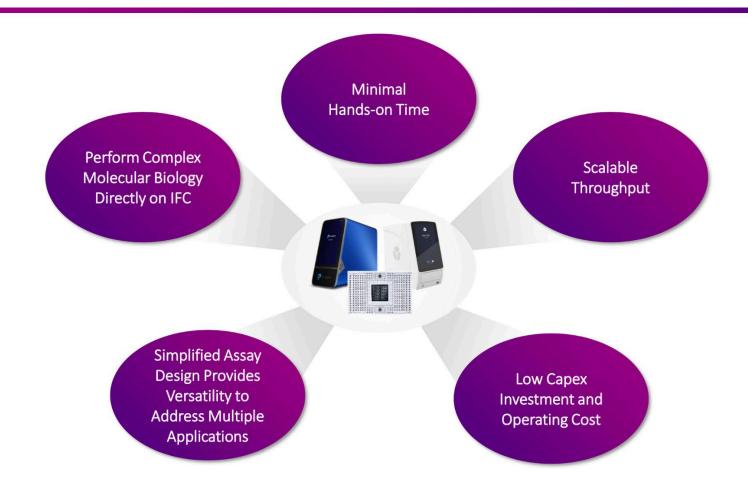
Microfluidics

Attractive markets that extend beyond COVID-19



Fluidigm Microfluidics Platform

Offers ease of workflow, cost savings and turnaround time without sacrificing performance





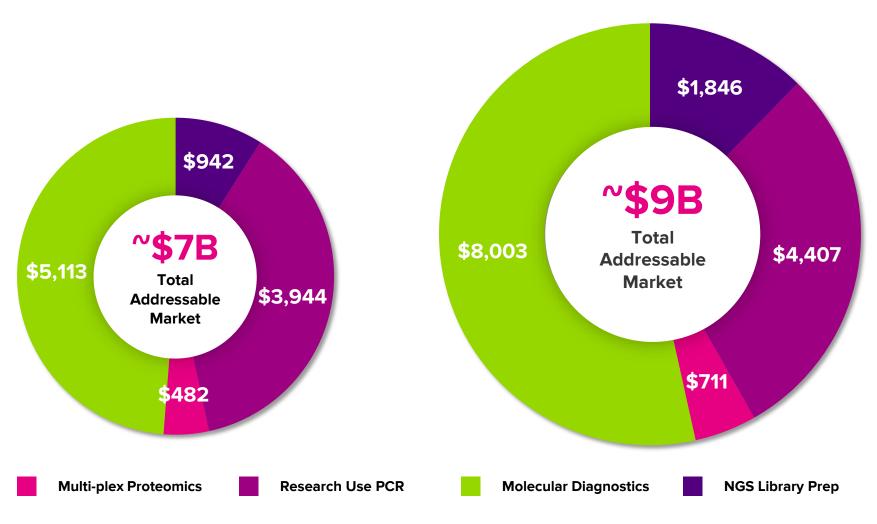
Used Non-Dilutive Funding to Upgrade Microfluidics Platform and Manufacturing Capacity

- Investment in new Biomark platform
- Expanded manufacturing capabilities
- Development of new Sample-to-Answer IFC to significantly expand installed base into midthroughput labs that value assay flexibility, scalable throughput, cost and data quality as their main driver to scale up their testing menu
- Experience bringing new tools through regulatory review and approval, building upon market experience
- Allows for Fluidigm's Microfluidics business to have an advantaged role in serving currently 200 specialty labs running high complexity molecular LDTs with an addressable market of \$2B





Large Market Opportunities



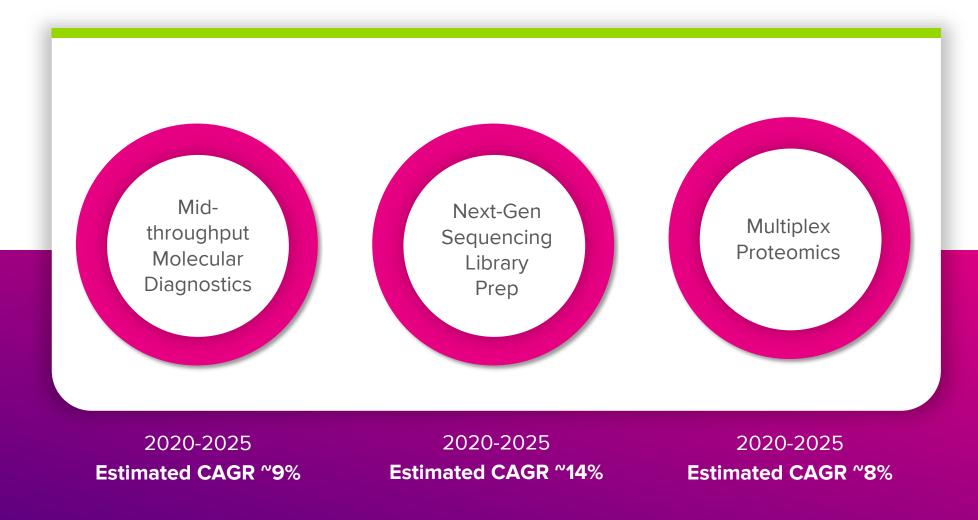


5%

Growth from 2020 to 2025



With Three Focus Areas





Platform Represents a Scalable Solution for Other Attractive Markets

	Consumer Genomics Providers	Clinical Research Laboratories		
Target Customers	Telemedicine and Walk in ClinicsPersonal Genomics	Clinical Laboratories developing novel LDTsClinical labs supporting Digital health		
Key Needs	 Ability to work with low or high sample volumes Customizable menu Flexible solutions with high test capacity 	Scalable solutionsFlexible solutionsIntegrated workflows		

Fluidigm's Solution

Fluidigm's Next Generation Biomark™ X and the Sample-to-Answer Chip

Highly Scalable Solutions

Flexible Assay Design

Lowest Cost

Works with Low Sample Volumes of Blood, Nasal Swabs and Saliva



Microfluidics OEM Opportunities

A key exemplar of future partnerships to grow revenue



Provided development revenue for new instrument (derivative of future next-gen Biomark) to be sold by Olink



Major milestone in Q2 2021 when Olink launched the instrument branded as the Signature Q100, a designated benchtop system for protein biomarker analysis



Signature Q100 will utilize proprietary chips from Fluidigm providing attractive recurring revenue stream and supporting margins



Partnerships like Olink will help us penetrate new markets and advance the field of proteomics and serve as a first-mover exemplar of our OEM strategy to propel Microfluidics growth.



OEM revenue is anticipated to expand rather significantly through 2025 as a result of current and contemplated partnerships with key industry players



Microfluidics Innovation

Innovative solutions to expand market opportunities

Integrated System

Platform will Open Up New Markets

- Molecular Diagnostics
- Expand Addressable Market for PCR
- Clinical Labs developing LDTs
- Personal Genomics

Sample-to-Answer IFC and Biomark[™] X



Key Highlights

- Integrated Fluidic Circuit (IFC) loading and qPCR function (no more Juno™ or Controller)
- 6x less volume (dimension) with same robustness of Biomark™ HD
- User installable
- Compatible with new IFC enabling Sample-to-Answer workflow

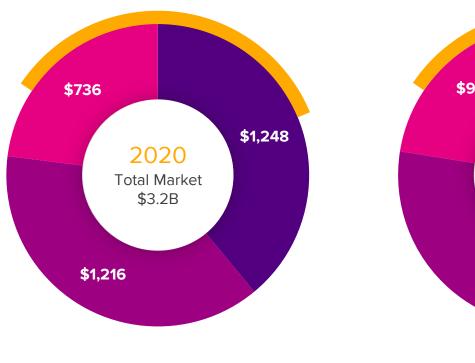


Mass Cytometry

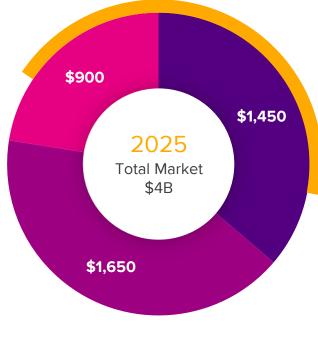
The world's most advanced single-cell proteomics technology



Focused on Highest Growing Cytometry Market Segments



- Academic and Research Institutions
- Pharma, CRO, Biotechnology



- Hospital and Clinical Testing
- Clinical and Translational Research

- Clinical and Translational Research Market \$700M-\$1,200M (2020) growing to \$1,300M-\$2B in 2025
- Clinical and Translational Research Market is growing at 10%-plus



Opportunity to Expand Market Penetration





Introducing CyTOF XT

Mass Cytometry Product Enablement Roadmap



CyT#F*XT

- Reduced total cost of ownership
- Automated setup and data acquisition
- Extended run times and system monitoring
- 22 systems placed since launch

Enables

- Site standardization
- Increased productivity
- Studies with larger sample sizes



CyTOF XT: Affordable High-Parameter Cytometry

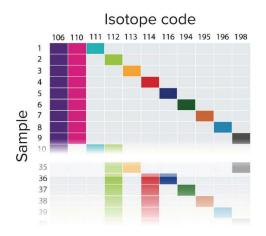
Anticipated ASP: \$365K to \$410K USD. Positioned to drive unit placements. High-margins Service offering in line with market expectations.



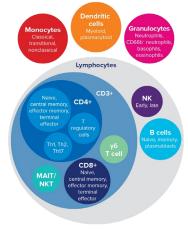


Consumables

Setting the Standard in Clinical Research







Live-cell barcoding

- Sample multiplexing for increased efficiency
- Enhanced data quality and workflow

Expansion modules for Maxpar[®] Direct[™] Immune Profiling Assay[™]

• Deep profiling of >35 immune cell populations with enhanced phenotyping of activation states, cytokine production

Enables

- Larger studies
- Access to more applied markets (infectious disease)
- Standardization across sites



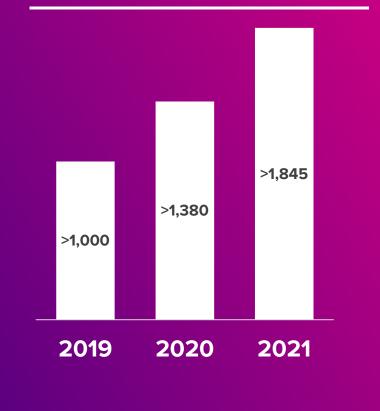
Accelerated Pace of Adoption

National Clinical Trials Citing CyTOF Technology*

By Study Start Date



Cumulative Publications*





Innovation Accelerates Segment Growth

Higher-Throughput Platforms

H₂ 2021

CyTOF XT™

2022

- Clinical cytometry entry in China via PLT partnership
- 2023 2025
 - Planned platform upgrades

Fixed and Flexible Assays

H1 2021

- 687 conjugates
- 28 panels
- 53 parameters

H2 2021

- ~750 conjugates
- 31 panels
- 57 parameters

2022

- 1,000–1,400 conjugates
- >35 panels
- 60-plus parameters

2023 - 2025

- >2,000 conjugates
- >50 panels
- 70-plus parameters

Automated Analysis

H2 2021

- · Instrument remote monitoring
- Maxpar® Pathsetter™ customization (automated analysis for immune monitoring)

2022

• CyTOF XT user interface upgrade

2023 - 2025

- Disease research specific modules
- Blood cancer diagnostic and immunotherapy guidance (PLT)
- Cloud analysis



Tissue Imaging

The most proven approach to high-multiplex imaging and single-cell protein analysis



Meeting the Needs of Target Markets

Translational and Clinical Research

Translational

- **Segment/Customer Need:** High-multiplexing, working with limited blood/tissue samples and inclusion of spatial information
- **Fluidigm Solution:** Mass Cytometry and Tissue Imaging for Fluidigm's customers has shown it provides the highest plexity for protein targets and is identifying new biomarkers associated with alternate disease prognoses and therapy guidance.

Clinical

- Segment/Customer Need: Automation, consistency and standardization, fixed and validated panels, unbiased analysis
- Fluidigm Solution: Foundational technology provides consistent and stable measurement/readout. Mass Cytometry for Fluidigm customers has shown it provides an ability to test new biomarkers associated with disease prognoses and therapy guidance.

Research Genre Description

Discovery Research

Systematic study directed toward greater understanding of fundamental mechanisms that drive disease

Translational Research

Transfers new understandings into the development of new methods for diagnosis, therapy and prevention in humans

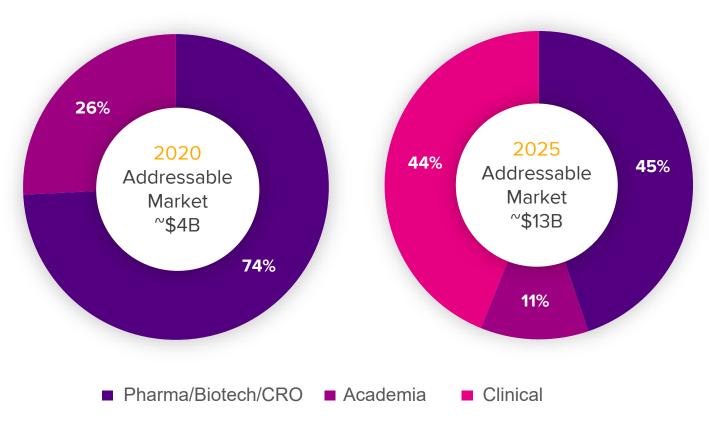
Clinical Research

The study of human subjects and samples, testing new methods of diagnosis, prevention and treatment



Fluidigm Strength in Translational Markets

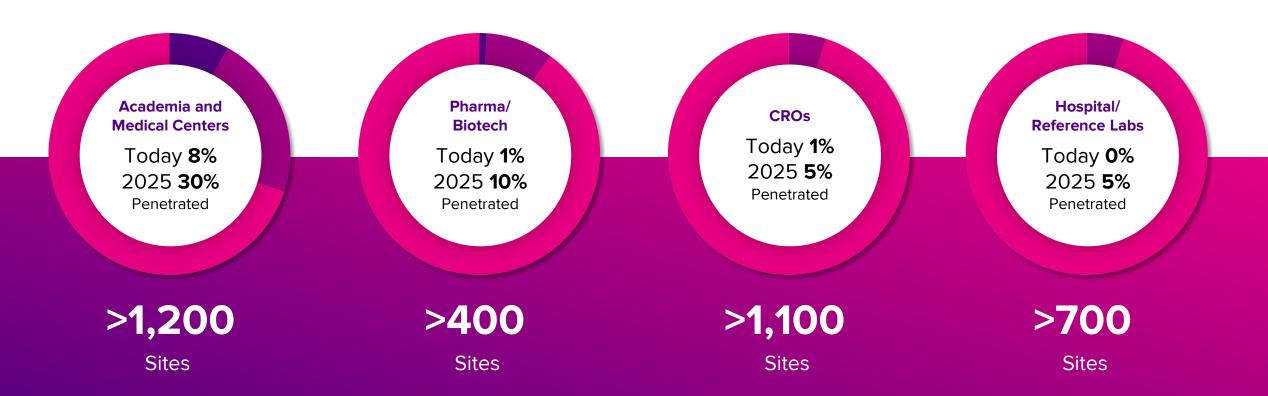
Spatial proteomics is largely translational today, but the potential for spatial in the clinical setting is growing rapidly.



- Total Tissue Imaging market is growing at ~12% CAGR.
- Translational segment is driving demand for high-plex platforms.
- Increased future addressable market:
 - Improved workflow by aggregating current immunohistochemistry biomarkers into one test
 - Improved predictive value compared to existing prognostic therapy guidance test with potential novel content unique to spatial



Opportunity to Expand Market Penetration





Fluidigm Offers the Most Complete Solution

Translational and Clinical Research

	Marker Type	Multiplexity	Resolution	Cost per Sample	Sensitivity	Verified Reagents
Spatial Proteomics (Fluidigm)						
Cyclic Immunofluorescence				0		
Spatial Transcriptomics						



Highlights

Fluidigm continues to be at the cutting edge of innovation.



Vision 2025: Innovation

To Penetrate Future Clinical Settings

Platforms

Q4 2021

• New Tissue Imager early access

H₁ 2022

 Commercial release of new Tissue Imager

2022-2025

Future platform development focused on:

- Increased speed, sensitivity, throughput and robustness
- Simplified user experience
- Automation

Fixed and Flexible Assays

H2 2021

- 150–200 conjugates
- 6 panels
- 39 channels

2022

- 400-600 conjugates
- 10-plus panels
- 40 channels

2022-2025

- >1,000 conjugates
- >20 panels
- 50-plus channels

Automated Analysis

H2 2021

Semi-automated analysis

2022

- · Application-specific output
- Al cell segmentation

2023-2025

- · Disease-specific modules
- Cloud-based personalized applications



