

H. C. Wainwright & Co. 2022 Global Invest Conference Pre-recorded May 20, 2022

Forward Looking Statements

The Private Securities Litigation Reform Act of 1995 provides a safe harbor from civil litigation for forward-looking statements accompanied by meaningful cautionary statements. Except for historical information, this presentation contains forward-looking statements within the meaning of Section 27A of the Securities Act of 1933, as amended, and Section 21E of the Securities Exchange Act of 1934, which may be identified by words such as "continues", "estimates", "anticipates", "projects", "plans", "seeks", "may", "will", "expects", "intends", "believes", "signals", "should", "can", "guidance" and similar expressions or the negative versions thereof and which also may be identified by their context. All statements that address operating performance or events or developments that Meridian Bioscience, Inc. ("Meridian" or "the Company") expects or anticipates will occur in the future, including, but not limited to, statements relating to per share diluted net earnings, sales, product demand, net revenues, operating margin, other guidance and the impact of COVID-19 on its business and prospects, are forward-looking statements. Such statements, whether expressed or implied, are based upon current expectations of the Company and speak only as of the date made. Specifically, Meridian's forward-looking statements are, and will be, based on management's then-current views and assumptions regarding future events and operating performance. Meridian assumes no obligation to publicly update or revise any forward-looking statements even if experience or future changes make it clear that any projected results expressed or implied therein will not be realized. These statements are subject to various risks, uncertainties and other factors that could cause actual results to differ materially, including, without limitation, the following:

Meridian's operating results, financial condition and continued growth depends, in part, on its ability to introduce into the marketplace enhancements of existing products that incorporate technological advances, meet customer requirements and respond to products developed by Meridian's competition, its ability to effectively sell such products and its ability to successfully expand and effectively manage increased sales and marketing operations. While Meridian has introduced a number of internally developed products and acquired products, there can be no assurance that it will be successful in the future in introducing such products on a timely basis or in protecting its intellectual property, and unexpected or costly manufacturing costs associated with its introduction of new products or acquired products could cause actual results to differ from expectations. Meridian relies on proprietary, patented and licenseed technologies. As such, the Company's ability to protect its intellectual property rights, as well as the potential for intellectual property litigation, would impact its results. Origing consolidations of reference laboratories and formation of multi-hospital alliances may cause adverse changes to pricing and distribution. Recessionary pressures on the economy and the markets in which the Company's customers operate, as well as adverse trends in buying patterns from customers, can change expected results. Costs and difficulties in complying with laws and regulations, including those administered by the United States Food and Drug Administration, and in complying with the original international states of the uncertainty of regulatory approvals and the regulatory process. The international scope of Meridian's operations, including changes in the relative strength or weakness of the U.S. delirance and product lines. There can be no assurance that additional acquisitions will be successfully integrated into Meridian's operations. There may be risks that acquisitions may disrupt operations and may pose p

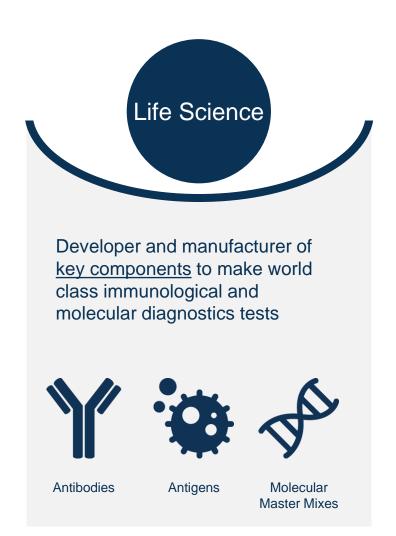


Intro to Meridian





Life Discovered. Life Diagnosed.







Life Science Products

Immunoassay Reagents Antigens & Antibodies





ToRCH







Tropical



Respiratory

Blockers





STD



Gastro



Microbial



Vet



Cancer



Cardiac



Autoimmune



DOA



Allergens







Air Dryable **Master Mixes**



Molecular Reagents

Inhibitor Tolerant Master Mixes



Extraction Controls



Specialized DNA Polymerases



Bst & Pfu



dNTPs

Key components supporting over 150 disease states and different specimen types





Diagnostics Products

Disease Areas

Gastrointestinal Pediatric & Neonatal C. difficile **Lead Poisoning** Calprotectin Congenital CMV H. pylori Group B Strep Campylobacter E. coli Crypto/Giardia **Healthcare Acquired Infections** Respiratory Flu Carba C Group A Strep C. difficile Mycoplasma Pneumoniae Pertussis Legionella RSV

Platforms & Product Families



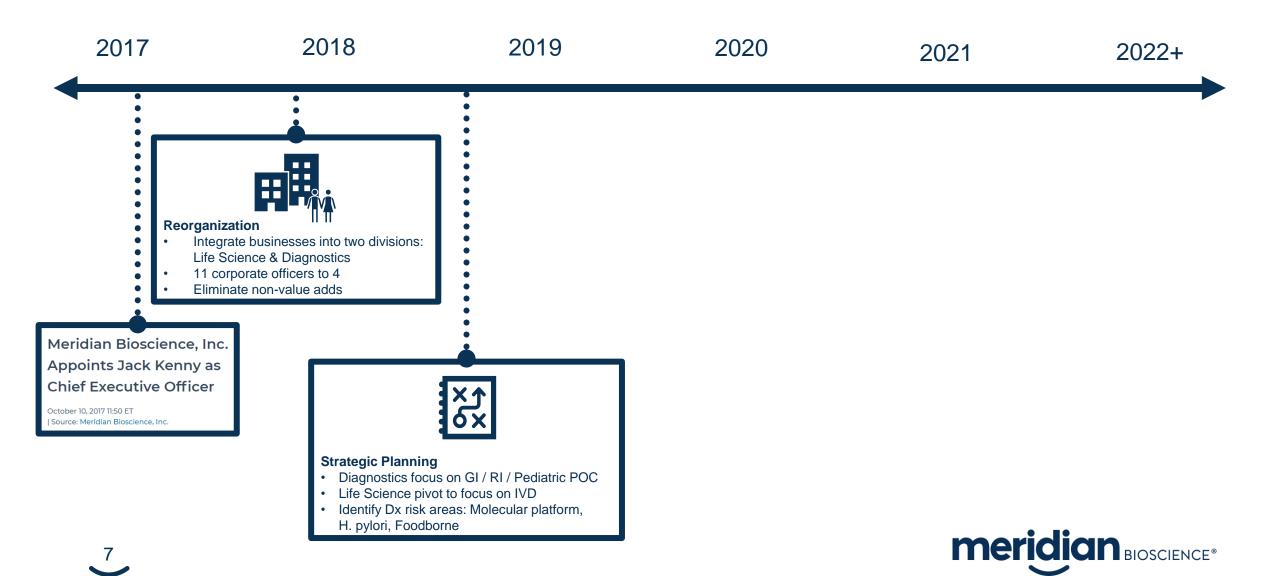


Historical Context





Transformation Process



Transformation Process

2017 2018 2019 2020 2021 2022+



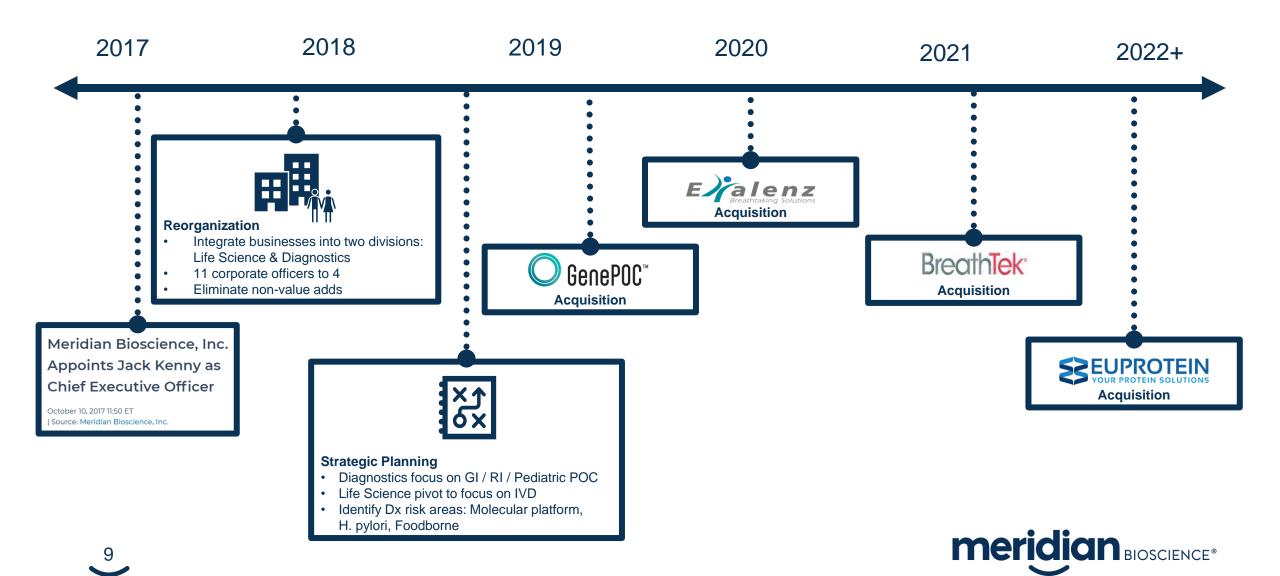
Strategic Planning

- Diagnostics focus on GI / RI / Pediatric POC
- Life Science pivot to focus on IVD
- Identify Dx risk areas: Molecular platform, H. pylori, Foodborne

Product Portfolio Investment: Immunoassay analyzer: Curian (March 2020) Next generation pediatric blood chemistry platform: (in-development) Menu expansion: Curian, Revogene, BreathID Acquisition: GenePoc, Exalenz, BreathTek Commercial Execution & Continuous Improvement Operational Pivot: Molecular sales to research & academia shifted to distributors Revamped commercial team to focus only on IVD Molecular new product development to solve development challenges of large IVD customers Acquisition: EUPROTEIN Commercial Execution & Continuous Improvement

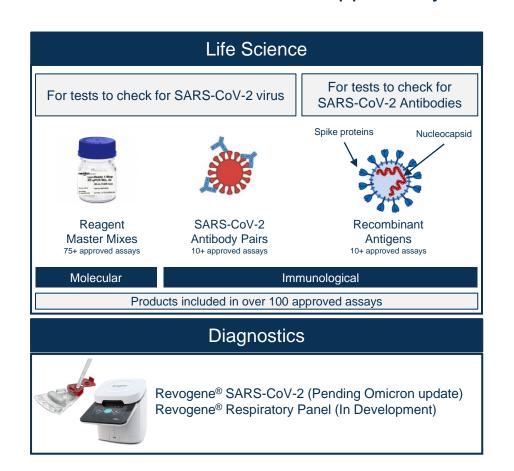


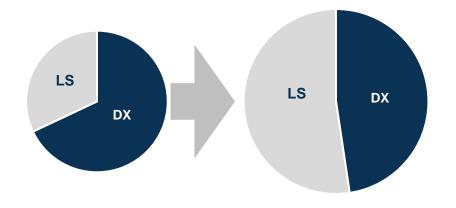
Transformation Process



COVID-19 Impact

STRATEGY: Maximize opportunity in Life Science / Keep Diagnostics focused on long-term plan





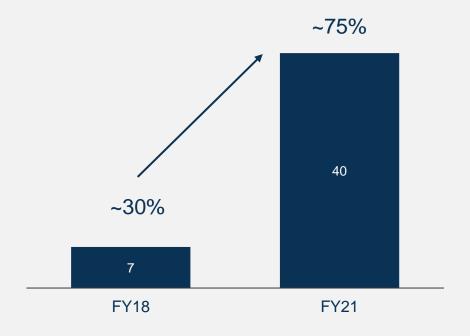
	FY2019 Actual		FY2020 Actual		FY2021 Actual
Diagnostics	\$137M	(11%)	\$121M	+5%	\$128M
Life Science	\$64M	+106%	\$133M	+43%	\$190M
Total	\$201M	+26%	\$254M	+25%	\$318M
Adj Op Margin %	19%		24%		30%



Life Science Major Customers

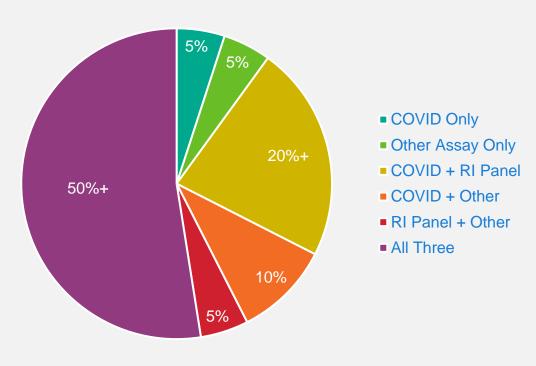
Number of Major Industry Customers⁽¹⁾

(% of total Life Science Net Revenues)



Registered Assay Diversity⁽²⁾

(% of total Major Industry Customers)



⁽¹⁾ IVD Customers generating \$1M+ in net revenues in select year, excludes academic, research, resellers and distributors unless customer has a registered assay



²⁾ Company estimates based on customer communications and analysis of sales trends

Growth Drivers





Disrupting Molecular Diagnostics Development

with innovative master mixes

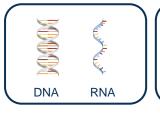
FAST TRACK YOUR R&D

Meridian's innovative sample-specific mixes (qPCR & LAMP)

SPECIMEN?



DETECTING?



KIT FORMAT?





INHIBITOR TOLERANT

Master Mixes have everything you need: Enzyme, Buffer, Excipients, Nucleotides, Mg

ONLY ADD

Primers & Probes

USE CRUDE

samples (no extraction)

PERFECT FOR

Multiplexing, fast cycling & POCT



Diagnostics R&D Pipeline

(as of 3/31/2022)

	Feasibility	Development	Clinicals	FDA
Breath				Liver MBT PMA Add'l Claim ⁽¹⁾
Immunoassay	Streptococcus pneumoniae / Legionella	C. difficile	Shiga Toxin	Campylobacter ⁽²⁾
Molecular		RI Panel GI Panel		COVID-19 EUA ⁽³⁾
Blood Chemistry		PediaStat Analyzer Lead		

Expected FY22 FDA Submissions

Add'l Claim

Shiga Toxin



⁽¹⁾ Submitted to FDA on 2/16/2022

⁽²⁾ Received FDA 510(k) clearance 12/23/2021

⁽³⁾ Received FDA STO(k) clearance 12/23/2021
(3) Received FDA Emergency Use Authorization (EUA) on 11/9/2021 – Not currently distributing pending changes to detect omicron variant

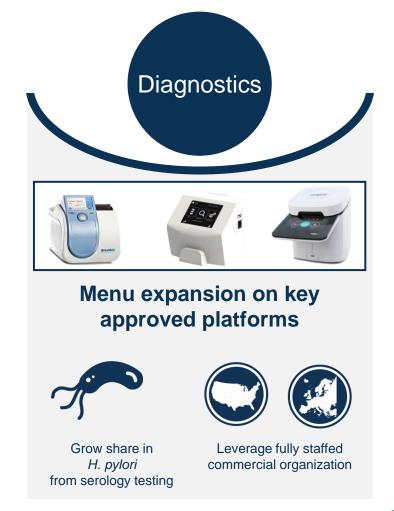
Investment Highlights





Key Growth Drivers







Meridian ESG Efforts Encompass Four Key Areas

Advancing Diagnostics to Enable Better Patient Healthcare Outcomes



- Accelerate diagnostic test development with high-quality Life Science products that power hundreds of diagnostic assays across 150+ disease states
- Supply hospitals, outpatient clinics, reference labs, and physician office labs with close to 200 diagnostics tests and transport media in more than 30 disease states
- First responder to the COVID-19 crisis, providing critical raw materials to enable rapid development of COVID-19 tests and now powering 100+ approved assays

Committing to Best-in-Class Governance Practices



- 88% of directors are considered independent
- Separated Chairman and CEO roles
- Aligned management compensation to clear financial targets

Reducing Environmental Impact with Product Innovation



- Reduce significant carbon emissions and Styrofoam packaging needs associated with cold chain shipping and storage of diagnostic assays through Lyo-Ready and Air-Dryable Mixes
- Enable broader distribution and use of diagnostics in resource-limited settings with limited cold chain management capabilities
- Lengthen shelf-life of molecular diagnostic assays to lower waste and improve assay availability

Creating a Diverse, Equitable, and Inclusive Workforce



- Introduced the One Meridian Inclusion Diversity and Equity Team to encourage and support an environment in which all employees feel included and empowered
- Approximately 55% of employees are women
- 25% of Meridian's US workforce is ethnically diverse





Investment Highlights

- COVID-19 pandemic beneficiary, not dependent
- Diagnostics business advancing turnaround poised to emerge with sustainable growth
- Life Science transformed with scale Building on pandemic, longterm customer relationships
- Strong balance sheet and cash generation to fuel organic and inorganic growth opportunities
- Potential trading multiple expansion in addition to earnings growth



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