1Q 2021 Results

May 4, 2021

Charles River Laboratories



Safe Harbor Statement

Caution Concerning Forward-Looking Statements. This presentation includes forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995. Forward-looking statements may be identified by the use of words such as "anticipate," "believe," "expect," "intend," "will," "may," "estimate," "plan," "outlook," and "project" and other similar expressions that predict or indicate future events or trends or that are not statements of historical matters. These statements also include statements about the impact of the COVID-19 pandemic for our business, financial condition and results of operations, including the long-term growth prospects and as compared to other companies, and the prospects for recovery therefrom; the effectiveness of our capital deployment strategy, in light of the COVID-19 pandemic and our ability to reduce capex, preserve jobs, support client research programs and sustain our financial position; our compliance with the maintenance covenants under our credit agreement; our projected 2021 and other future financial performance whether reported, constant currency, organic, and/or factoring acquisitions, with respect to Charles River as a whole and/or any of our reporting or operating segments or business units; our annual guidance and two-year targets; the assumptions that form the basis for our revised annual guidance; the expected performance of our venture capital and other strategic investments; the future demand for drug discovery and development products and services and our portfolio; the impact of foreign exchange; our expectations regarding stock repurchases and debt repayment; the development and performance of our services and products; market and industry conditions including industry consolidation, outsourcing of services and identification of spending trends by our clients and funding available to them; the potential outcome of, and impact to, our business and financial operations due to litigation and legal proceedings and tax law changes; our business

Forward-looking statements are based on Charles River's current expectations and beliefs, and involve a number of risks and uncertainties that are difficult to predict and that could cause actual results to differ materially from those stated or implied by the forward-looking statements. Those risks and uncertainties include, but are not limited to: the COVID-19 pandemic, its duration, its impact on our business, results of operations, financial condition, liquidity, business practices, operations, suppliers, third party service providers, clients, employees, industry, ability to meet future performance obligations, ability to efficiently implement advisable safety precautions, and internal controls over financial reporting; the COVID-19 pandemic's impact on client demand, the global economy and financial markets; the ability to successfully integrate businesses we acquire (including Cognate BioServices and risks and uncertainties associated with Cognate BioServices and reporting; the COVID-19 pandemic; industry, ability to meet future terms in a research and development spending, negative trends in areas that the Company did not previously operate); the timing and magnitude of our share repurchases; negative trends in research and development spending, negative trends in the level of outsourced services, or other cost reduction actions by our clients; the ability to convert backlog to revenue; special interest groups; contaminations; industry trends; new displacement technologies; USDA and FDA regulations; c

Regulation G

This presentation includes discussion of non-GAAP financial measures. We believe that the inclusion of these non-GAAP financial measures provides useful information to allow investors to gain a meaningful understanding of our core operating results and future prospects, without the effect of often one-time charges, consistent with the manner in which management measures and forecasts the Company's performance. The non-GAAP financial measures included in this presentation are not meant to be considered superior to or a substitute for results of operations prepared in accordance with GAAP. The company intends to continue to assess the potential value of reporting non-GAAP results consistent with applicable rules and regulations. In accordance with Regulation G, you can find the comparable GAAP measures and reconciliations to those GAAP measures on our website at ir.criver.com.



Estimates of COVID-19 Impact in 2020

The Company has provided its estimates for the impact from the COVID-19 pandemic in 2020, including on the Company's revenue. These estimates were determined using methodologies and assumptions that vary depending on the specific reporting segment and situation. For the Research Models and Services segment, estimates were primarily based on comparisons to daily historical research model sales volumes prior to the COVID-19 pandemic and the subsequent reduction in research model order activity associated with our clients' COVID-19 pandemic-related site closures and/or their reduced on-site activity, as well as our discussions with clients, particularly of our research model services and HemaCare businesses, with regard to revenue expectations and operational impacts from the COVID-19 pandemic. For the Discovery and Safety Assessment segment, estimates were based on multiple factors including, but not limited to, discussions with clients with regard to the cause of delays to discovery projects and safety assessment studies, location-specific actions to ensure employee safety in our facilities, the impact of remote versus in-person activities and services, and supply chain delays and other resource constraints. For the Manufacturing Support segment, estimates were based on multiple factors including, but not limited to, analysis of the sales impact due to the COVID-19 pandemic, assessments of idle instruments and the related revenue streams due to the inability to access clients' sites, as well as discussions with clients with regard to their revenue expectations and operations. The estimated revenue loss related to COVID-19 was also expected to be partially offset by incremental work on clients' COVID-19 programs. Because these estimates and assumptions involve risks and uncertainties, actual events and results may differ materially from these estimates and assumptions, and Charles River assumes no obligation and expressly disclaims any duty to update them.



Robust 1Q21 Performance

- Robust 1Q21 financial performance 13% organic revenue growth and 170 bps of non-GAAP operating margin improvement YOY – demonstrates:
 - Strength of the biopharmaceutical market environment
 - Power of our unique portfolio
 - Both of which we believe are as strong as they have ever been
- Clients are increasingly choosing to partner with CRL for:
 - Flexible and efficient outsourcing solutions
 - Scientific depth and breadth of our portfolio
 - Unwavering focus on seamlessly serving their diverse needs
- Clients are opting to work with a smaller number of CROs who offer broad scientific capabilities
 - Enables them to drive greater efficiency and accelerate speed of their research, non-clinical development, and manufacturing programs
- Complexity of scientific research is also increasing clients' reliance on a high-science outsourcing partner like CRL



Strategically Differentiating from the Competition

- To further differentiate CRL from competition, strategically expanding portfolio in areas that deliver greatest value to clients and offer significant growth potential
- Already in 2021, have enhanced scientific capabilities for advanced drug modalities through acquisitions of:
 - Distributed Bio
 - Cognate BioServices
 - Retrogenix
- Distributed Bio and Retrogenix strengthened Discovery portfolio
- Cognate (completed on March 29th) provides excellent growth opportunity by enabling us to offer CDMO services in high-growth, high-science, cell and gene therapy sector
- Very pleased to welcome the talented staff of each organization
- Believe that by continuing to invest in our portfolio and people, we are maintaining and enhancing our position as the leading, non-clinical CRO



Strong Trends Support Outlook for 2021

- Believe strength of our portfolio and robust industry fundamentals are leading to unprecedented client demand across most of our businesses
- In 1Q21, experienced continuation of robust demand from the end of 2020, including, new, record booking and proposal levels in Safety Assessment business
- Organic revenue growth above 10% for the second consecutive quarter, even after normalizing for last year's COVID-19 impact
- Overall, believe robust 1Q21 performance and solid business trends support our improved outlook for the year



1Q21 Revenue

(\$ in millions)	1Q21	1Q20	ΥΟΥ Δ
Revenue, reported	\$824.6	\$707.1	16.6%
(Increase)/decrease due to FX			(2.9)%
Contribution from acquisitions			(0.7)%
Revenue growth, organic			13.0%

- Quarterly revenue surpassed \$800M for first time
- Organic revenue growth driven by double-digit growth across all three business segments
- YOY comparison to last year's COVID-related revenue impact contributed ~140 bps to the 1Q21 revenue growth rate
 - Primarily affected the RMS segment
- Broad-based growth across all client segments, with biotech leading the way as they continued to benefit from robust funding

1Q21 Operating Margin

	1Q21	1Q20	Δ ΥΟΥ
GAAP OM%	15.0%	13.3%	170 bps
Non-GAAP OM%	20.7%	19.0%	170 bps

- Improvement driven by RMS and DSA
- Reflected operating leverage from robust revenue growth and continued efforts to drive efficiency
- Expect the same factors will drive non-GAAP operating margin improvement in 2021
 - Believe operating margin will approach 21%, above our target



1Q21 EPS

	1Q21	1Q20	ΥΟΥ Δ
GAAP EPS	\$1.20	\$1.02	17.6%
Non-GAAP EPS	\$2.53	\$1.84	37.5%

- Outstanding earnings growth principally reflected:
 - Double-digit revenue growth
 - Meaningful operating margin improvement



Increasing 2021 Guidance

- Based on 1Q21 performance and positive outlook for remainder of the year, meaningfully increasing revenue growth and non-GAAP EPS for 2021
- Normalized for last year's COVID impact, would still expect low-double-digit organic revenue growth
- Non-GAAP EPS represents 20%-23% growth YOY, an increase of \$0.75 at midpoint from prior outlook



Increasing 2021 Guidance Including Cognate

	CURRENT	PRIOR
Revenue growth, reported	19%-21%	16%-18%
Contribution from acquisitions	(4.5%)-(5.0%)	(4.5%)-(5.0%)
Decrease/(Increase) due to FX	<u>(~2.5%)</u>	<u>(2.0%)-(2.5%)</u>
Revenue growth, organic	12%-14%	9%-11%
GAAP EPS	\$5.95-\$6.20	
Acquisition-related amortization	\$2.15-\$2.40	
Acquisition-related adjustments	\$0.75-\$0.80	
Other items	~\$0.55	
Venture capital investment losses/(gains)	<u>\$0.25</u>	<u>==</u>
Non-GAAP EPS	\$9.75-\$10.00	\$9.00-\$9.25



DSA Results – Revenue

(\$ in millions)	1Q21	1Q20	Δ ΥΟΥ
Revenue, reported	\$501.2	\$438.7	14.2%
(Increase)/decrease due to FX			(2.3)%
Contribution from acquisitions			(0.3)%
Revenue growth, organic			11.6%

Revenue growth driven by broad-based demand for both Discovery and Safety Assessment (SA) services



DSA Results - Safety Assessment (SA)

- Performance reflected robust demand from biotech and biopharma clients and price increases
- Bookings and proposal volume reach record highs in 1Q21, with strength across all regions and major service areas
- Bookings increased substantially more than target
- Clients are expanding preclinical pipelines and intensifying focus on complex biologics
 - Provides us with greater visibility
- Believe they are securing space with CRL further in advance to ensure no delays to research
- Believe these trends position SA extremely well; support low-double-digit organic revenue growth for DSA segment
 - Higher than prior outlook
- Pleased with extensive depth and breadth of SA portfolio and remain intently focused on continuing to enhance value we provide to clients



DSA Results – Discovery Services

- Discovery Services had another exceptional quarter, led by broad-based demand for oncology, early discovery, and CNS services
- Efforts to broaden and strengthen discovery capabilities and enhance scientific expertise are enabling us to expand support for clients' discovery research
- Clients increasingly view CRL as a premier scientific partner who can support efforts to identify new drug targets and discover novel therapeutics
- Intend to continue to build Discovery portfolio so clients can outsource complex discovery projects to us, including advanced modalities
- Distributed Bio and Retrogenix enhanced our large molecule discovery capabilities
 - Retrogenix, through proprietary cell microarray technology, offers target receptor identification and off-target screening services
 - Will enhance clients' early discovery efforts and enable them to explore potential preclinical safety liabilities
- Combination of Distributed Bio and Retrogenix further strengthens our integrated, end-to-end solution for therapeutic antibody and cell and gene therapy discovery and development
- Continuing to add cutting-edge technologies through strategic partnership strategy, most recently with Valence Discovery, a new artificial intelligence (AI) drug discovery partner



DSA Results – Operating Margin

	1Q21	1Q20	Δ ΥΟΥ
DSA GAAP OM%	18.1%	16.5%	160 bps
DSA Non-GAAP OM%	23.8%	22.0%	180 bps

- Leverage from robust DSA revenue growth was primary driver of margin improvement
- Expect this trend to continue to propel DSA non-GAAP operating margin into the mid-20% range for 2021



RMS Results – Revenue

(\$ in millions)	1Q21	1Q20	ΥΟΥ Δ
Revenue, reported	\$176.9	\$146.0	21.2%
(Increase)/decrease due to FX			(4.2)%
Contribution from acquisitions			(2.2)%
Revenue growth, organic			14.8%

- Robust demand for research models in China was primary driver of 1Q21 revenue growth
- Higher revenue for research model services, including GEMS and CRADL, also contributed to revenue growth
- ~620 bps of growth was attributable to the comparison to last year's COVIDrelated revenue impact from client site closures and associated disruptions



RMS Results – Research Models

- Demand trends for research models largely consistent with trends prior to pandemic
- Growth in China widely outpacing mature markets
 - Research models business in China had an exceptional 1Q21, even normalizing for last year's COVID-19 impact
 - Driven by resurgence of demand across all client segments
- Biomedical research in China has returned to pre-COVID levels, and in some areas, even greater levels
- In U.S. and Europe, client order activity has also rebounded



RMS Results – RM Services

- RM Services also continued to perform well
- GEMS benefited from renewed outsourcing demand as clients seek greater flexibility and efficiency afforded them when we manage their proprietary colonies, as we did for many clients during the pandemic
- Also continuing to generate substantial interest for CRADL (Charles River Accelerator Development Labs), as both small and large biopharma clients seek turnkey research capacity
 - Allows them to invest in people and research rather than infrastructure
- To accommodate demand, actively expanding existing capacity in Boston/Cambridge and South San Francisco (SSF) biohubs
- Utilizing CRADL also provides clients with collaborative opportunities to seamlessly access other CRL services, from Discovery to GEMS, further enhancing the speed and efficiency of their research programs



RMS Results – HemaCare & Cellero

- Cell supply revenue growth remained below targeted level in 1Q21, due to some limitations on donor access
 - Believe revenue will increase during 2021 as donor availability continues to improve
- Continuing to work diligently to expand donor base in the U.S. and add more comprehensive capabilities at all sites to accommodate robust demand in broader cell therapy market
- Believe Cognate acquisition is particularly timely because it creates new business opportunities for HemaCare and Cellero in cell and gene therapy development
- Expanded capabilities are establishing CRL as a trusted partner who can move clients' programs forward using the same cellular products through each step of research and early-stage development phases and into CGMP production



RMS Results – Operating Margin

	1Q21	1Q20	Δ ΥΟΥ
RMS GAAP OM%	25.4%	18.7%	670 bps
RMS Non-GAAP OM%	28.7%	23.0%	570 bps

- Significant margin improvement is due to:
 - Comparison to 23% non-GAAP operating margin in 1Q20 which was depressed by onset of COVID-related disruptions and resulting impact on research model order activity
 - 1Q21 performance reflects operating leverage on robust revenue growth, particularly for research models in China



Manufacturing Results – Revenue

(\$ in millions)	1Q21	1Q20	ΥΟΥ Δ
Revenue, reported	\$146.5	\$122.4	19.7%
(Increase)/decrease due to FX			<u>(4.1)%</u>
Revenue growth, organic			15.6%

Revenue increase driven by double-digit growth in both Biologics Testing Solutions (Biologics) and Microbial Solutions



Manufacturing – Operating Margin

	1Q21	1Q20	Δ ΥΟΥ
Manufacturing GAAP OM%	33.8%	33.6%	20 bps
Manufacturing Non-GAAP OM%	35.5%	35.6%	(10) bps

- Stable operating margin YOY is consistent with historical Q1 trends
- Non-GAAP operating margin outlook in line with revised expectations for 2021 of mid-30% when factoring in Cognate acquisition



Manufacturing Results – Microbial Solutions

- Microbial Solutions growth rate rebounded >10% in 1Q21, reflecting strong demand for Endosafe® endotoxin testing systems, cartridges, and core reagents in all geographic regions
- Continue to work through delayed instrument installations that resulted from COVID-19 restrictions
 - Gaining access to more client sites
- Pleased with strength of underlying demand for endotoxin testing platform
 - Performs FDA-mandated, lot release testing for clients' critical quality-control testing needs
- Clients prefer our comprehensive and efficient microbial testing solutions because of quality, speed, and accuracy of testing platform



Mfg Results – Biologics Testing Solutions

- Biologics reported another exceptional quarter of strong, double-digit revenue growth
 - Principally driven by robust market demand for testing cell and gene therapies and COVID-19 therapeutics
- Believe cell and gene therapies will continue to drive significant growth for years to come
- Demand for COVID-19 vaccine testing is intensifying as therapies move on to commercial production phase, even as some early-stage testing activity subsides
- Given strength of demand, continuing to build our extensive portfolio of services to support the safe manufacture of biologics and ensure available capacity to accommodate client demand



Cognate Acquisition

- Believe Cognate will be highly complementary to Biologics business and portfolio as a whole
- Cognate establishes CRL as a premier scientific partner for cell and gene therapy development, testing, and manufacturing
- Broader services will provide clients with integrated solution from basic research through CGMP production, enabling them to outsource CGMP cell therapy production and required analytical testing to one scientific partner
 - Reduces bottlenecks and inefficiencies of utilizing multiple outsource providers
- Because CRL already provides extensive non-clinical services for cell and gene therapies, integration process is particularly focused on unlocking new business opportunities across our portfolio



Broadening Portfolio & Scientific Expertise

- Acquisition of Cognate is part of ongoing strategy to broaden our unique portfolio and scientific expertise in order to support new paradigms and therapeutic areas of research
- Biopharma clients seek to drive greater efficiency and leverage scientific benefits by working with fewer, trusted partners who have broad, integrated capabilities
- Transformed our business over the last decade to accommodate client needs through M&A, strategic partnerships, internal investment, and promoting a culture of continuous improvement in everything we do
- Built the leading safety assessment franchise in the world and established integrated, end-to-end discovery offering for both small and large molecules
- Given emerging importance of complex biologics and cell and gene therapies, adding CDMO capabilities is a logical extension for the portfolio



Importance of Growth Strategy

- Will continue to move our growth strategy forward
- Disciplined M&A and strategic partnerships remain vital components, as we endeavor to further enhance scientific expertise, global reach, and innovative technologies we can offer clients across all three business segments
- Investing in scientific capabilities, as well as internally in necessary staff and resources, will help ensure that we can meet clients' needs and support robust growth in our markets
- Biotech funding has never been stronger, clients are investing more in research and development, and it is incumbent on CRL to be the scientific partner who can help move client programs forward from concept, to non-clinical development, to safe manufacture of life-saving therapeutics

1Q21 Results

(\$ in millions)	1Q21	1Q20	ΥΟΥ Δ	Organic Δ
Revenue	\$824.6	\$707.1	16.6%	13.0%
GAAP OM%	15.0%	13.3%	170 bps	
Non-GAAP OM%	20.7%	19.0%	170 bps	
GAAP EPS	\$1.20	\$1.02	17.6%	
Non-GAAP EPS	\$2.53	\$1.84	37.5%	

- Delivered strong revenue growth, well above the 10% level on an organic basis
- Encouraging operating margin performance reflects our efforts to:
 - Build a more scalable and efficient infrastructure
 - Leverage the robust growth in our end markets



Increased 2021 Guidance Including Cognate

Revenue growth, reported	19%-21%
Revenue growth, organic	12%-14%
GAAP EPS	\$5.95-\$6.20
Non-GAAP EPS	\$9.75-\$10.00

- Increased 2021 guidance reflects the enhanced growth profile, including the strong
 1Q21 performance and the addition of Cognate and other acquisitions
- Given the robust top-line performance, we expect to drive meaningful operating margin improvement this year
 - 2021 non-GAAP margin approaching 21%
- Increased non-GAAP EPS guidance represents YOY growth above 20%



Revised 2021 Segment Revenue Outlook

	2021 Reported Revenue Growth	2021 Organic Revenue Growth
RMS	Low-20% range	High teens
DSA	Mid-teens	Low-double digits
Manufacturing	High-30% range	Mid-teens
Consolidated CRL	19%-21%	12%-14%

- Outlook for 2021 continues to reflect the strong business environment and the differentiated capabilities we provide to support our clients' needs
 - RMS segment reflects COVID-19 recovery, exceptional growth in China, and expectation that cell supply revenue growth improves during the year
 - DSA segment reflects strong 1Q21 performance and intensified early-stage research activity
 - MFG segment reflects both Biologics and Microbial contributions



Revised 2021 Segment Operating Margin Outlook

- RMS segment will continue to be a primary contributor to CRL's overall margin improvement in 2021
 - RMS non-GAAP operating margin meaningfully above 25% this year
- DSA non-GAAP operating margin is expected to increase over the prior year, into the mid-20% range
- Factoring in Cognate's margin, MFG non-GAAP operating margin is expected to be in the mid-30% range, or moderately below its 2020 level



Unallocated Corporate Expenses

(\$ in millions)	1Q21	4Q20	1Q20
GAAP	\$61.6	\$45.7	\$46.5
Non-GAAP	\$51.2	\$41.4	\$39.8

- Non-GAAP unallocated corporate costs were slightly higher than our expectations
 - 6.2% of total revenue in 1Q21, compared to 5.6% of total revenue in 1Q20
- 1Q21 increase was primarily the result of:
 - Continued investments to support the growth of our businesses
 - Higher performance-based compensation costs, due in part to the 1Q21 operating outperformance
- Despite higher 1Q21 expenses, continue to expect non-GAAP unallocated corporate costs to be in the mid-5% range as percent of revenue for 2021
- GAAP unallocated corporate costs expected to be ~6% of total revenue in 2021



Tax Rate

	1Q21	4Q20	1Q20
GAAP	3.6%	16.4%	8.3%
Non-GAAP	14.5%	17.8%	14.3%

- 1Q21 non-GAAP tax rate was 20 bps higher YOY and consistent with our outlook in February
 - Prior outlook (February): Mid-teens tax rate due to the gaiting of the excess tax benefit from stock-based compensation
- Continue to expect full-year 2021 tax rate will be in the low-20% range (GAAP and non-GAAP)
 - Unchanged from our outlook provided in February



Net Interest Expense

(\$ in millions)	1Q21	4Q20	1Q20
GAAP interest expense, net	\$29.7	\$33.1	\$14.8
Non-GAAP interest expense, net	\$3.7	\$33.1	\$14.8
Adjustments for foreign exchange forward contract and related interest expense ⁽¹⁾	<u>\$13.4</u>	<u>(\$16.1)</u>	<u>\$4.2</u>
Adjusted net interest expense	\$17.1	\$17.0	\$19.0

- Total adjusted net interest expense for 1Q21 was essentially flat sequentially and a decrease of nearly \$2M YOY
 - Lower average debt levels resulted in interest rate savings based on leverage ratio
- At the end of 1Q21, total outstanding debt was \$2.2B, representing a gross leverage ratio of 2.3x and a net leverage ratio of 1.9x



Capital Structure

- In March, issued \$1B senior notes to further optimize capital structure and take advantage of attractive interest rate environment
- Proceeds of the bond offering were used to:
 - Redeem a previously issued, higher-rate \$500M bond
 - Pay down the existing term loan and a portion of the revolving credit facility
 - Finance a portion of the Cognate acquisition
- In April, amended existing credit agreement to establish a new revolver with borrowing capacity of up to \$3B
- Refinancing activities reduced average interest rate on debt by ~50 bps to 2.65%



Capital Structure (con't.)

- On a pro forma basis, including Cognate and Retrogenix, gross leverage ratio was just under 3x and total outstanding debt was ~\$3B
- For 2021, total adjusted net interest expense expected in the range of \$83-\$86M
 - Higher debt balances due primarily to Cognate, partially offset by the lower average interest rate from refinancing activities
- For 2021, GAAP interest expense expected to be in a range of \$96-\$99M

CRL Capitalization (\$ in MM)	4/24/2021
4.25% Senior notes due 2028	\$500
3.75% Senior notes due 2029	\$500
4.00% Senior notes due 2031	\$500
Revolving credit facility	\$1,452
Finance leases & other	\$11
Total debt (short & long-term)	\$2,963
Additional borrowing capacity	\$1,522



Cash Flow

(\$ in millions)	1Q21	1Q20	2021 Outlook
Free cash flow (FCF)	\$142.2	\$42.9	~\$435
Capital expenditures	\$28.0	\$25.7	~\$220
Depreciation	\$32.7	\$29.4	\$145-\$150
Amortization	\$28.8	\$27.9	\$145-\$160

- Significant increase in FCF from last year was due to the strong 1Q21 operating performance, along with continued focus on working capital management
- Increasing capex guidance for 2021 by \$40M to ~\$220M, reflecting investments in Cognate to support this high-growth business
 - Expect capex to remain below 7% of total revenue this year, which is consistent with 2019 Investor Day target
- Updating 2021 FCF guidance to ~\$435M, the upper end of prior guidance range
 - Increase due to strong 1Q21 performance, even after incorporating Cognate transaction costs and capital needs

2021 Revised Guidance Summary (Including Cognate)

	GAAP	Non-GAAP
Revenue growth	19%-21% reported	12%-14% organic ⁽¹⁾
Operating margin	Comparable to 2020 level	Approaching 21%
Unallocated corporate	~6% as a % of revenue	Mid-5% range as a % of revenue
Net interest expense (total)	\$96M-\$99M	\$83M-\$86M
Tax rate	Low-20% range	Low-20% range
EPS	\$5.95-\$6.20	\$9.75-\$10.00
Cash flow	Operating cash flow ~\$655M	Free cash flow ~\$435M
Capital expenditures	~\$220M	~\$220M



2Q21 Outlook (Including Cognate)

	2Q21 Outlook
Reported revenue growth YOY	At or near 30% including Cognate
Organic revenue growth YOY	At or near 20%
Non-GAAP EPS growth YOY	More than 50% vs. 2Q20

- 2Q21 updated outlook reflects a continuation of strong demand environment
- 2Q21 revenue growth reflects the YOY comparison to last year's COVIDrelated revenue impact, which will contribute ~700 bps in 2Q21
- As a result of COVID-19 impact on 2Q20, expect 2Q21 non-GAAP operating margin and EPS to increase significantly vs. the prior year



Concluding Remarks

- Very pleased with our strong 1Q21 performance, which included robust revenue, earnings and free cash flow growth
- Remain confident about our prospects for the year, and our ability to consistently grow the top-line, bottom-line, and cash flow generation
 - Reflected in the substantial improvement in our 2021 outlook
- Hosting a virtual Investor Day on Thursday, May 27th via ir.criver.com
 - Plan to update our longer-term financial targets, which we believe will reflect the strong demand environment



1Q21 Regulation G Financial Reconciliations



CHARLES RIVER LABORATORIES INTERNATIONAL, INC.

RECONCILIATION OF GAAP TO NON-GAAP SELECTED BUSINESS SEGMENT INFORMATION (UNAUDITED)⁽¹⁾

(in thousands, except percentages)

	Three Months Ended								
	Mar	ch 27, 2021	Mai	rch 28, 2020					
Research Models and Services									
Revenue	\$	176,910	\$	145,996					
Operating income		44,935		27,373					
Operating income as a % of revenue		25.4 %		18.7 %					
Add back:									
Amortization related to acquisitions		5,339		5,652					
Severance		7		(9)					
Acquisition related adjustments (2)		456		285					
Site consolidation costs, impairments and other items		_		229					
Total non-GAAP adjustments to operating income	\$	5,802	\$	6,157					
Operating income, excluding non-GAAP adjustments	\$	50,737	\$	33,530					
Non-GAAP operating income as a % of revenue		28.7 %		23.0 %					
Depreciation and amortization	\$	9,679	\$	8,752					
Capital expenditures	\$	2,983	\$	5,412					
Discovery and Safety Assessment									
Revenue	\$	501,178	\$	438,683					
Operating income		90,949		72,283					
Operating income as a % of revenue		18.1 %		16.5 %					
Add back:									
Amortization related to acquisitions		22,648		23,007					
Severance		412		83					
Acquisition related adjustments (2)		5,270		1,289					
Site consolidation costs, impairments and other items		147		-					
Total non-GAAP adjustments to operating income	\$	28,477	\$	24,379					
Operating income, excluding non-GAAP adjustments	\$	119,426	\$	96,662					
Non-GAAP operating income as a % of revenue		23.8 %		22.0 %					
Depreciation and amortization	\$	44,608	\$	41,330					
Capital expenditures	\$	17,040	\$	14,729					
Manufacturing Support									
Revenue	\$	146,478	\$	122,380					
Operating income		49,437		41,112					
Operating income as a % of revenue		33.8 %		33.6 %					
Add back:									
Amortization related to acquisitions		2,214		2,247					
Severance		294		256					
Acquisition related adjustments (2)		42		2					
Site consolidation costs, impairments and other items		40		-					
Total non-GAAP adjustments to operating income	\$	2,590	\$	2,505					
Operating income, excluding non-GAAP adjustments	\$	52,027	\$	43,617					
Non-GAAP operating income as a % of revenue		35.5 %		35.6 %					
Depreciation and amortization	\$	6,569	\$	6,366					
Capital expenditures	\$	7,110	\$	5,161					



CHARLES RIVER LABORATORIES INTERNATIONAL, INC. RECONCILIATION OF GAAP TO NON-GAAP SELECTED BUSINESS SEGMENT INFORMATION (UNAUDITED)⁽¹⁾

(in thousands, except percentages)

		Three Mo	nths End	ed
	Mar	rch 27, 2021	Ma	rch 28, 2020
CONTINUED FROM PREVIOUS SLIDE				
Unallocated Corporate Overhead	\$	(61,618)	\$	(46,487)
Add back:				
Severance		(151)		-
Acquisition related adjustments (2)		10,560		6,983
Other items (3)				(287)
Total non-GAAP adjustments to operating expense	\$	10,409	\$	6,696
Unallocated corporate overhead, excluding non-GAAP adjustments	\$	(51,209)	\$	(39,791)
Total				
Revenue	\$	824,566	\$	707,059
Operating income		123,703		94,281
Operating income as a % of revenue		15.0 %		13.3 %
Add back:				
Amortization related to acquisitions		30,201		30,906
Severance		562		330
Acquisition related adjustments (2)		16,328		8,559
Site consolidation costs, impairments and other items (3)		187		(58)
Total non-GAAP adjustments to operating income	\$	47,278	\$	39,737
Operating income, excluding non-GAAP adjustments	\$	170,981	\$	134,018
Non-GAAP operating income as a % of revenue		20.7 %		19.0 %
Depreciation and amortization	\$	61,508	\$	57,260
Capital expenditures	\$	28,030	\$	25,721

- (1) Charles River management believes that supplementary non-GAAP financial measures provide useful information to allow investors to gain a meaningful understanding of our core operating results and future prospects, without the effect of often-one-time charges and other items which are outside our normal operations, consistent with the manner in which management measures and forecasts the Company's performance. The supplementary non-GAAP financial measures included are not meant to be considered superior to, or a substitute for results of operations prepared in accordance with U.S. GAAP. The Company intends to continue to assess the potential value of reporting non-GAAP results consistent with applicable rules, regulations and guidance.
- (2) These adjustments are related to the evaluation and integration of acquisitions, which primarily include transaction, third-party integration, and certain compensation costs, and fair value adjustments associated with contingent consideration.
- (3) Other items relate to third-party costs, net of insurance reimbursements, incurred during the three months ended March 28, 2020 associated with the remediation of the unauthorized access into the Company's information systems which was detected in March 2019.



CHARLES RIVER LABORATORIES INTERNATIONAL, INC.

RECONCILIATION OF GAAP EARNINGS TO NON-GAAP EARNINGS (UNAUDITED)⁽¹⁾

(in thousands, except per share data)

	Three Months Ended						
	Mai	rch 27, 2021	Ma	rch 28, 2020			
Net income attributable to common shareholders	\$	61,530	\$	50,769			
Add back:							
Non-GAAP adjustments to operating income (Refer to previous schedule)		47,278		39,737			
Write-off of deferred financing costs and fees related to debt financing		25,979		-			
Venture capital and strategic equity investment losses, net		16,719		12,035			
Other (2)		(2,370)		-			
Tax effect of non-GAAP adjustments:							
Non-cash tax provision related to international financing structure (3)		1,035		1,073			
Tax effect of the remaining non-GAAP adjustments		(21,013)		(11,804)			
Net income attributable to common shareholders, excluding non-GAAP adjustments	\$	129,158	\$	91,810			
Weighted average shares outstanding - Basic		49,980		49,189			
Effect of dilutive securities:							
Stock options, restricted stock units and performance share units		1,095		777			
Weighted average shares outstanding - Diluted		51,075		49,966			
Earnings per share attributable to common shareholders:							
Basic	\$	1.23	\$	1.03			
Diluted	\$	1.20	\$	1.02			
Basic, excluding non-GAAP adjustments	\$	2.58	\$	1.87			
Diluted, excluding non-GAAP adjustments	\$	2.53	\$	1.84			

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- (2) This adjustment relates to the gain on an immaterial divestiture which occurred in the three months ended March 27, 2021.
- (3) This adjustment relates to the recognition of deferred tax assets expected to be utilized as a result of changes to the Company's international financing structure.



CHARLES RIVER LABORATORIES INTERNATIONAL, INC. RECONCILIATION OF GAAP REVENUE GROWTH TO NON-GAAP REVENUE GROWTH, ORGANIC (UNAUDITED) (1)

Three Months Ended March 27, 2021	Total CRL	RMS Segment	DSA Segment	MS Segment
Revenue growth, reported	16.6 %	21.2 %	14.2 %	19.7 %
Increase due to foreign exchange	(2.9)%	(4.2)%	(2.3)%	(4.1)%
Contribution from acquisitions (2)	(0.7)%	(2.2)%	(0.3)%	- %
Non-GAAP revenue growth, organic (3)	13.0 %	14.8 %	11.6 %	15.6 %

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- (2) The contribution from acquisitions reflects only completed acquisitions.
- (3) Organic revenue growth is defined as reported revenue growth adjusted for acquisitions and foreign exchange.



CHARLES RIVER LABORATORIES INTERNATIONAL, INC. RECONCILIATION OF GAAP TO NON-GAAP REVENUE AND EARNINGS PER SHARE (EPS)

Guidance for the Twelve Months Ended December 25, 2021E

2021 GUIDANCE INCLUDING COGNATE	CURRENT	PRIOR
Revenue growth, reported	19% – 21%	16% – 18%
Less: Contribution from acquisitions (1)	(4.5%) - (5.0%)	(4.5%) - (5.0%)
Unfavorable/(favorable) impact of foreign exchange	~(2.5%)	(2.0%) - (2.5%)
Revenue growth, organic (2)	12% – 14%	9% – 11%
GAAP EPS estimate	\$5.95 – \$6.20	_
Acquisition-related amortization (3)	\$2.15 - \$2.40	_
Acquisition-related adjustments (4)	\$0.75 - \$0.80	_
Other items ⁽⁵⁾	~\$0.55	_
Venture capital and other strategic investment losses/(gains), net (6)	\$0.25	_
Non-GAAP EPS estimate	\$9.75 – \$10.00	\$9.00 - \$9.25
Free cash flow ⁽⁷⁾	~\$435 million	_

Footnotes to Guidance Table:

- (1) The contribution from acquisitions reflects only those acquisitions that have been completed.
- (2) Organic revenue growth is defined as reported revenue growth adjusted for acquisitions and foreign currency translation.
- (3) Acquisition-related amortization includes an estimate of \$0.45-\$0.65 for the impact of the Cognate acquisition and \$0.05-\$0.10 for other acquisitions completed in 2021 because the preliminary purchase price allocation has not been completed.
- (4) These adjustments are related to the evaluation and integration of acquisitions, and primarily include transaction, advisory, and certain third-party integration costs, as well as certain costs associated with acquisition-related efficiency initiatives.
- (5) These items primarily relate to charges of a) approximately \$0.15 associated with U.S. and international tax legislation, and b) approximately \$0.40 associated with debt extinguishment costs and the write-off of deferred financing costs related to debt refinancing.
- (6) Venture capital and other strategic investment performance only includes recognized gains or losses. The Company does not forecast the future performance of these investments.
- (7) Reconciliation of the current 2021 free cash flow guidance is as follows: Cash flow from operating activities of approximately \$655 million, less capital expenditures of approximately \$220 million, equates to free cash flow of approximately \$435 million.

CHARLES RIVER LABORATORIES INTERNATIONAL, INC. RECONCILIATION OF GAAP TO NON-GAAP

SELECTED BUSINESS SEGMENT INFORMATION (UNAUDITED)⁽¹⁾

(in thousands)

	Three Months Ended						
	Decen	nber 26, 2020					
Unallocated Corporate Overhead	\$	(45,747)					
Add back:							
Severance and executive transition costs		375					
Acquisition related adjustments (2)		4,020					
Total non-GAAP adjustments to operating expense	\$	4,395					
Unallocated corporate overhead, excluding non-GAAP adjustments	\$	(41,352)					

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- (2) These adjustments are related to the evaluation and integration of acquisitions, which primarily include transaction, third-party integration, and certain compensation costs, and fair value adjustments associated with contingent consideration.



CHARLES RIVER LABORATORIES INTERNATIONAL, INC. RECONCILIATION OF GAAP TAX RATE TO NON-GAAP TAX RATE (UNAUDITED) (1) (in thousands)

	Three Months Ended							
		March 27, 2021	Dece	mber 26, 2020	Ma	rch 28, 2020		
Income from operations before income taxes & noncontrolling interests \$		66,302	\$	172,427	\$	55,459		
Add back:								
Amortization related to acquisitions		30,201		28,097		30,906		
Severance		562		1,051		330		
Acquisition related adjustments (2)		16,328		5,724		8,559		
Site consolidation costs, impairments and other items (3)		187		877		(58)		
Write-off of deferred financing costs and fees related to debt financing		25,979		-		-		
Venture capital and strategic equity investment losses (gains), net		16,719		(68,635)		12,035		
Loss due to U.S. Pension termination		-		10,283		-		
Other (4)		(2,370)						
Income before income taxes & noncontrolling interests, excluding specified charges								
(Non-GAAP)	\$	153,908	\$	149,824	\$	107,231		
and the second s								
Provision for income taxes (GAAP)	\$	2,367	\$	28,237	\$	4,622		
Non-cash tax benefit related to international financing structure (5)		(1,035)		(1,454)		(1,073)		
Tax effect of the remaining non-GAAP adjustments		21,013		(87)		11,804		
Provision for income taxes (Non-GAAP)	\$	22,345	\$	26,696	\$	15,353		
Total rate (GAAP)		3.6 %		16.4 %		8.3 %		
Total rate, excluding specified charges (Non-GAAP)		14.5 %		17.8 %		14.3 %		

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⁽²⁾ These adjustments are related to the evaluation and integration of acquisitions, which primarily include transaction, third-party integration, and certain compensation costs, and fair value adjustments associated with contingent consideration.

⁽³⁾ Other items relate to third-party costs, net of insurance reimbursements, incurred during the three months ended March 28, 2020 associated with the remediation of the unauthorized access into the Company's information systems which was detected in March 2019.

⁽⁴⁾ This adjustment relates to the gain on an immaterial divestiture which occurred in the three months ended March 27, 2021.

⁽⁵⁾ This adjustment relates to the recognition of deferred tax assets expected to be utilized as a result of changes to the Company's international financing structure.

CHARLES RIVER LABORATORIES INTERNATIONAL, INC. RECONCILIATION OF GAAP TO NON-GAAP NET INTEREST EXPENSE(1)

(in thousands)

		Fiscal Year Ended			
	March 27, 2021	_	December 26, 2020	March 28, 2020	December 25, 2021E
GAAP Interest expense, net	\$ 29,684	\$	33,084	\$ 14,751	\$96,000-\$99,000
Exclude:					
Write-off of deferred financing costs and fees related to debt financing	 (25,979)		-	 =	(26,000)
Non-GAAP Interest expense, net	3,705		33,084	14,751	70,000-73,000
Adjustments for foreign exchange forward contract and related interest expense, net (2)	 13,356		(16,068)	4,213	13,000
Adjusted Interest expense, net	\$ 17,061	\$	17,016	\$ 18,964	\$83,000-\$86,000



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⁽²⁾ Amounts reported in total adjusted interest expense include a \$14.0 million gain on a forward contract and \$0.1 million of additional interest expense for the three months ended March 27, 2021; a \$15.4 million loss on a forward contract and \$0.1 million of additional interest expense for the three months ended December 26, 2020; and a \$6.1 million gain on a forward contract partially offset by \$1.4 million of additional interest expense for the three months ended March 28, 2020.

CHARLES RIVER LABORATORIES INTERNATIONAL, INC.

RECONCILIATION OF GROSS/NET LEVERAGE RATIO. INCLUDING GAAP NET INCOME TO ADJUSTED ERITDA (1)

(dollars in thousands, except for per share data)

	N	Iarch 27,	December 26, I		6, December 28,		December 29,		December 30,		December 31,		December 26,		December 27,		December 28,		December 29,	
		2021		2020	2019		2018		2017		2016		2015		2014		2013			2012
<u>DEBT (2):</u>																				
Total Debt & Finance Leases	\$	2,205,266	\$	1,979,784	\$	1,888,211	\$	1,668,014	\$	1,145,104	\$	1,235,009	\$	863,031	\$	777,863	\$	663,789	\$	666,520
Plus: Other adjustments per credit agreement	\$	33,163	\$	2,328	\$	712	\$	3,033	\$	298	\$	3,621	\$	1,370	\$	2,828	\$	9,787	\$	9,680
Less: Unrestricted Cash and Cash Equivalents up to \$150M	\$	(150,000)				_				_				_		_		_		
Total Indebtedness per credit agreement	\$	2,088,429	\$	1,982,112	\$	1,888,924	\$	1,671,047	\$	1,145,402	\$	1,238,630	\$	864,401	\$	780,691	\$	673,576	\$	676,200
Less: Cash and cash equivalents (net of \$150M above)		(315,411)		(228,424)		(238,014)		(195,442)		(163,794)		(117,626)		(117,947)		(160,023)		(155,927)		(109,685)
Net Debt	\$	1,773,018	\$	1,753,688	\$	1,650,910	\$	1,475,605	s	981,608	\$	1,121,004	\$	746,454	\$	620,668	\$	517,649	\$	566,515

	March 27, I		Dec	December 26, De 2020		cember 28, December 29, 2019 2018		December 30, 2017		December 31, 2016		December 26, 2015		December 27, 2014		December 28, 2013		December 29, 2012	
ADJUSTED EBITDA (2):																			
Net income attributable to common shareholders	\$	375,064	\$	364,304	\$	252,019	\$ 226,373	\$	123,355	\$	154,765	\$	149,313	\$	126,698	\$	102,828	\$	97,295
Adjustments:																			
Adjust: Non-cash gains/losses of VC partnerships & strategic investments		26,148		_		_	_		_		_		_		_		_		_
Less: Aggregate non-cash amount of nonrecurring gains		(1,423)		(1,361)		(310)	_		_		(685)		(9,878)		(2,048)		_		_
Plus: Interest expense		99,647		76,825		79,586	65,258		29,777		27,709		15,072		11,950		20,969		33,342
Plus: Provision for income taxes		79,553		81,808		50,023	54,996		171,369		66,835		43,391		46,685		32,142		24,894
Plus: Depreciation and amortization		239,172		234,924		198,095	161,779		131,159		126,658		94,881		96,445		96,636		81,275
Plus: Non-cash nonrecurring losses		13,783		16,810		427	559		17,716		6,792		10,427		1,615		4,202		12,283
Plus: Non-cash stock-based compensation		58,570		56,341		57,271	47,346		44,003		43,642		40,122		31,035		24,542		21,855
Plus: Permitted acquisition-related costs		26,183		18,750		34,827	19,181		6,687		22,653		13,451		6,285		1,752		3,676
Plus: Pro forma EBITDA adjustments for permitted acquisitions		5,420		8		12,320	15,648		690		18,573		9,199		10,787				253
Adjusted EBITDA (per the calculation defined in compliance certificates)	\$	922,117	S	848,408	\$	684,259	\$ 591,140	\$	524,756	\$	466,942	\$	365,978	\$	329,452	\$	283,071	\$	274,873

	March 27,	December 26,	December 28,	December 29,	December 30,	December 31,	December 26,	December 27,	December 28,	December 29,
	2021	2020	2019	2018	2017	2016	2015	2014	2013	2012
LEVERAGE RATIO:										
Gross leverage ratio per credit agreement (total debt divided by adjusted										
EBITDA)	2.26x	2.34x	2.76x	2.83x	2.2x	2.7x	2.4x	2.4x	2.4x	2.5x
Net leverage ratio (net debt divided by adjusted EBITDA)	1.9x	2.1x	2.4x	2.5x	1.9x	2.4x	2.0x	1.9x	1.8x	2.1x
	March 27, 2021	December 26, 2020								
INTEREST COVERAGE RATIO:										
Capital Expenditures	166,578	166,560	_	_	_	_	_	_	_	_
Cash Interest Expense Interest Coverage ratio per the credit agreement (Adjusted EBITDA minus	99,814	77,145	_	_	_	_	_	_	_	_
Capital Expenditures divided by cash interest expense)	7.57x	8.84x	_	_	_	_	_	_	_	_

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(2) Pursuant to the definition in its credit agreement dated April 21.2021, the Company has defined its pro forma leverage ratio as total debt divided by adjusted EBITDA for the trailing-twelve-month period. The Company has defined interest coverage ratio as adjusted EBITDA for the trailing-twelve-month period less the aggregate amount of capital expenditures for the trailing-twelve-period: divided by the consolidated interest expense for the period of four consecutive fiscal quarters.

Total Debt represents third-party debt and financial lease obligitations minus up to \$150M of unrestricted cash and cash equivalents. Adjusted EBITDA represents net income, prepared in accordance with accounting principles generally accepted in the U.S. (GAAP), adjusted for interest, taxes, depreciation and amortization, and certain items that management believes are not reflective of the operational performance of the business. These adjustments include, but are not limited to, non-cash gains/loss on venture capital portfolios and strategic partnerships, acquisition-related expenses including transaction and advisory costs; asset impairments; changes in fair value of contingent consideration obligations; employee stock compensation; historical EBITDA of companies acquired during the period; and other items items items (and other items item).

Total Debt and EBITDA have not been restated for periods prior to Q1-2021.



CHARLES RIVER LABORATORIES INTERNATIONAL, INC. RECONCILIATION OF FREE CASH FLOW (NON-GAAP) $^{(1)}$

(in thousands)

		Three Mo	Fiscal Year Ended				
	Ma	rch 27, 2021	N	Iarch 28, 2020	December 25, 2021E		
Net cash provided by operating activities	\$	170,229	\$	68,590	~\$655,000		
Less: Capital expenditures		(28,030)		(25,721)	(~220,000)		
Free cash flow	\$	142,199	\$	42,869	~\$435,000		

⁽¹⁾ Charles River management believes that supplementary non-GAAP financial measures provide useful information to allow investors to gain a meaningful understanding of our core operating results and future prospects, without the effect of often-one-time charges and other items which are outside our normal operations, consistent with the manner in which management measures and forecasts the Company's performance. The supplementary non-GAAP financial measures included are not meant to be considered superior to, or a substitute for results of operations prepared in accordance with U.S. GAAP. The Company intends to continue to assess the potential value of reporting non-GAAP results consistent with applicable rules, regulations and guidance.





