UNITED STATES SECURITIES AND EXCHANGE COMMISSION

WASHINGTON, D.C. 20549

FORM 8-K

CURRENT REPORT

PURSUANT TO SECTION 13 OR 15(d) OF THE SECURITIES EXCHANGE ACT OF 1934

March 6, 2018

(Date of earliest event reported)

LABORATORY CORPORATION OF AMERICA HOLDINGS

(Exact Name of Registrant as Specified in its Charter)

Delaware		1-11353	13-3/5/3/0		
(State or other jurisdiction of Inc	corporation)	(Commission File Number)	(I.R.S. Employer Identification No.)		
358 South Main Stre	et,				
Burlington, North Carolina		27215	336-229-1127		
(Address of principal executiv	re offices)	(Zip Code)	(Registrant's telephone number including area code)		
[] Written communication pursuant to Ru [] Soliciting material pursuant to Rule 14 [] Pre-commencement communications p	le 425 under the Securities a-12 under the Exchange A ursuant to Rule 14d-2(b) u	s Act (17 CFR 230.425)			
Item 7.01 Regulation FD	Disclosure				
Summary information of the Company date	d March 6, 2018.				
Exhibit Index Exhibit 99.1					

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

LABORATORY CORPORATION OF AMERICA HOLDINGS

Registrant

By: /s/ F. SAMUEL EBERTS III

F. Samuel Eberts III

Chief Legal Officer and Secretary

March 6, 2018







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MARCH 6, 2018 | ORLANDO, FL

FORWARD LOOKING STATEMENT AND USE OF ADJUSTED MEASURES

This presentation contains forward-looking statements including but not limited to statements with respect to estimated 2018 guidance and the related assumptions, the impact of various factors on operating and financial results, expected savings and synergies (including from the LaunchPad initiative and as a result of acquisitions), and the opportunities for future growth.

This presentation contains forward-looking statements which are subject to change based on various important factors, including without limitation, competitive actions and other unforeseen changes and general uncertainties in the marketplace, changes in government regulations, including health care reform, customer purchasing decisions, including changes in payer regulations or policies, other adverse actions of governmental and third-party payers, changes in testing guidelines or recommendations, adverse results in material litigation matters, the impact of changes in tax laws and regulations, failure to maintain or develop customer relationships, our ability to develop or acquire new products and adapt to technological changes, failures in information technology systems or data security, challenges in implementing business process changes, employee relations, and the effect of exchange rate fluctuations on international operations.

Actual results could differ materially from those suggested by these forward-looking statements. The Company has no obligation to provide any updates to these forward-looking statements even if its expectations change. Further information on potential factors, risks and uncertainties that could affect the operating and financial results of Laboratory Corporation of America Holdings (the "Company") is included in the Company's Form 10-K for the year ended December 31, 2017, and Forms 10-Q, including in each case under the heading risk factors, and in the Company's other filings with the SEC.

This presentation contains "adjusted" financial information that has not been prepared in accordance with GAAP, including Adjusted EPS, and Free Cash Flow, and certain segment information. The Company believes these adjusted measures are useful to investors as a supplement to, but not as a substitute for, GAAP measures, in evaluating the Company's operational performance. The Company further believes that the use of these non-GAAP financial measures provides an additional tool for investors in evaluating operating results and trends, and growth and shareholder returns, as well as in comparing the Company's financial results with the financial results of other companies. However, the Company notes that these adjusted measures may be different from and not directly comparable to the measures presented by other companies. Reconciliations of these non-GAAP measures to the most comparable GAAP measures are included in this presentation.

LabCorp

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AGENDA



WHO WE ARE

LabCorp is a leading global life sciences company

that is deeply integrated in guiding patient care

Our Mission

is to

improve health and improve lives

Our Strategic Objectives are to:

Deliver World-Class Diagnostics

Bring Innovative Medicines to Patients Faster

Use Technology to Improve the Delivery of Care

LabCorp

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LABCORP OVERVIEW

LabCorp Diagnostics

- \$6.9B revenue in 2017 (1)
- Patient database reaching ~50% of U.S. population
- Proprietary data sets with >30 billion lab test results across a growing menu of nearly 5,000 assays
- Broad physician, health system and managed care relationships
- Consumer engagement through ~1,900 PSC/retail locations,
 5,000+ in-office phlebotomists
- Proprietary decision-support and reporting tools



- \$10.4B revenue in 2017 (1)
- Global footprint with business in 127 countries; 60,000 employees
- Unmatched real-world data and patient intelligence
- Deep scientific and therapeutic experience
- Leader in Companion Diagnostics (CDx)
- Innovative technology-enabled solutions for customers

Covance Drug Development

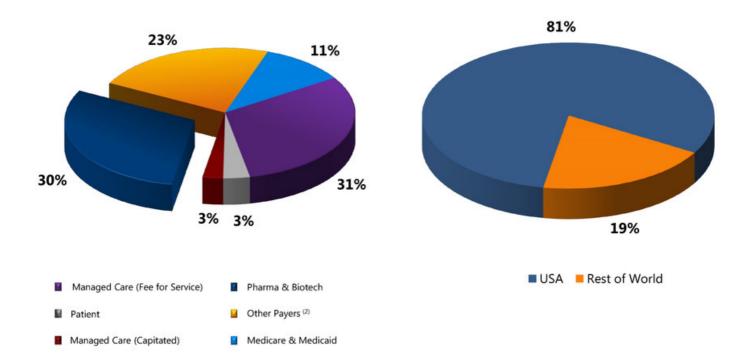
- \$3.6B revenue in 2017 (1)
- · Serving the top 20 biopharma
- Serving high-growth emerging and mid-market segments through Chiltern
- Working on ~50% of clinical trials
- >175,000 unique investigators
- Involved in all top 50 bestselling drugs on the market
- Supported 70% of all CDx on the market today
- Robust technology suite for trial planning and execution

Includes the estimated impact from adoption of the new revenue recognition accounting standard (ASC 606).
 See Appendix for details of the preliminary reconciliation of 2017 results.



EXPANDED REVENUE BASE

Attractive Customer Mix and Geographic Presence(1)



Based on full year 2017 results, which include results from Chiltern as of September 1, 2017. Does not tie due to rounding.
 Includes physicians and hospitals, occupational testing services, non-U.S. clinical diagnostic laboratory operations, nutritional chemistry and food safety operations, and Beacon LBS.



AGENDA



OUR 2018 PRIORITIES

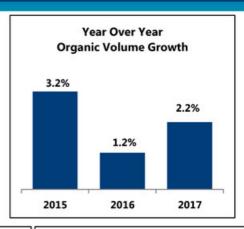




DRIVING PROFITABLE GROWTH

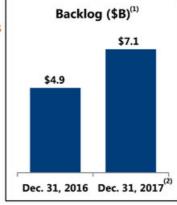
Diagnostics: Capitalize on Growth Opportunities

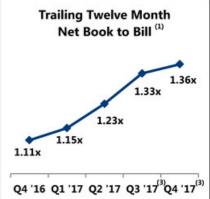
- Health systems, large physician groups and managed care partnerships
- · Expand 23andMe collaboration
- Mitigate pricing impact of PAMA
- Women's health, genetics and medical drug monitoring portfolio and capabilities
- · Pursue accretive acquisitions
- Continue focus on quality, service and innovation



Drug Development: Build on Existing Momentum to Exceed Historical Growth Rates

- Convert backlog into profitable revenue growth
- · Maintain broad-based strength in net orders
- Capitalize on strategic investments in leadership, sales force and technology





⁽¹⁾ Beginning with the fourth quarter of 2016, the Company began reporting net orders, net book-to-bill and backlog based upon fully-executed contracted awards. The Company believes this methodology is a more conservative and objective practice, providing greater visibility into its revenue conversion from the backlog. Results shown include the impact from cancellations and foreign currency translation.

Includes backlog from the acquisition of Chiltern.

(3) Includes results from Chiltern following the closing of the acquisition on September 1, 2017.



INTEGRATING KEY ACQUISITIONS

Maximize Value Through Flawless Integration

- Generates approximately \$500 million in profitable revenue growth in 2018 from Chiltern, PAML, and Mount Sinai transactions
- Successful "Best of the Best" approach to selecting and retaining talent
- Dedicated and experienced integration teams, focused on customer retention and synergies







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OPTIMIZING ENTERPRISE MARGINS

Continue Value Creation Through the LaunchPad Business Process Improvement Initiative

Covance LaunchPad

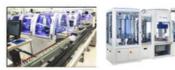
- Applying LaunchPad principles to Drug Development
- Rightsizing implemented in mid-2017 generating \$20 million in savings
- Expect \$130 million in additional net savings over the three-year period ending in 2020
- Aided by Chiltern capabilities and expertise

Diagnostics LaunchPad

- Ongoing benefit from reengineering projects
- Additional opportunities, including streamlining delivery of services

Opportunities for Productivity Gains

Automation



Global Service Delivery Model



Procurement



New Tools and Technology





AGENDA





HEALTHCARE IS UNDERGOING A PERIOD OF UNPRECEDENTED CHANGE

Transition to Value-Based Care

Enhance Drug Development Process

Role of the Consumer

- Improve efficiency in care delivery
- Reduce the overall cost of patient care
- Utilize advanced tools and analytics to deliver better outcomes via personalized medicine and population health
- Address increased trial complexity, and competition for patients and investigators
- Greater need for scalable tools and processes to initiate and manage trials
- Increased sponsor demand for data-driven study design and execution, as well as access to relevant analytes, biomarkers and tests
- Increased interest in and influence over healthcare decision-making
- Technology advances driving expectation of convenience
- Consumer satisfaction increasingly important to other healthcare stakeholders



Value-Based Care Solutions



Leading Laboratory Services

- · National access
- · Comprehensive test menu
- · Sales and service organization
- · Scientific innovation
- · Power of scale



Clinical Decision Support



- · Programs on key disease states
- · Lab reports support care guidelines
- · Developed by physicians
- Data monitoring drives cost-effective care management





Payer and Provider Collaboration

- Help stakeholders achieve total cost of care metrics in value-based care contracts
- · Actionable lab results
- · Global patient results data
- · MACRA, HEDIS, and ACO quality metrics
- · Care Intelligence® population health



Drug Development Solutions



- · Companion diagnostics leadership
- Potential provider revenue stream from increased participation in clinical trials
- · Cost savings to patients and payers
- "Real World" data



Maximizing Drug Development Tools and Technologies for Innovative Solutions Focused on Client Needs



PharmAcuity

Leverages proprietary and public data providing insights and guiding decisions; optimizes trial planning.

Includes metrics benchmarking, trial forecasting, and protocol optimization.



Patient Recruitment

Providing insights into site selection, patient recruitment and resource allocation.

Using custom
analytics to
leverage the power
of Covance's
unparalleled
patient and
investigator
databases.



Endpoint

Best-in-class interactive response system

Continuous innovation and investment including new high-value physical sample management solutions.



Xcellerate

Most modern, end-to-end clinical trial solution. Decreases risk, increases patient safety and data quality.

Includes advanced clinical data management, risk-based monitoring, and dashboards.



Global Specimen Solutions

Reduces the time, cost and risk of specimen based research.

Increases value of existing assets with advanced analytics, visualizations, and first-to-market patient consent mapping.



Patient Intelligence

Voice of patient insights from industry leading patient panel.

Understand patient view of trial participation, enhance protocol design and recruiting tactics.

Creates patientcentric development approach.

Contributed to \$1B+ of Revenue Across Clinical Development in 2017



Power of the LabCorp Data for Trial Design, Site Selection, and Patient Recruitment



Real World Data

- Not biased and represents people as they live with their disease
- Patient data is granular and identifiable



Vast Test Menu

- 30+ billion test results across thousands of diagnostic assays
- >2.5 million samples collected (>30% by LabCorp phlebotomists) and processed weekly across many diseases and therapeutic areas

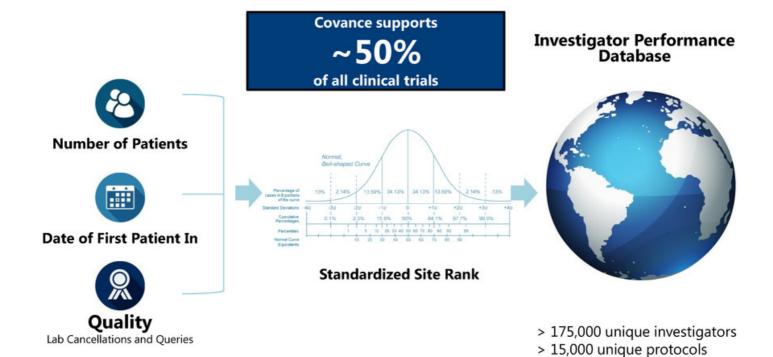


Population Level Disease Analysis

- Surveillance of disease spread to enable just in time recruitment
- Unlike other types of real world data, lab data can be easily accessed near real time



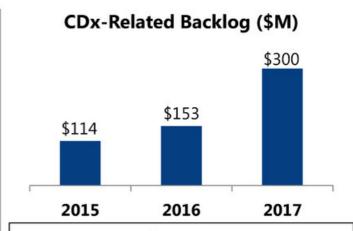
Power of the Xcellerate Investigator Database





Increasing Our Leadership in Companion and Complementary Diagnostics (CDx)

- Dedicated CDx organization with capabilities across development, validation, testing, regulatory support, commercialization and market access
- Opened purpose built, state of the art CDx laboratory, with focus on genomics and molecular pathology
- Supported ~70% of CDx on the market⁽¹⁾
- Customizable offering -- in vitro diagnostic (IVD) partnerships and single site pathway
- Collaborated with over 40 clients on more than 165 CDx projects in 2017
- ~\$135M in CDx-related enterprise revenue in 2017; 3-year CAGR of ~20%



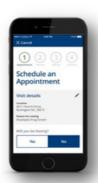
On pace to significantly exceed \$100 million

in cumulative new CDx-related revenue from the acquisition of Covance through 2018



Developing a Broad Consumer Platform to Create Deep Relationships

- Organizing around the empowered consumer
- Creating a convenient, seamless experience
- Providing easy access to lab test results and personalized content
- Offering price transparency to highlight access to highest quality, low-cost diagnostics















Bringing our High-Quality Offering to Consumers



- Patient Service Centers opened inside Walgreens stores
- · Strong patient volume, net promoter scores, and patient feedback
- Start to roll out to new markets in 2018



 Launch new at-home, self-collection offering built on LabCorp's 10 years of experience with dried blood spot testing



 Invest in expanded capacity and enhanced automation to support 23andMe strategic collaboration



 Collaborate on new delivery models, such as telehealth and on-demand phlebotomy



OUR DIFFERENTIATED SOLUTIONS ARE RESONATING WITH CUSTOMERS

Value-Based Care Solutions Streamlining Clinical Studies

Consumer Platform

Reference Laboratory Testing United the Laboratory Testing United the Laboratory Testing United the Laboratory Management

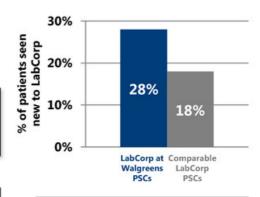
PAML and its Joint Venture Interests

Mount Sinai Health System

Novant Health

Cumulative new orders won through the combination of LabCorp patient data and Covance capabilities:

2016 2017 >\$200 million ~\$500 million



Completed
3 marquee transactions
in 2017

On track to deliver \$150 million n cumulative new reve

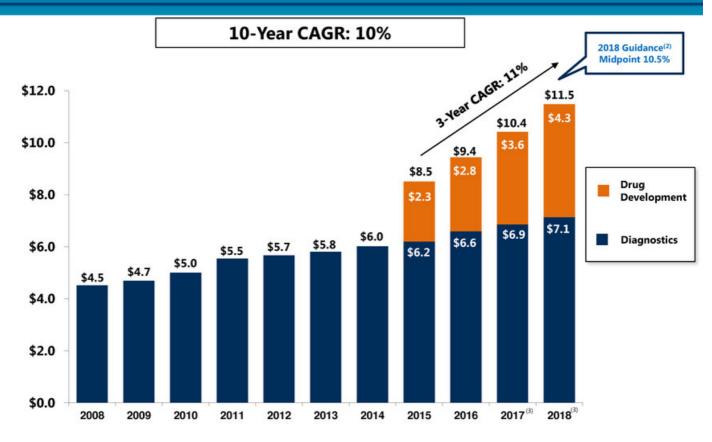
in cumulative new revenue from the acquisition of Covance through 2018 LabCorp PSCs in Walgreens stores are attracting new patients



AGENDA



REVENUE⁽¹⁾ GROWTH (DOLLARS IN BILLIONS)



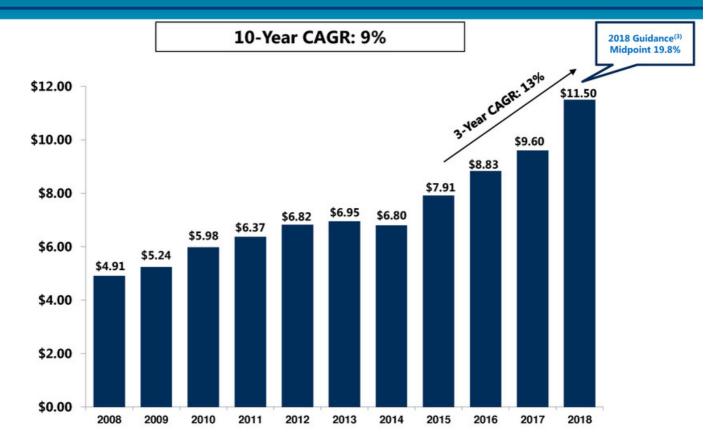
2008-2014 revenues excludes Covance results. 2008 revenue includes a \$7.5 million adjustment relating to certain historic overpayments made by Medicare for claims submitted by a subsidiary of the Company.

Guidance issued on February 6, 2018.

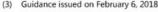
Includes the estimated impact from adoption of the new revenue recognition accounting standard (ASC 606). See Appendix for details of the preliminary reconciliation of 2017 results.



ADJUSTED EPS(1)(2) GROWTH

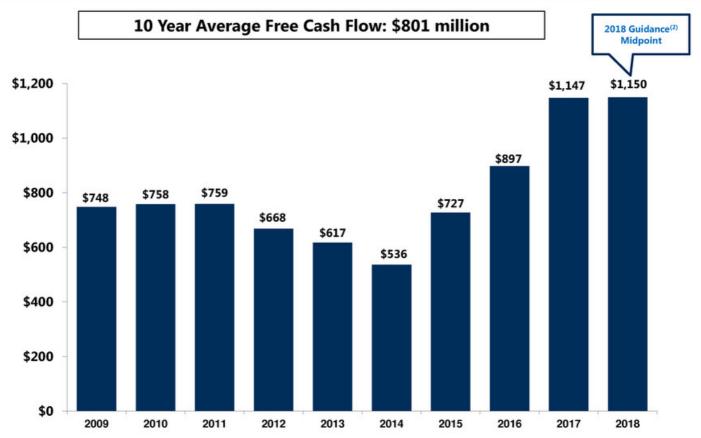


- EPS, as presented, represents adjusted, non-GAAP financial measures (excludes amortization, restructuring and other special charges).
- See Appendix for non-GAAP reconciliation. 2008-2014 figures exclude Covance results. Guidance issued on February 6, 2018.





STRONG FREE CASH FLOW⁽¹⁾ (DOLLARS IN MILLIONS)

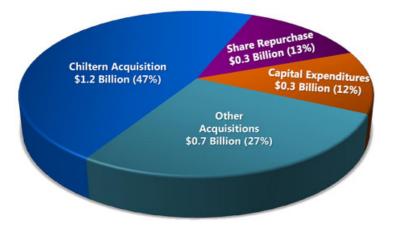


⁽¹⁾ Free Cash Flow represents Operating Cash Flow less Capital Expenditures in each of the years presented. 2009-2014 figures exclude Covance results, and other items discussed in the Appendix.
(2) Guidance issued on February 6, 2018.



TRACK RECORD OF EFFECTIVE AND BALANCED CAPITAL DEPLOYMENT TO BUILD SHAREHOLDER VALUE

Approximately \$2.5 Billion in Capital Deployment in 2017



2018 Free Cash Flow

M&A Priorities

Diagnostics "tuck-in" transactions

Return of Capital to Shareholders

· Continue share repurchases

Debt Reduction

Pay down debt to reduce leverage



OUR MARKET OPPORTUNITY AND DIFFERENTIATED CAPABILITIES POSITION US TO DELIVER UNPARALLELED VALUE

Beyond Lab

- Engage consumers through clinical study participation and care management tools
- Provide health systems, physicians, and ACOs with greater access to cutting-edge care through clinical trials
- Expand our leadership in companion diagnostics development and commercialization

Beyond CRO

- Engage consumers directly with >115 million patient encounters per year in LabCorp Diagnostics
- Apply insights from rich patient database of granular, timely, real world evidence to clinical study planning and execution
- Utilize patient service center footprint for virtual real world evidence studies

Value of the Enterprise

Actionable data for everyone

Leading proprietary IT solutions and analytics

Positioned to outperform in the healthcare system of the future









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Appendix



PRELIMINARY RECONCILIATION FOR THE NEW REVENUE RECOGNITION ACCOUNTING STANDARD (ASC 606)

Effective January 1, 2018, the Company adopted the FASB-issued converged standard on revenue recognition, using the full retrospective method. Although the Company has not completed all of the analysis required to finalize this restatement of revenues and expenses relating to this new standard, the table below presents the Company's current best estimate of the potential financial impact on its two segments' revenue, and is being provided as a reference point for the Company's guidance in 2018. The 2017 calculation is preliminary and will be finalized upon adoption in the first quarter of 2018 and the amounts are therefore subject to change. The Company does not believe the application of the new standard will have any significant impact on previously reported Adjusted Operating Income. However, the change in reported revenue will impact previously reported operating margins as shown in the table provided.

(Dollars in millions)	Decen	Months Ended aber 31, 2017 Reported	Twelve Months Ended December 31, 2017 Preliminary Restatement		
LabCorp Diagnostics (1)	72		0.00		
Net Revenue	\$	7,170.5	\$	6,858.0	
Adjusted Operating Income	\$	1,446.3	\$	1,446.3	
Adjusted Operating Margin		20.2%		21.1%	
Covance Drug Development (2)					
Net Revenue	\$	3,037.2	\$	3,562.4	
Adjusted Operating Income	\$	422.4	\$	425.7	
Adjusted Operating Margin		13.9%		11.9%	
Consolidated (1) (2)					
Net Revenue	\$	10,205.9	\$	10,418.6	
Adjusted Segment Operating Income	\$	1,868.7	\$	1,872.0	
Unallocated corporate expense	\$	(137.4)	\$	(137.4)	
Consolidated Adjusted Operating Income	\$	1,731.3	\$	1,734.6	
Adjusted Operating Margin		17.0%		16.6%	

(1)In LabCorp Diagnostics, the impact of the accounting change will reduce revenue and increase margins, as bad debt will be treated as a reduction in revenue rather than selling, general and administrative expense

(2)In Covance Drug Development, the impact of this accounting change will increase revenue and cost of revenue, resulting in lower margins due to the inclusion of investigator fees and other pass-through expenses in both categories



RECONCILIATION OF NON-GAAP FINANCIAL MEASURES(1)

(in millions, except per share data)

Twelve Months Ended

	35				Decer	nber 31,	8363			
Adjusted EPS	2008	2009	2010	2011	2012	2013	2014	2015	2016	2017
Diluted earnings per common share	\$4.16	\$4.98	\$5.29	\$5.11	\$5.99	\$6.25	\$5.91	\$4.35	\$ 7.02	\$12.21
One-time benefit from Tax Cuts and Jobs Act	-	-	-	-	100	-	-	-		(5.00)
Restructuring and special items	0.44	(0.09)	0.26	0.72	0.29	0.15	0.34	2.44	0.64	0.98
Loss on the divestiture of assets	-	-	-	0.03	-	-	-	-	-	-
Amortization expense	0.31	0.35	0.43	0.51	0.54	0.55	0.55	1.12	1.17	1.41
Adjusted EPS	\$ 4.91	\$5.24	\$ 5.98	\$ 6.37	\$ 6.82	\$ 6.95	\$ 6.80	\$ 7.91	\$ 8.83	\$ 9.60
Free Cash Flow										
Net cash provided by operating activities	\$ 781	\$ 862	\$ 884	\$ 905(2)	\$ 841	\$ 819	\$ 739	\$ 982	\$1,176	\$1,459
Less: Capital expenditures	\$(157)	\$(115)	\$(126)	\$(146)	\$(174)	\$(202)	\$(204)	\$(256)	\$ (279)	\$ (313)
Free cash flow	\$ 624	\$ 748	\$ 758	\$ 759(2)	\$ 668	\$ 617	\$ 536	\$ 727	\$ 897	\$1,147



 ²⁰⁰⁸⁻²⁰¹⁴ figures exclude Covance results.
 Operating Cash Flow and Free Cash Flow in 2011 exclude the \$49.5 million Hunter Labs settlement.