



# Myriad Genetics Fiscal Second-Quarter 2018 Earnings Call

02/06/2018



# Forward Looking Statements

## Forward Looking Statements

Some of the information presented here today may contain projections or other forward-looking statements regarding future events or the future financial performance of the Company. These statements are based on management's current expectations and the actual events or results may differ materially and adversely from these expectations. We refer you to the documents the Company files from time to time with the Securities and Exchange Commission, specifically, the Company's annual reports on Form 10-K, its quarterly reports on Form 10-Q, and its current reports on Form 8-K. These documents identify important risk factors that could cause the actual results to differ materially from those contained in the Company's projections or forward-looking statements.

## Non-GAAP Financial Measures

In this presentation, the Company's financial results and financial guidance are provided in accordance with accounting principles generally accepted in the United States (GAAP) and using certain non-GAAP financial measures. The Company's financial measures under GAAP include substantial one-time charges related to its acquisitions and ongoing amortization expense related to acquired intangible assets that will be recognized over the useful lives of the assets and charges related to executive severance. Management believes that presentation of operating results that excludes these items provides useful supplemental information to investors and facilitates the analysis of the Company's core operating results and comparison of operating results across reporting periods. Management also uses non-GAAP financial measures to establish budgets and to manage the Company's business. A link to reconciliation of the GAAP to non-GAAP financial guidance is provided above.

Financial Guidance	Fiscal Year 2018
GAAP diluted earnings per share	\$1.82 - \$1.87
Acquisition – amortization of intangible assets	\$0.52
Change in contingent consideration	(\$0.85)
Tax reform impact on deferred taxes	(\$0.46)
One time charges	\$0.08
<b>Non-GAAP diluted earnings per share</b>	<b>\$1.11 - \$1.16</b>
	Fiscal Third-Quarter 2018
GAAP diluted earnings per share	\$0.11 - \$0.13
Acquisition – amortization of intangible assets	\$0.12
One time charges	\$0.03
<b>Non-GAAP diluted earnings per share</b>	<b>\$0.26 - \$0.28</b>

For additional information on GAAP to non-GAAP reconciliation see:

<https://www.myriad.com/investors/gaap-to-non-gaap-reconciliation/>



# FY 2018 Second-Quarter Financial Results

*Significantly Exceed Expectations*

	2Q18 Actual Results	2Q17 Actual Results	YoY Change
Revenue (in mil.)	\$194.0	\$196.5	(1%)
GAAP EPS	\$0.45	\$0.09	400%
Adjusted EPS	\$0.31	\$0.26	19%



# Critical Success Factors to Achieving Strategic Goals

## STRATEGIC GOALS

**>10%**

Revenue Growth

**>30%**

Operating Margin

7 Products

**>\$50M**

**>10%**

International Revenue

## CRITICAL SUCCESS FACTORS

**Build upon solid hereditary cancer foundation**

**Grow new product volume**

**Expand reimbursement coverage for new products**

**Increase RNA kit revenue internationally**

**Improve profitability with Elevate 2020**



# Solid Hereditary Cancer Foundation

*4<sup>th</sup> Straight Quarter With Year-Over-Year Volume Growth*

- Hereditary cancer revenue up slightly sequentially in-line with expectations
- 4<sup>th</sup> straight quarter with YoY volume growth
- Exceeded 3% volume growth target in 2Q18
- Successful riskScore™ launch led to accelerating growth in Preventive Care

## Key Drivers of Volume Trends

Competitor Quality Concerns

Customizable Panels

U.S. Oncology & ION

Digital Integration

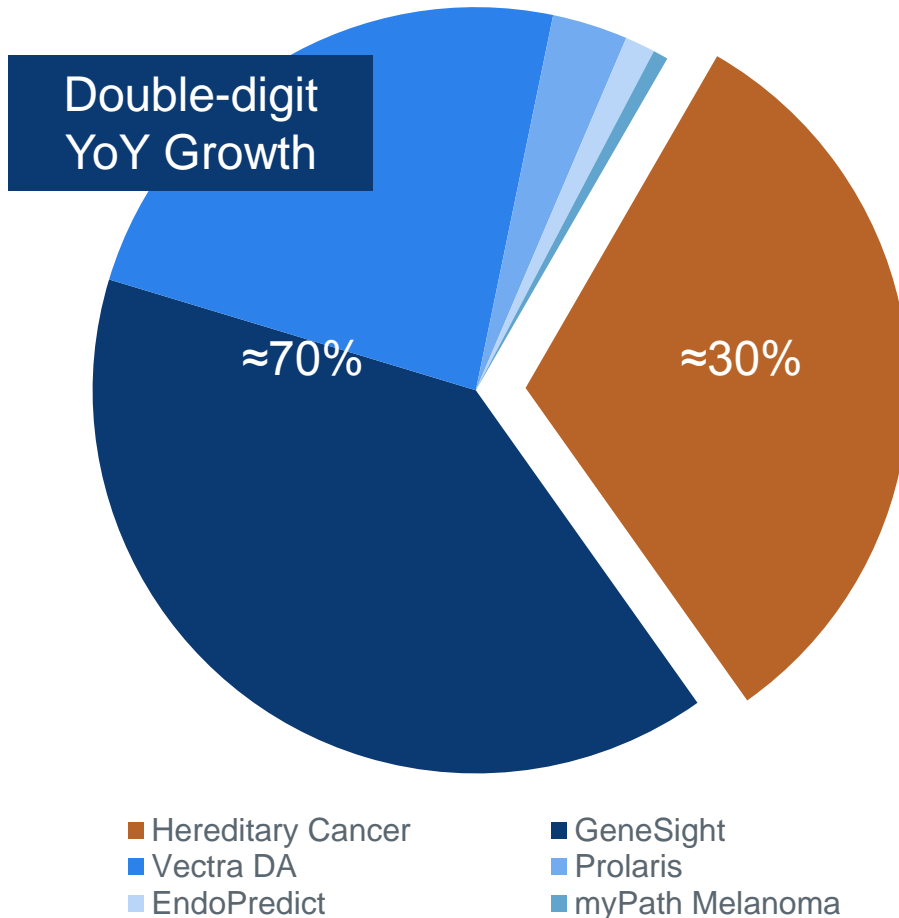
riskScore®



# Grow New Product Volume

*New Product Volume Grows Double-Digits; Revenue Sets New Record*

## Test Volume



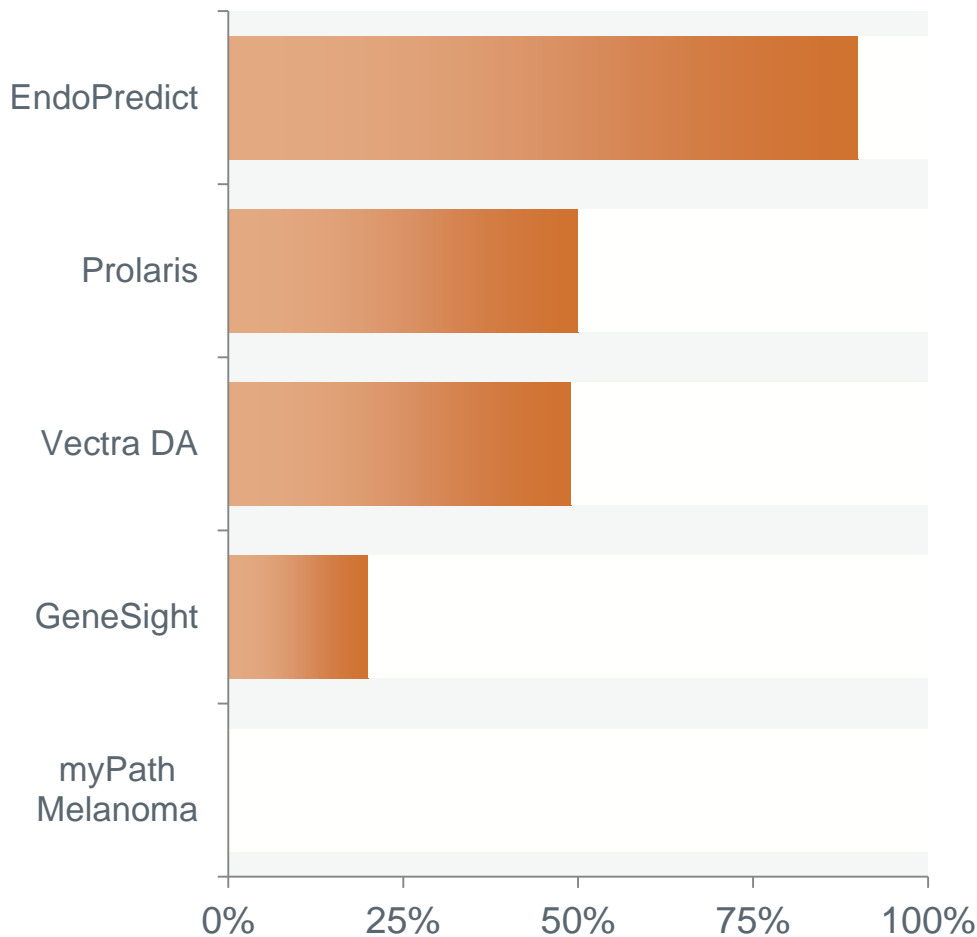
- New products comprise ≈70% of test volume
- New product YoY volume grew at double-digit rate
- New products set new record at 35% of total revenue
- EndoPredict volume up >70% sequentially



# Expand Reimbursement Coverage For New Products

*Several Key Reimbursement Catalysts in Fiscal Year 2018*

Current U.S. Reimbursement

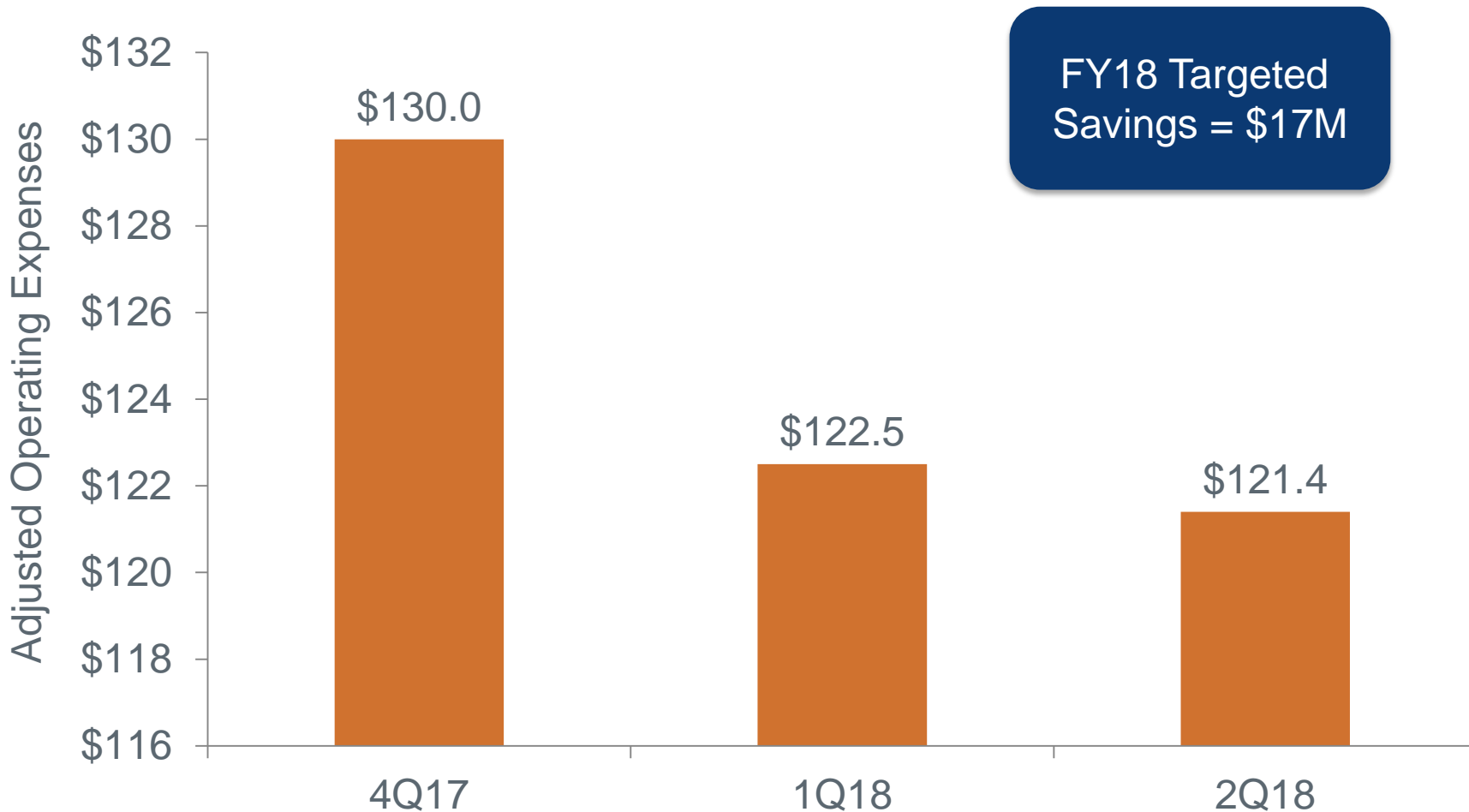


- Final Medicare LCD for EndoPredict became effective Jan. 30
- Prolaris favorable intermediate covered by Medicare
- Potential NCCN guidelines for Prolaris
- New Vectra DA utility guiding studies at ACR
- GeneSight trial showed statistically significant improvement in response and remission in MDD
- IMPACT study shows primary care MDs have even better outcomes than psychiatrists with GeneSight
- Potential NCCN guidelines for myPath Melanoma



# Improve Profitability With Elevate 2020

2<sup>nd</sup> Straight Quarter of Operating Expense Declines



FY18 Targeted Savings = \$17M





# FY 2018 Second-Quarter Revenue By Product

(in millions)

Product	2Q18	2Q17	YoY Growth
Hereditary Cancer	\$126.9	\$143.9	(12%)
GeneSight	\$31.7	\$21.7	46%
Vectra DA	\$11.1	\$10.7	4%
Prolaris	\$5.0	\$3.1	61%
EndoPredict	\$2.0	\$1.6	25%
Other	\$2.5	\$2.9	(14%)
<b>Total Molecular Diagnostic Revenue</b>	<b>\$179.2</b>	<b>\$183.9</b>	<b>(3%)</b>
Pharmaceutical & Clinical Services	\$14.8	\$12.6	18%
<b>Total Revenue</b>	<b>\$194.0</b>	<b>\$196.5</b>	<b>(1%)</b>



# Fiscal Second-Quarter Financial Results

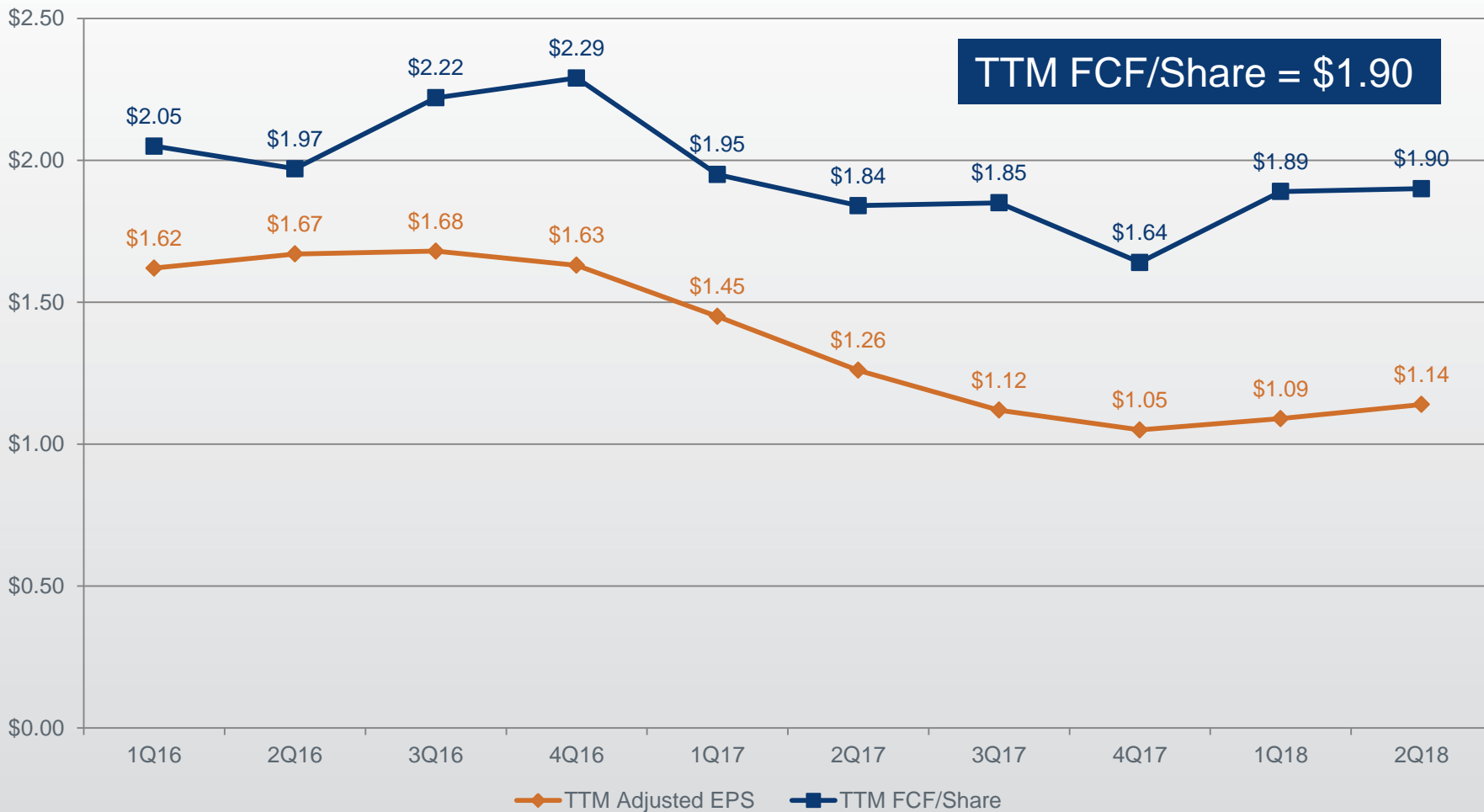
*Adjusted Earnings Per Share Increase 19% Over Q2 FY2017*

	2Q18	2Q17	YoY Growth
Total Revenue	\$194.0	\$196.5	(1%)
Gross Profit	\$149.6	\$152.1	(2%)
Gross Margin	77.1%	77.4%	-30 bps
Operating Income	\$4.4	\$17.0	(74%)
<b>Adjusted Operating Income</b>	<b>\$28.2</b>	<b>\$23.6</b>	<b>20%</b>
Adjusted Operating Margin	14.5%	12.0%	+250 bps
Net Income	\$32.1	\$5.9	444%
Diluted EPS	\$0.45	\$0.09	400%
<b>Adjusted EPS</b>	<b>\$0.31</b>	<b>\$0.26</b>	<b>19%</b>



# Comparison of Adjusted EPS and FCF/Share

*Adjusted EPS Significantly Understates Cash Earnings Power*





# FY18 and 3Q18 Financial Guidance







*Raising FY18 Financial Outlook*

Metric	Fiscal Year 2018	Fiscal Third-Quarter 2018
Revenue	\$760 to \$770 million	\$186 to \$188 million
GAAP Diluted EPS	\$1.82 to \$1.87	\$0.11 to \$0.13
Adjusted EPS	\$1.11 to \$1.16	\$0.26 to \$0.28



# Potential Financial Catalysts

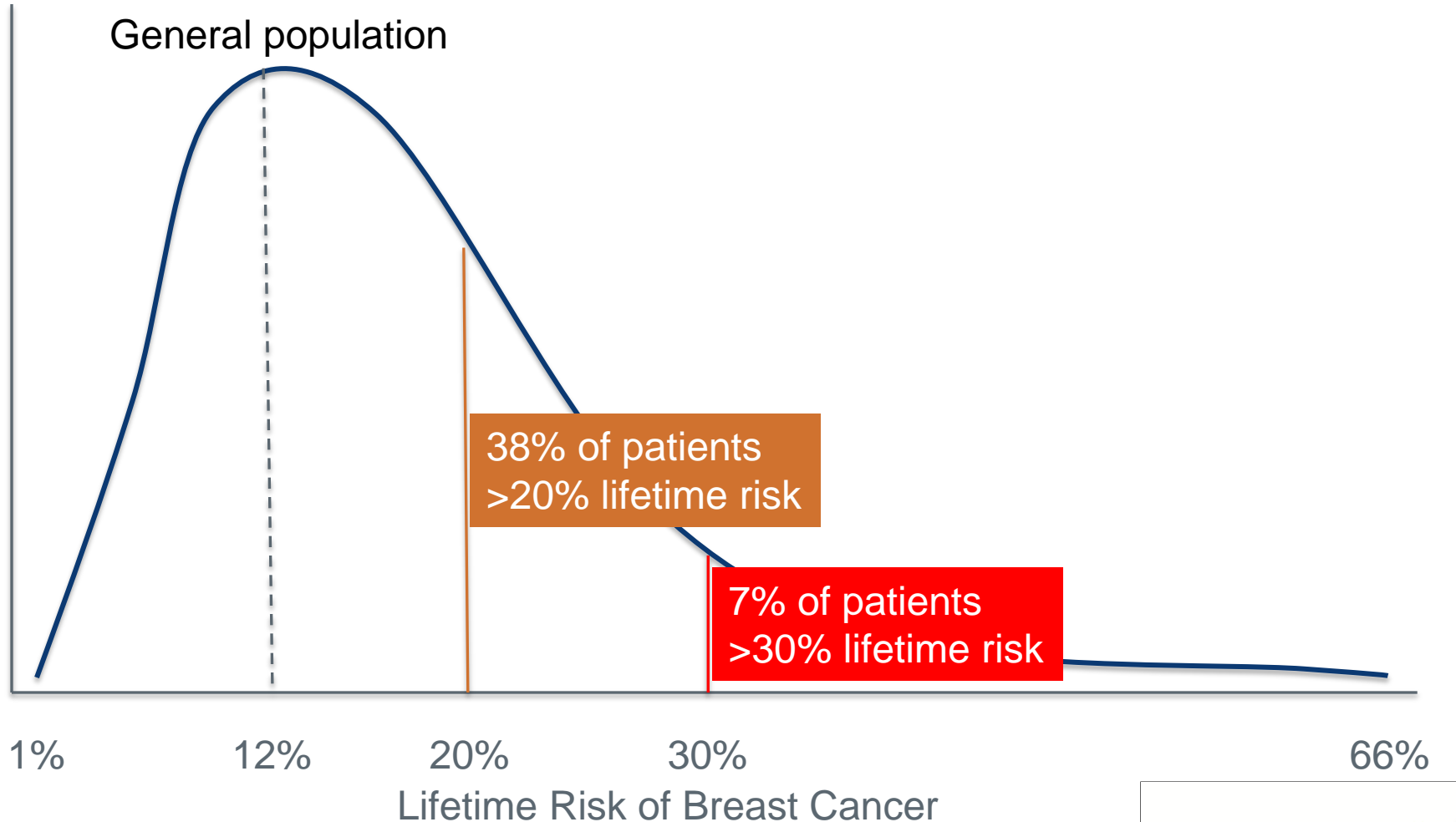
*Good Progress on Multiple Possibilities for Material Upsides*

Product	Potential Catalyst	Progress
 	Continued Improvements in Volume Growth	>3% growth in 1H18
<b>BRACAnalysisCDx</b>	Metastatic Breast Cancer Indication	FDA approval in 3Q18
	Additional Reimbursement	Successful RCT study Successful IMPACT study
	ACR Guidelines & Reimbursement	Increased Medicare rate under PAMA
	Additional Reimbursement	Increased Medicare rate under PAMA; new NCCN guidelines
<b>EndoPredict</b>	Increased Adoption in U.S.	2% U.S. Market Share
	Additional Reimbursement	New NCCN guidelines



# riskScore Validation Presented at SABCS

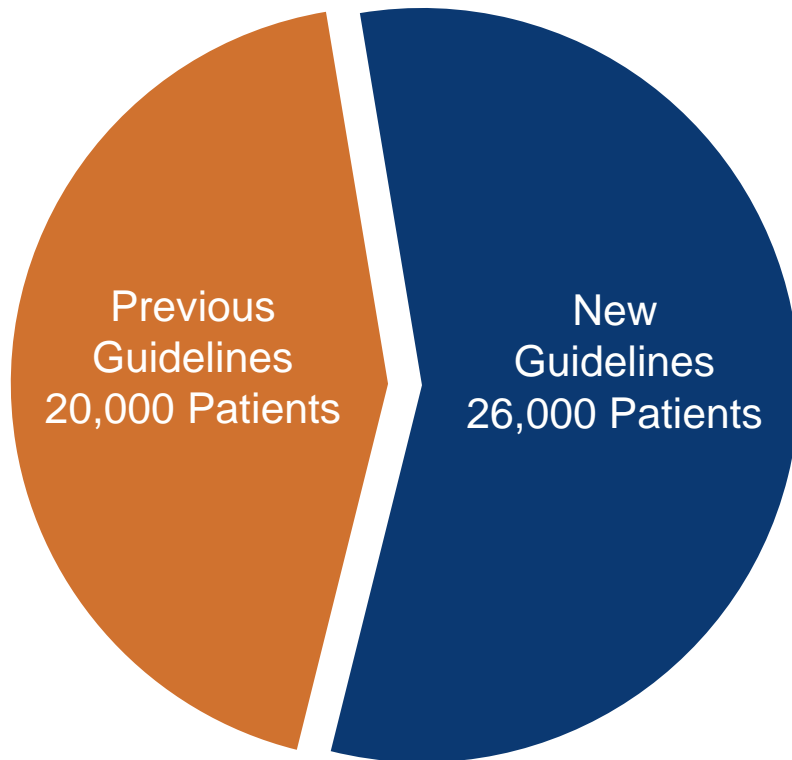
*Identifies More Elevated Risk Patients Than BRCA Testing*



# Expanded Opportunity in Hereditary Prostate Cancer

Testing Volume Up 10x in Last Year

46,000 Patient Opportunity

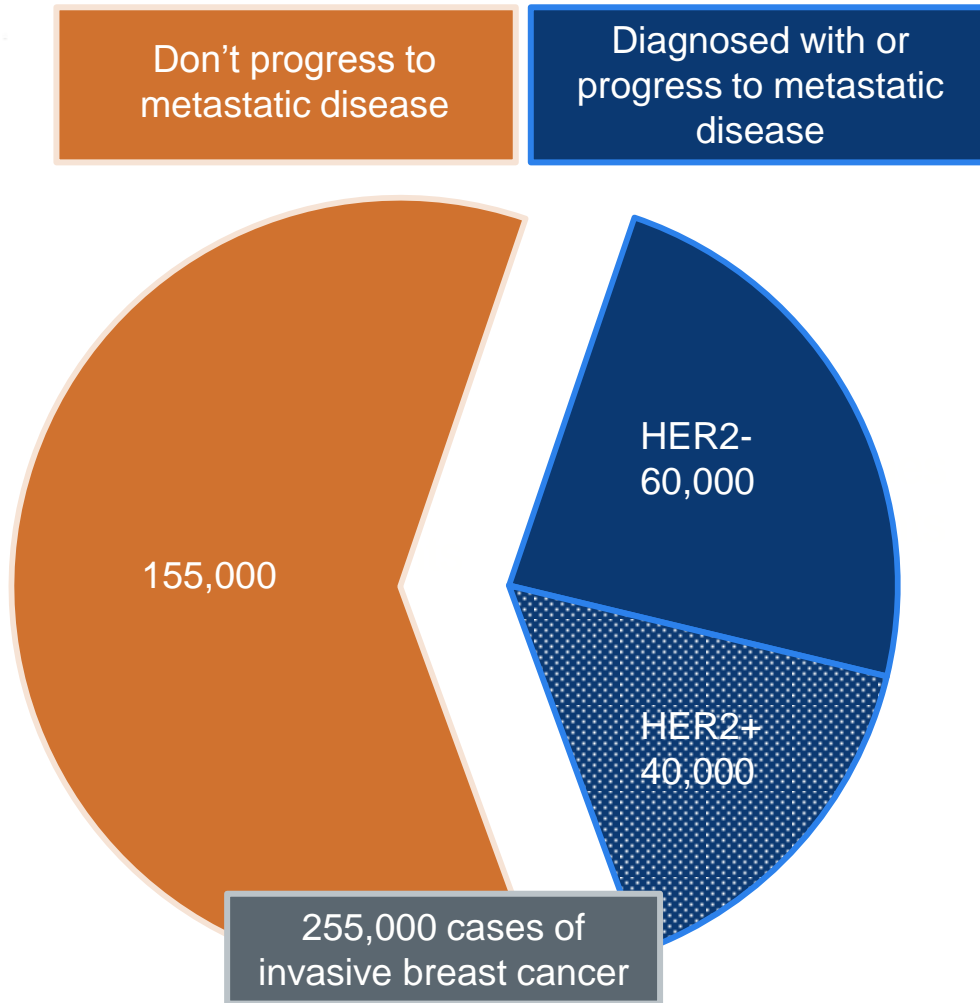


- Gleason Score  $\geq 7$  & Family History
- Metastatic Prostate Cancer

- New consensus guidelines published in the *Journal of Clinical Oncology* recommend routine counseling for prostate patients on hereditary cancer risks
- NCCN guidelines recommend all metastatic prostate cancer patients tested for hereditary mutations
- HOXB13 added to myRisk Hereditary Cancer Panel
- Commercializing through both Oncology and Urology channels

# Additional FDA Approval for BRACAnalysis CDx

125,000 Metastatic Breast Cancer Patients Now Eligible for Testing



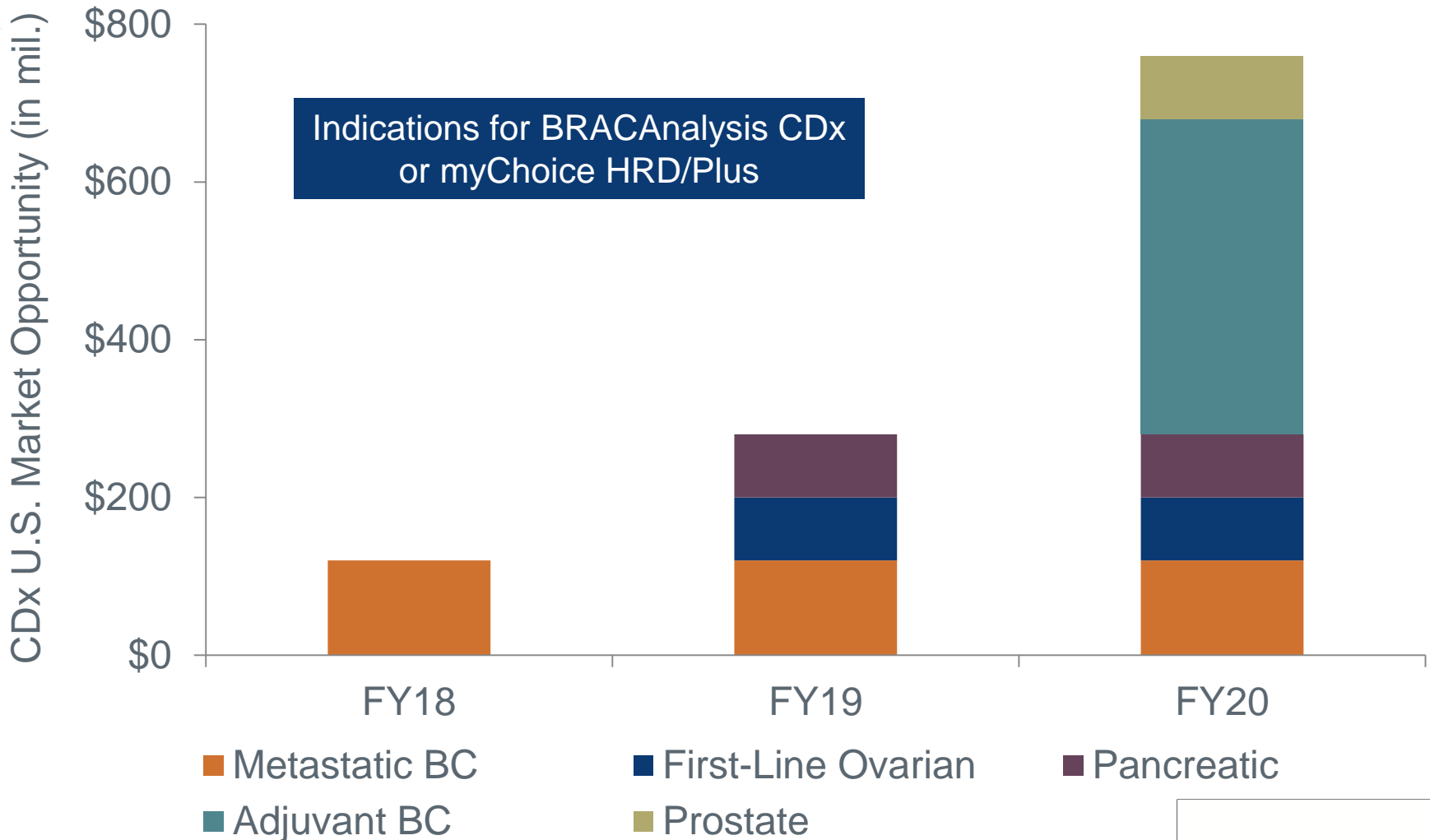
- 125,000 patients highly motivated to know BRCA status
- Collaborating with AstraZeneca and Merck
- Launched one of our largest digital marketing campaigns to drive patient awareness
- Increased targeting to 3,300 oncologists who treat 75% of patients with metastatic breast cancer





# Continued Opportunity for Market Expansion With CDx

*Additional Indications Could Total \$700M Opportunity*





# GeneSight Study Significant for Most Important Endpoints

*Beginning Discussions With Commercial Payers*

Study endpoint	What it Means	Study Result	Importance to Clinicians and Payers
<b>Remission hardest to achieve</b>	Patient no longer depressed	Highly statistically significant (p<0.01)	Very important
<b>Response difficult to achieve</b>	Patient feels a lot better	Highly statistically significant (p=0.01)	Very important
<b>Symptom Improvement most likely to achieve</b>	Patient feels somewhat better	Approaching statistical significance (p=0.1)	Meaningful

- Remission, response, and symptom improvement were durable and continued to improve over the 24-week study period
- 40 antidepressant FDA registration studies in the last 20 years:
  - All were compared to placebo, not active drug like GeneSight
  - Only 13% showed statistical significance for Remission
  - Only 33% showed statistical significance for Response



## Two Additional Major Studies on GeneSight Underway

*Will Further Strengthen Supporting Clinical Evidence for the Test*

### IMPACT Study

- Conducted in conjunction with the Canadian Centre for Addiction and Mental Health
- Open-label study with 8,000 patients enrolled to date with any mental health disorder
- Data on 2,000 depressed patients comparing outcomes between primary care and psychiatrists
- Primary care had even better outcomes than psychiatrists ( $p=0.0005$ )

### PRIME Care (VA) Study

- Conducted in conjunction with the Veterans Affairs Administration
- Randomized controlled trial planning to enroll 2,000 patients at 21 VA medical centers
- VA has committed \$12M to fund the study
- Important for establishing coverage for Dept. of Defense personnel



# Pathway To Expanded Vectra DA Reimbursement

## *Additional Data and ACR Guidelines by End of Fiscal Year*

Payer Questions	Progress
<input type="checkbox"/> Is Vectra DA included in guidelines?	<ul style="list-style-type: none"> <li>✓ Included in United Rheumatology guidelines in Dec. 2016 (represents 10% of rheumatologists)</li> <li>✓ Considered for ACR guidelines H2FY18</li> </ul>
<input type="checkbox"/> How does Vectra DA compare to historical measures of disease activity?	<ul style="list-style-type: none"> <li>✓ Two major studies presented at ACR 2017 showing Vectra DA was more than three times better than conventional disease activity measures</li> </ul>
<input type="checkbox"/> How should physicians modify treatment based upon Vectra DA score?	<ul style="list-style-type: none"> <li>✓ Publication submitted in Q3FY18 on meaningful change in Vectra DA score</li> <li>○ Medical management protocol to be added to the test report</li> </ul>
<input type="checkbox"/> When doctors follow the Medical Management Protocol does it lead to improved outcomes?	<ul style="list-style-type: none"> <li>○ Retrospective data available by end of FY18</li> <li>○ Prospective data from ongoing demonstration studies in two years</li> </ul>

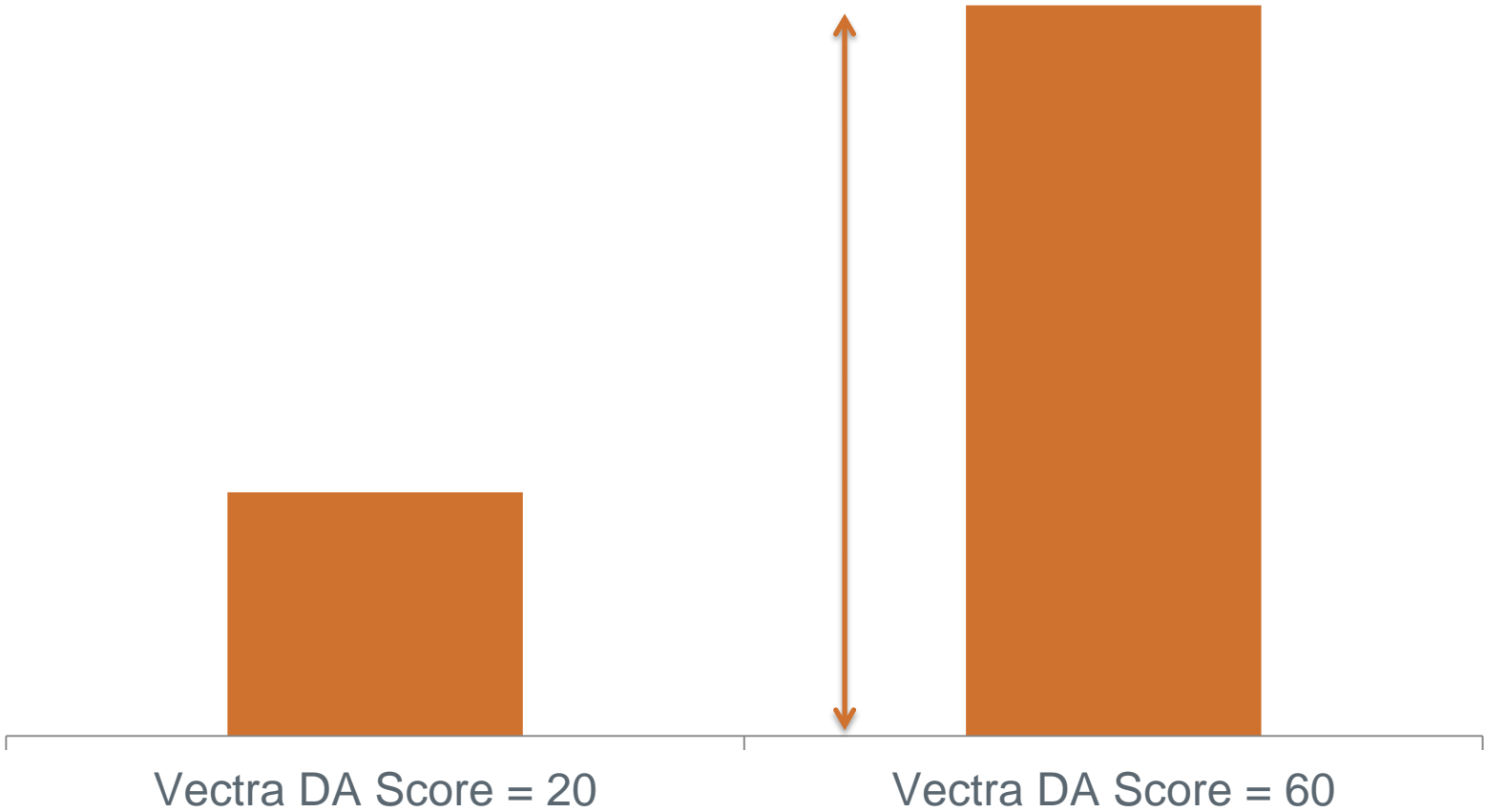


# New Data Supports Vectra DA Use for Assessing Cardiac Risk

*Potential Future Expanded Indication on Test Report*

Risk of Major Cardiac Event

≈3x Increased Risk



Source: Ann Rheumatic Disease Dec. 2017 - Biomarker-related risk for myocardial infarction and serious infections in patients with rheumatoid arthritis: a population-based study



# First Chemopredictive Data Presented on EndoPredict

*Highly Predictive of Response to Neoadjuvant Chemotherapy*

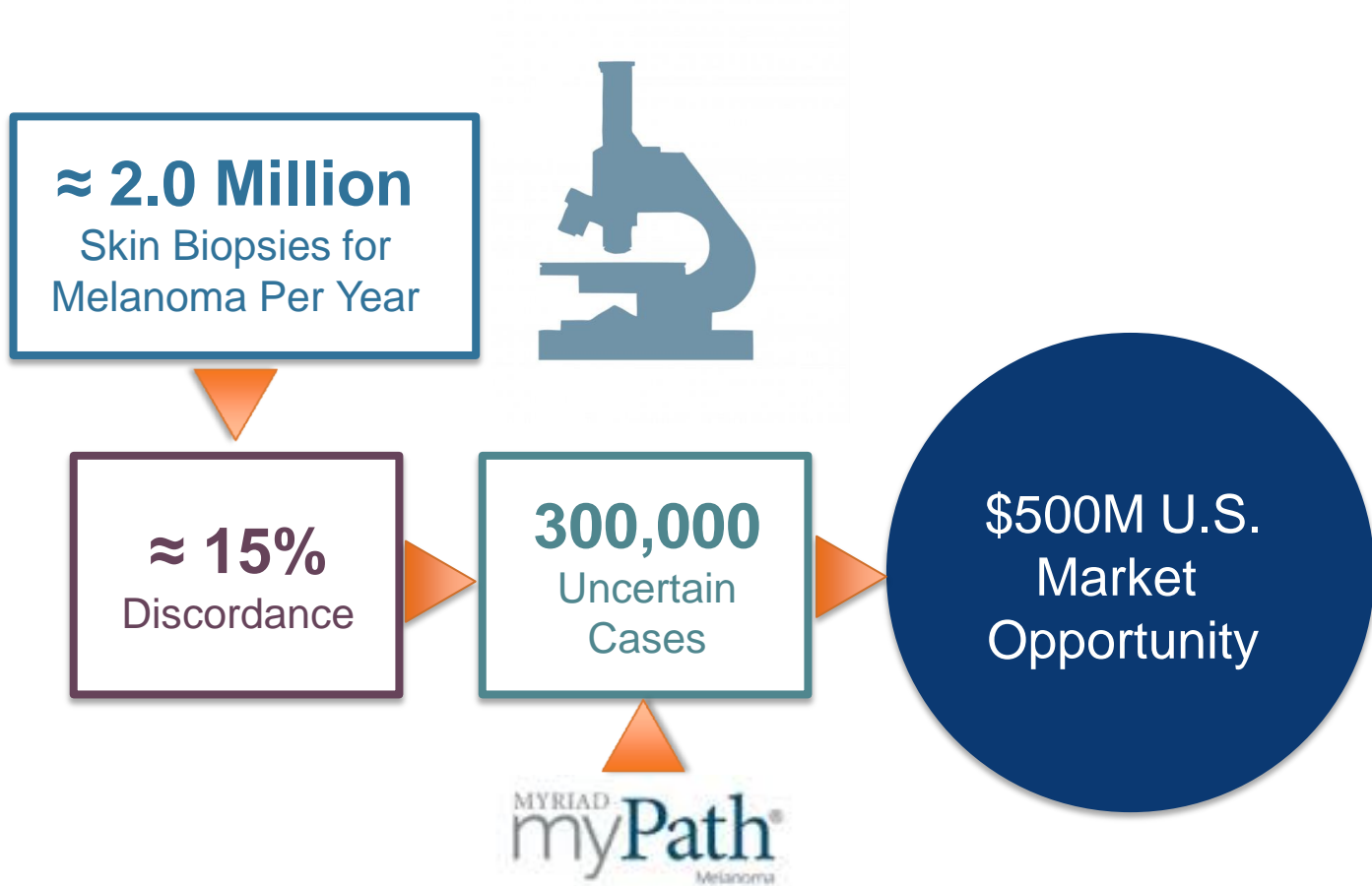
	<b>EndoPredict Low Score</b>	<b>EndoPredict High Score</b>	<b>p value</b>
Response to neo-adjuvant chemotherapy	0%	26.4%	0.0001
Response to endocrine therapy	27.3%	7.7%	0.015

Source: SABCs Presentation 2017 - The EndoPredict score predicts residual cancer burden to neoadjuvant chemotherapy and to neuroendocrine therapy in HR+/HER2- breast cancer patients from ABCSG34.



# myPath Melanoma Represents Significant Market

*300,000 Patients Per Year in the U.S. have Uncertain Diagnosis*



# Worldwide Leader in Personalized Medicine

- We are entering the **golden age** for personalized medicine
- Molecular diagnostics are the keystone for improving patient outcomes while eliminating waste in healthcare spending
- Myriad is the pioneer of “research-based” and “education-centric” business model for molecular diagnostics
- We are the best positioned company to lead this revolution in healthcare

