



Morgan Stanley Healthcare Conference

September 10, 2019

Forward-Looking Statements

This presentation contains statements that are not historical and that are based on our beliefs and assumptions and on information currently available to us. These statements constitute forward-looking statements within the meaning of the Safe Harbor provisions of the Private Securities Litigation Reform Act of 1995. Forward-looking statements are subject to known and unknown risks, uncertainties, assumptions, and other factors that could cause actual results to differ materially from our expectations.

Forward-looking statements can be identified by words such as: "anticipate," "intend," "plan," "expect," "believe," "should," "may," "will" and similar references to future periods. Examples of forward-looking statements include, among others, statements we make regarding our ability to achieve milestones under the collaboration agreement with Johnson & Johnson; our ability to achieve and maintain Medicare coverage for our tests; the benefits of our tests and the applicability of clinical results to actual outcomes; the size of our addressable market; the laws and regulations applicable to our business, including potential regulation by the Food and Drug Administration or other regulatory bodies; our ability to successfully achieve and maintain adoption of and reimbursement for our products; the amount by which use of our products are able to reduce invasive procedures and misdiagnosis, and reduce healthcare costs; the occurrence and outcomes of clinical studies; the timing and publication of study results; the applicability of clinical results to actual outcomes; the continued application of clinical guidelines to our products and their inclusion in such clinical practice guidelines; our ability to compete; our ability to obtain capital when needed; and other risks set forth in our filings with the Securities and Exchange Commission, including the risks set forth in our Quarterly Report on Form 10-Q for the quarter ended June 30, 2019. These forward-looking statements speak only as of the date hereof and Veracyte specifically disclaims any obligation to update these forward-looking statements.

Veracyte, Afirma, Percepta, Envisia, Know by Design, the Veracyte logo and Afirma logo are trademarks of Veracyte, Inc. This presentation also contains trademarks and trade names that are the property of their respective owners.



A Genomics Leader Creating Value Through Innovation

FOUNDED WITH A MISSION TO IMPROVE DIAGNOSTIC ACCURACY **Expanding to advance early detection and inform treatment decisions**

Comprehensive scientific approach using whole-transcriptome sequencing coupled with machine learning to develop diagnostic tests that we believe can change clinical care

Clinical evidence published in top-tier journals to facilitate test adoption, coverage and reimbursement

Market leader with first-to-market tests in large, untapped clinical areas: thyroid cancer, lung cancer and idiopathic pulmonary fibrosis (IPF)

Robust pipeline, including non-invasive test for lung cancer, and biopharma partnerships to augment future growth

Experienced management team with deep expertise and proven track record









Execution Driving Momentum

Strong Commercial Growth

Driven by multi-product sales strategy

Revenue growth*

32%

Genomic test volume*

26%

- Afirma revenue grew by 16%*
- Percepta classifier volume increased 142%*
- **Envisia** volume increased to 130 tests from 5 in prior year

*2Q19 compared with 2Q18

Continued Reimbursement Expansion

All classifiers covered by Medicare

(Afirma, Percepta and Envisia)

Achieved in-network status with nearly all major U.S. health plans as a service provider

Strategic Collaborations

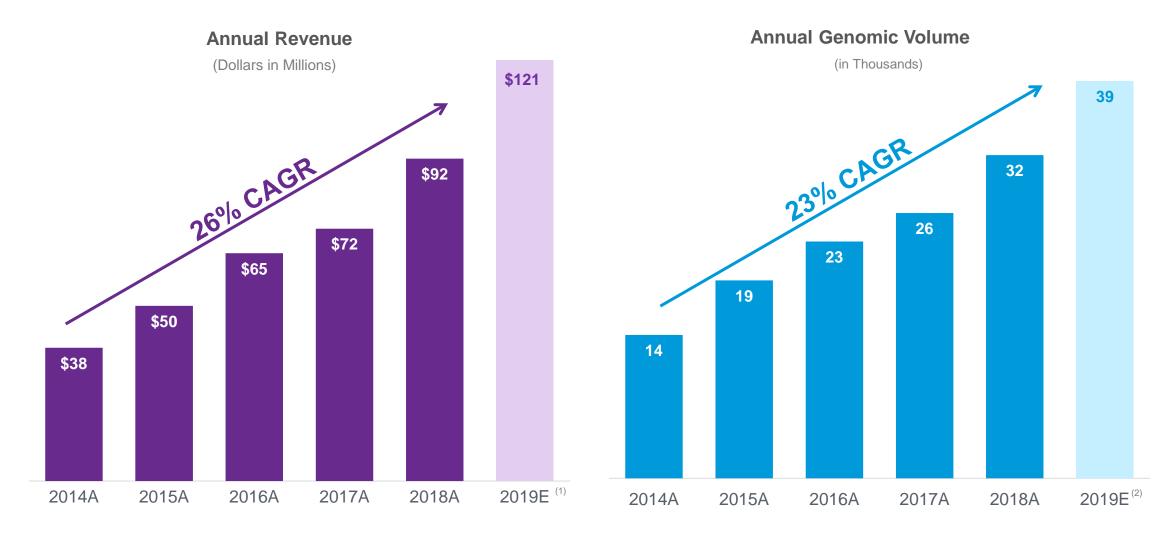
Advancing Pipeline and Driving Value Creation

Strategic collaboration with Johnson & Johnson

- Launched Percepta[®] Genomic Sequencing Classifier
- Accelerating development of first nasal swab test for early lung cancer detection
- Global addressable market for lung cancer estimated at more than \$30 billion



Robust Annual Revenue and Genomic Volume Growth



⁽¹⁾ Midpoint of 2019 revenue guidance as of July 30, 2019

(2) As of July 30, 2019, the \$121 million midpoint of 2019 revenue guidance is supported by an estimated 20% to 25% growth in genomic test volume over 2018, or a midpoint of approximately 39,000 reported tests in 2019



Building on a Firm Foundation

Following a Proven Formula for Success



Relevance

ANSWER QUESTIONS THAT MATTER!

INTEGRATE INTO CURRENT CARE PATHWAY TO CHANGE PRACTICE AND REDUCE COSTS



Rigor

BUILD ROBUST SCIENTIFIC AND CLINICAL EVIDENCE; INFORM GUIDELINES



Rationale

PROVIDE ANSWERS THAT CHANGE CARE WITH REAL CLINICAL UTILITY AND ECONOMIC VALUE

Reimbursement

Extensive experience and coverage policies with payors pave way for reimbursement expansion



A Powerful Scientific Platform: Multiple Vectors for Value Creation

DISCOVERY

Unique Biorepositories

DEVELOPMENT & VALIDATION

Proprietary Technology & Bioinformatics

COMMERCIAL

Informing Clinical Questions

BIOPHARMA

Monetizing Assets

Extensive Clinical Cohorts

RNA Whole-Transcriptome Sequencing "Unified Assay" **Deep Machine Learning**

Genomic Variants "Xpression Atlas"

AFIRMA, PERCEPTA, ENVISIA "DIAGNOSTIC CLASSIFIERS"

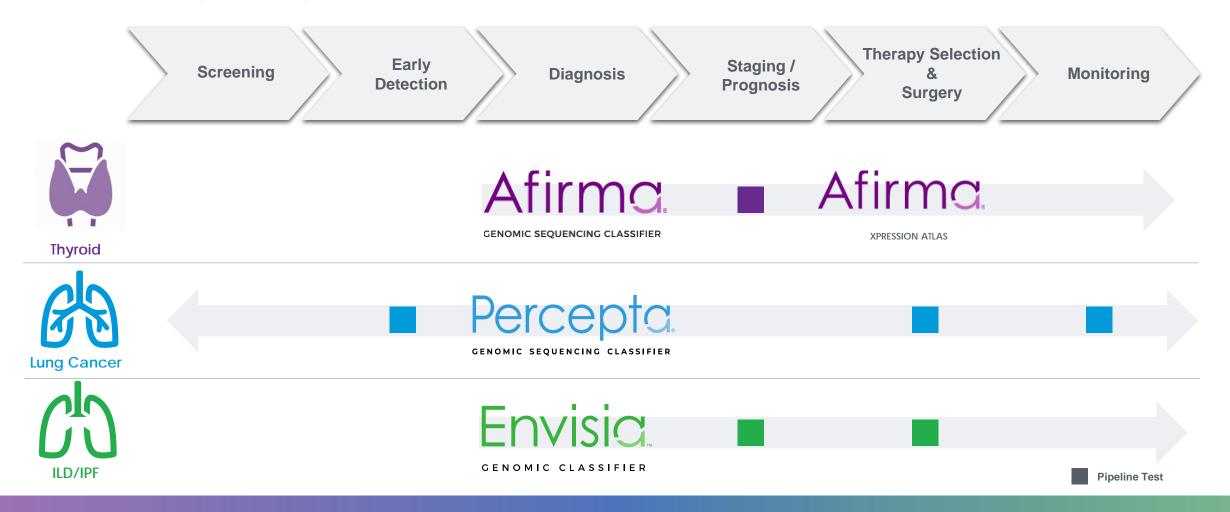
VARIANTS INFORM SURGICAL AND TREATMENT DECISIONS AT TIME OF DIAGNOSIS New Target Discovery

Clinical Trial Eligibility

Treatment Selection



Expanding Along the Value Chain



DELIVERING VALUE TO PATIENTS, CLINICIANS AND PAYORS ACROSS THE CLINICAL CARE CONTINUUM



Leading in the Age of Evidence

Deep library of clinical evidence published in top-tier journals



3 clinical validation

2 analytical verification

24 clinical utility, including 3 long-term clinical outcome

2 cost-effectiveness and quality-of-life

Percepta

BRONCHIAL GENOMIC CLASSIFIER

2 clinical validation

1 analytical verification

2 clinical utility

1 cost-effectiveness

PERCEPTA Registry with ~ 775 enrolled patients

Envisia

GENOMIC CLASSIFIER

3 clinical validation

1 analytical verification

1 clinical utility

BRAVE ongoing clinical trial ~ 450 patients





THE LANCET
Respiratory Medicine







GENOMIC SEQUENCING CLASSIFIER

Percepta.

GENOMIC SEQUENCING CLASSIFIER





SALES SPECIALIST ACROSS ALL PRODUCTS, PULMONARY CLINICAL EXPERTS, ACCOUNT SPECIALISTS FOR TOP ACCOUNTS

Evidence

SCIENTIFIC RIGOR FOCUSED ON GENERATING DATA, PUBLICATIONS, GUIDELINES AND COMMERCIAL ADOPTION

Reimbursement

IN-NETWORK STATUS
WITH NEARLY ALL
MAJOR U.S. HEALTH PLANS

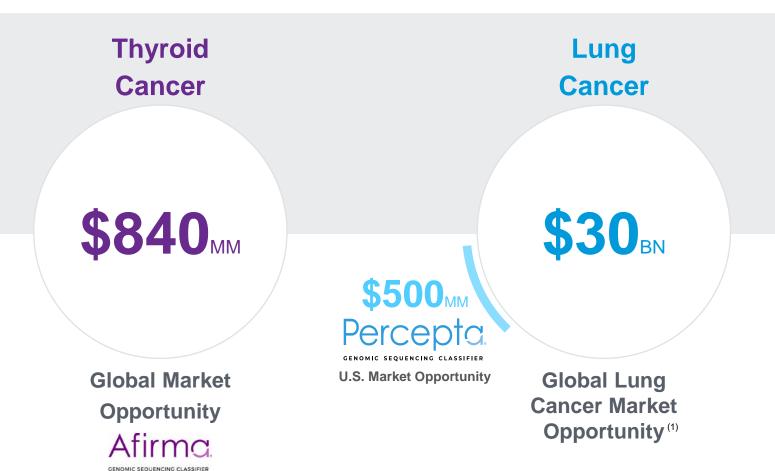
EXPECT TO ACHIEVE OPERATING CASH FLOW BREAKEVEN BEFORE END OF 2019



Our Founding Strategy: Improve Diagnostic Accuracy Hundreds of thousands of patients evaluated for suspected disease



Robust Market Opportunity Across Multiple Diseases



Idiopathic Pulmonary Fibrosis (IPF)

\$560_{MM}

U.S. + EU Market
Opportunity
Envision

Estimated market sizes based on Company estimates

(1) Market size based on Company estimates and includes nasal swab test for early detection



In Thyroid Cancer

An Incredibly Inefficient and Avoidable Diagnostic Paradigm

~15% to 30%

YIELD INCONCLUSIVE RESULTS

525_K
FINE NEEDLE ASPIRATIONS
PER YEAR TO EVALUATE
THYROID NODULES

Majority

OF PATIENTS WITH INDETERMINATE RESULTS UNDERGO SURGERY

~75%

DEEMED BENIGN POST-OPERATIVELY

Challenging diagnosis with

100_K +

UNNECESSARY SURGERIES PERFORMED IN U.S. ANNUALLY (1)

Source: Company estimates (1) ~180k surgeries performed to diagnose ~60K cancers

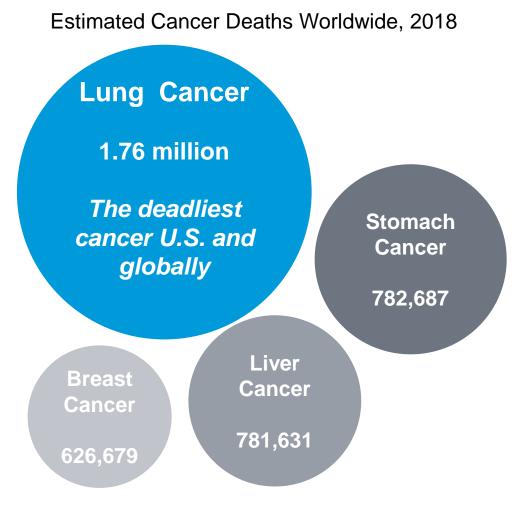
Improving Patient Outcomes in Thyroid Cancer

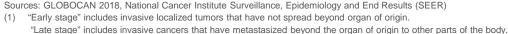
525,000 patients in U.S. evaluated for suspected thyroid cancer annually **Ultrasound-guided FNA Diagnosis Uncertain** >100,000 **Xpression Atlas** Reduce Afirma Suspicious Benign Unnecessary Inform treatment decisions to improve surgeries by patient outcomes ~70% GENOMIC SEQUENCING CLASSIFIER

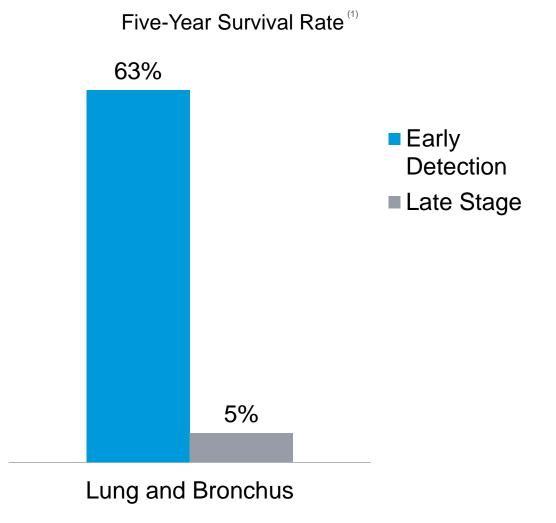


In Lung Cancer

Early Detection and Improved Diagnosis are Key to Saving Lives



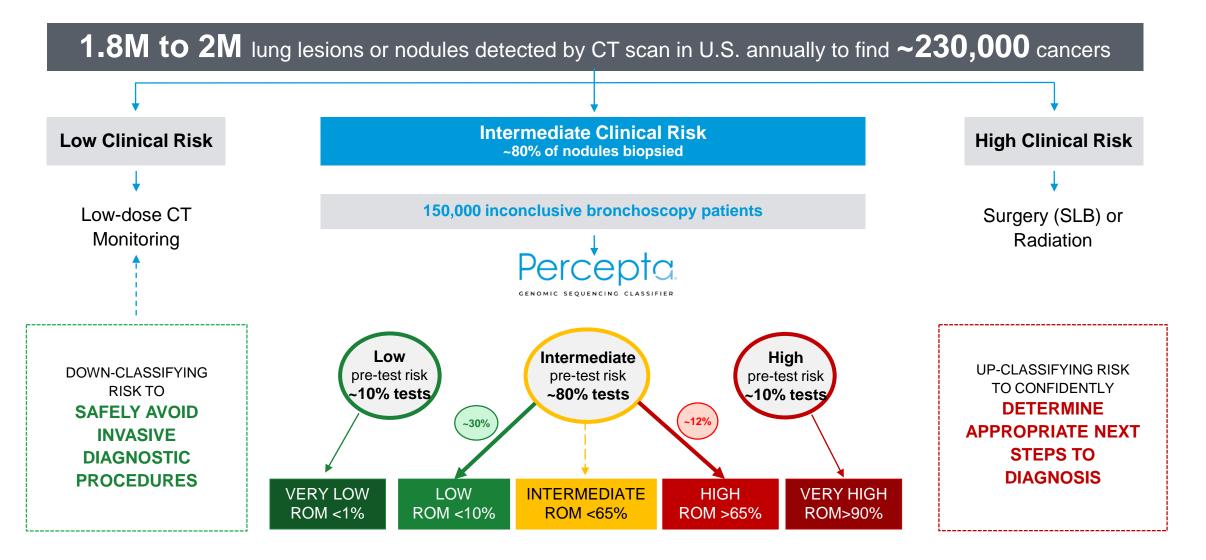








Improving the Efficiency of Lung Cancer Diagnosis





In Idiopathic Pulmonary Fibrosis (IPF)

A Progressive, Life-Threatening, Interstitial Lung Disease



Notoriously difficult to diagnose

LEADING TO TREATMENT DELAYS, PROLONGED MISDIAGNOSIS, PATIENT DISTRESS AND ADDED HEALTHCARE EXPENSE

Median survival time 2.5 years

LIFE EXPECTANCY WITH IPE IS WORSE THAN THAT OF MANY CANCERS.

IMPROVING DIAGNOSIS TO INFORM TREATMENT THAT COULD SAVE LIVES

Sources: American Lung Association; Ley B, et al. Clinical Course and Prediction of Survival in Idiopathic Pulmonary Fibrosis. *AJRCCM* 2011; Hutchison J, et al. Increasing Global Mortality from Idiopathic Pulmonary Fibrosis in the Twenty-First Century. *Annals ATS* 2014



IPF: Accelerate Diagnosis to Get Patients Life-Extending Treatment Faster

~200,000 patients in U.S. and Europe evaluated for suspected ILD

High Resolution CT

Lack highly confident Dx

~150,000 up to 75%*



Classic UIP pattern improves confidence in IPF diagnosis without surgery



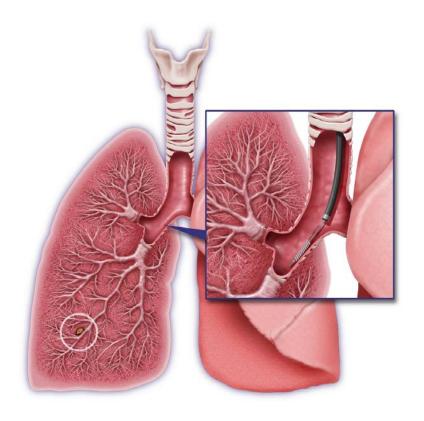
^{*} Based on Company estimates

[®] Veracyte, Inc. All rights reserved. September 2019

Pipeline Highlight: Nasal swab test for early lung cancer detection

Using novel, proprietary field of injury science that powers Percepta.





Peripheral lung nodules **difficult to biopsy** leading to late-stage diagnosis

Exposures such as smoking leads to genomic alterations in airway detected from a simple brushing or swab

Percepta classifier is based on bronchial airway brushing

New nasal swab test is designed to **detect lung cancer** from genomic alterations in the nose



Shared Gene Expression Alterations in Nasal and Bronchial Epithelium for Lung Cancer Detection - Perez-Rogers J, et al. JNCI J Natl Inst. 2017

NON-INVASIVE NASAL SWAB TEST DESIGNED FOR EARLY DETECTION TO SAVE LIVES



Strategic Collaboration Accelerates Pipeline and Expands Market Opportunity



Accelerates two key programs for Veracyte:

- Nasal swab test designed for early lung cancer detection (expect preliminary data by end of 2019)
- ✓ Commercialization of second-generation Percepta classifier, deploying RNA whole-transcriptome sequencing platform
- +\$50M in monetary and non-monetary value*

*\$20MM cash and estimated \$30M value on cohorts

FURTHER STRENGTHENS OUR LEADING POSITION IN LUNG CANCER DIAGNOSIS



Experienced Team with Track Record of Success

Bonnie Anderson Chairman and Chief Executive Officer	BECKMAN
Keith Kennedy Chief Financial Officer and Chief Operating Officer	A C P GE Capital
Giulia Kennedy, Ph.D. Chief Scientific and Medical Officer	CHIRON MILLENNIUM THE TAKEDA ONCOLOGY COMPANY THE TAKEDA ONCOLOGY COMPANY
John Hanna Chief Commercial Officer	IBM. Humana.
Freddie Bowie, Ph.D. Vice President, Corporate and Business Development	FOUNDATION DANAHER BOSTON CONSULTING GROUP



Catalysts to Drive Continued Momentum in 2019







Revenue Growth



Launch of next-generation
Percepta classifier
in mid-2019



Evidence Development



Spotlight clinical utility data



Pipeline Advancement

Field of injury advances; early data on nasal swab test







VCYT: A Compelling Value Proposition

Proven Model of Success

Answering clinical questions that matter

Novel RNA whole-transcriptome sequencing and machine learning scientific platform

Clinical evidence published in top-tier journals

First-to-market, first-to-coverage

Clinically impactful tests

Address large, underserved thyroid cancer, lung cancer and idiopathic pulmonary fibrosis markets

Significant growth opportunity

Current and pipeline products address market opportunity of more than \$30 billion

Experienced management team

with deep expertise and proven track record

Continued strong momentum

positions VCYT for near- and long-term success

