Ouanterix The Science of Precision Health

Earnings Call Q3 2019

November 6, 2019



This presentation contains "forward-looking" statements that are based on our beliefs and assumptions and on information available to us as of the date of this presentation. Forward-looking statements include all statements that are not historical facts. We may not actually achieve the plans, intentions or expectations disclosed in our forward-looking statements. Forward-looking statements involve known and unknown risks, uncertainties, assumptions and other factors that may cause our actual results, performance or achievements to be materially different from any future results, performance or achievements expressed or implied by the forward-looking statements. The risks and uncertainties that we face are described in our most recent filings with the Securities and Exchange Commission. Except as required by law, we assume no obligation to update these forward-looking statements, even if new information becomes available in the future.

To supplement the Company's financial statements presented on a GAAP basis, the Company has provided certain non-GAAP financial measures, including Non-GAAP revenue, Non-GAAP gross profit and Non-GAAP gross margin. Our non-GAAP adjustments exclude (i) the impact of \$1.3M in revenue recognized in Q3 2018 in connection with the termination of our license agreement with bioMerieux SA; and (ii) acquisition-related purchase accounting adjustments in connection with our acquisitions of Aushon Biosystems in Q1 2018 and Uman Diagnostics in Q3 2019. Management uses these non-GAAP measures to evaluate the Company's operating performance in a manner that allows for meaningful period-to-period comparison and analysis of trends in its business. Management believes that such measures are important in comparing current results with prior period results, and are useful to investors and financial analysts in assessing the Company's operating performance. The non-GAAP financial information presented here should be considered in conjunction with, and not as a substitute for the financial information presented in accordance with GAAP. Investors are encouraged to review the reconciliation of these non-GAAP measures to their most directly comparable GAAP financial measures set forth on Slide #27.

Today's Agenda



- I. Strategic and Financial Progress Kevin Hrusovsky Chairman, CEO
 - i. Q3 Highlights, Direction & 2019 Goals
 - ii. Neurology Momentum & PPH
- II. Financial Report Amol Chaubal CFO
- III. Q&A

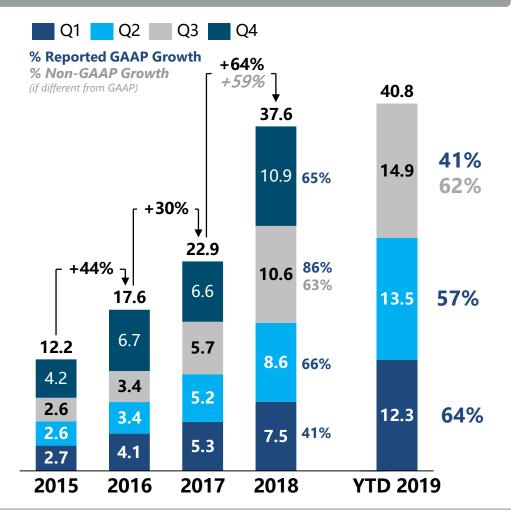
Disruptive Market, Technology and Growth



Q	3 Highlights	Q3 2019	Q3 2018
	Revenue (\$million)		
	Adjusted*	\$14.9	\$9.2
	GAAP	\$14.9	\$10.6
	Gross Margin (%)		
	Adjusted*	51.8%	46.2%
	GAAP	47.1%	52.8%

- HD-X shipped early
- Closed Uman acquisition
- Raised \$69m growth capital
- Nf-L assay highlighted in a record 50 publications at ECTRIMS (Multiple Sclerosis) conference in Stockholm
- Dr. Tatiana Plavina 18 year Biogen veteran joined QTRX
- Completed Siemens Nf-L license & Techne supply agreement

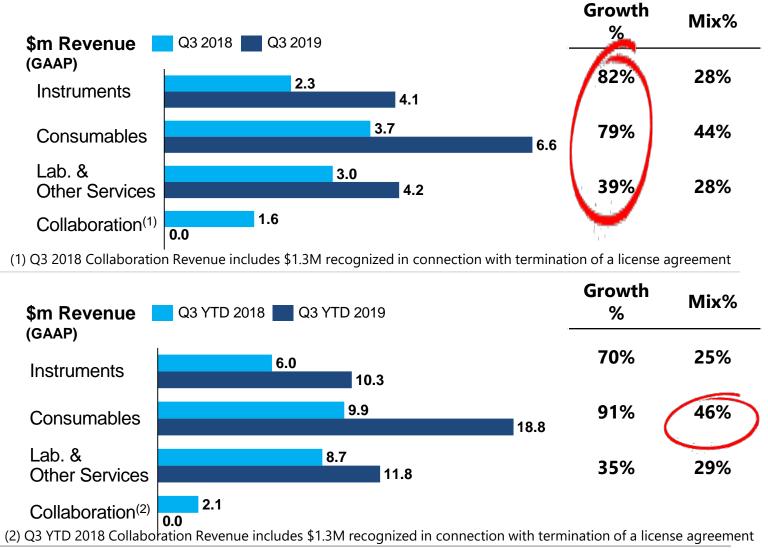
Total Quanterix Revenue Growth



Q3 2019 Growth Led by Consumables and Services



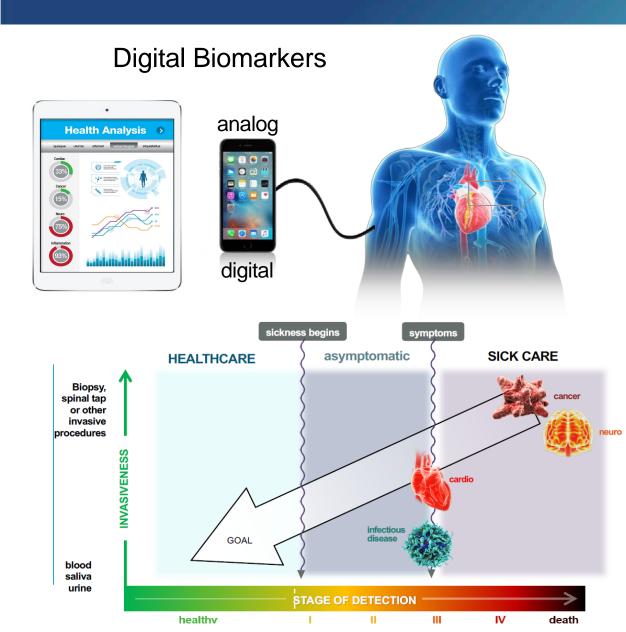
	Q3 2019	Q3 2018	YoY
Revenue (\$million)			Growth
Adjusted*	\$14.9	\$9.2	+62%
Gross Margin			
Adjusted*	51.8%	46.2%	+560 bpc
Revenue (\$million)	YTD Sept. 2019	YTD Sept. 2018	YoY Growth
Adjusted*	\$40.8	\$25.4	+61%
Gross Margin			
Adjusted*	50.7%	45.0%	+570



* Non-GAAP item. Reconciliations are included in the Appendix to this presentation and in our Q3 2019 press release.

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Convergence of Technology and Healthcare



Late; Invasive







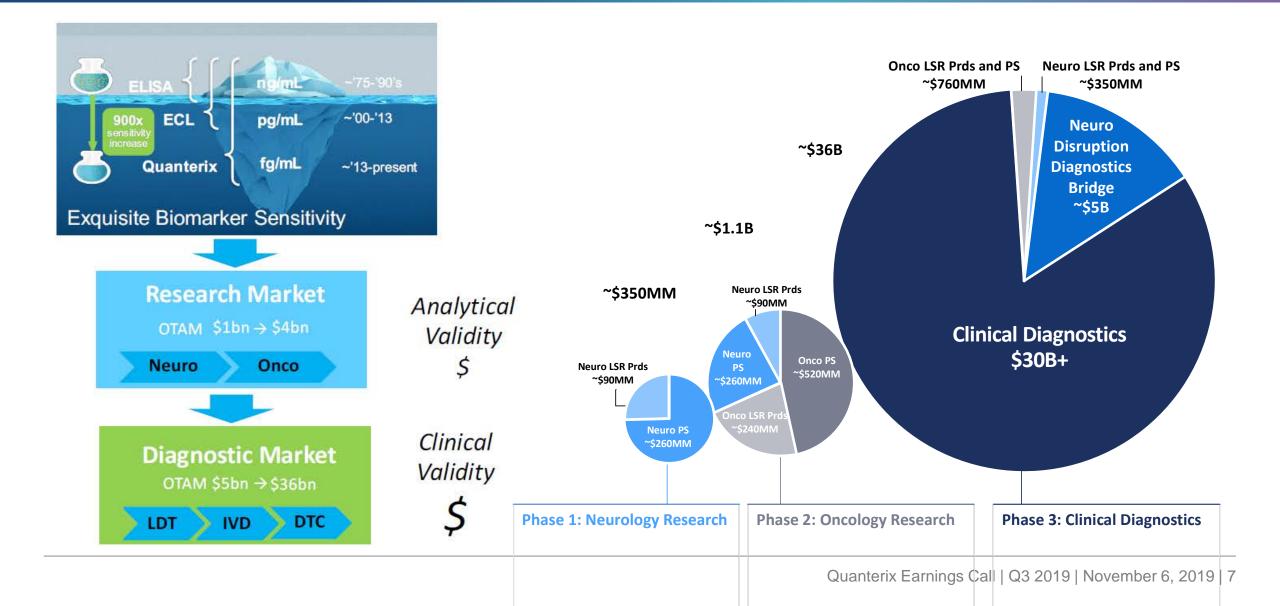






Early; Non-Invasive

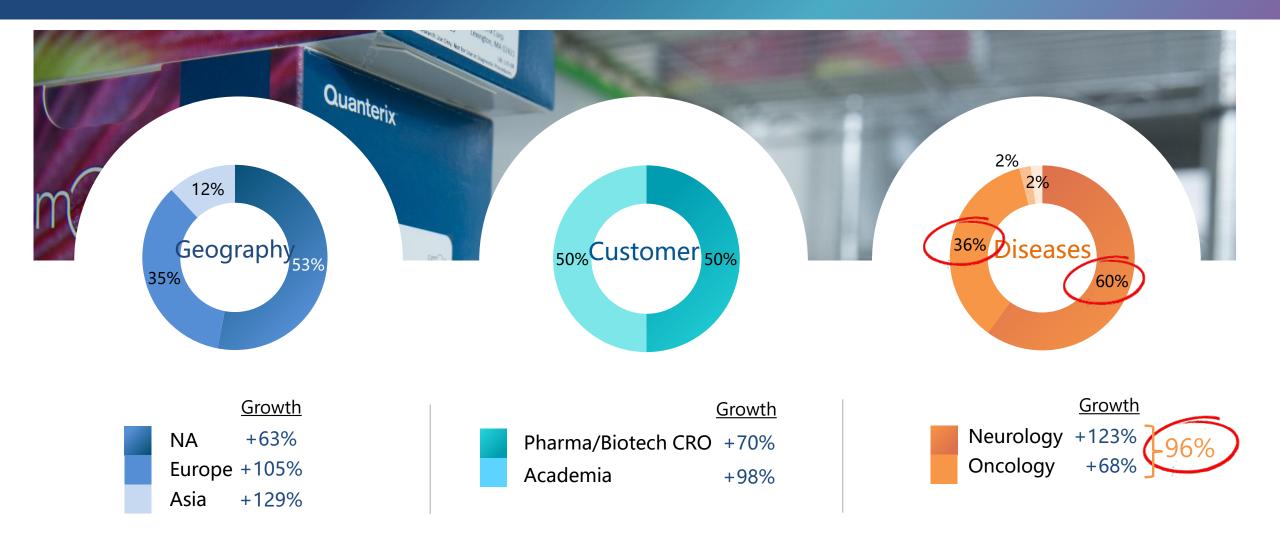
1. Neuro Beachhead 2. Penetrate Onco – Inflammation



Quanterix

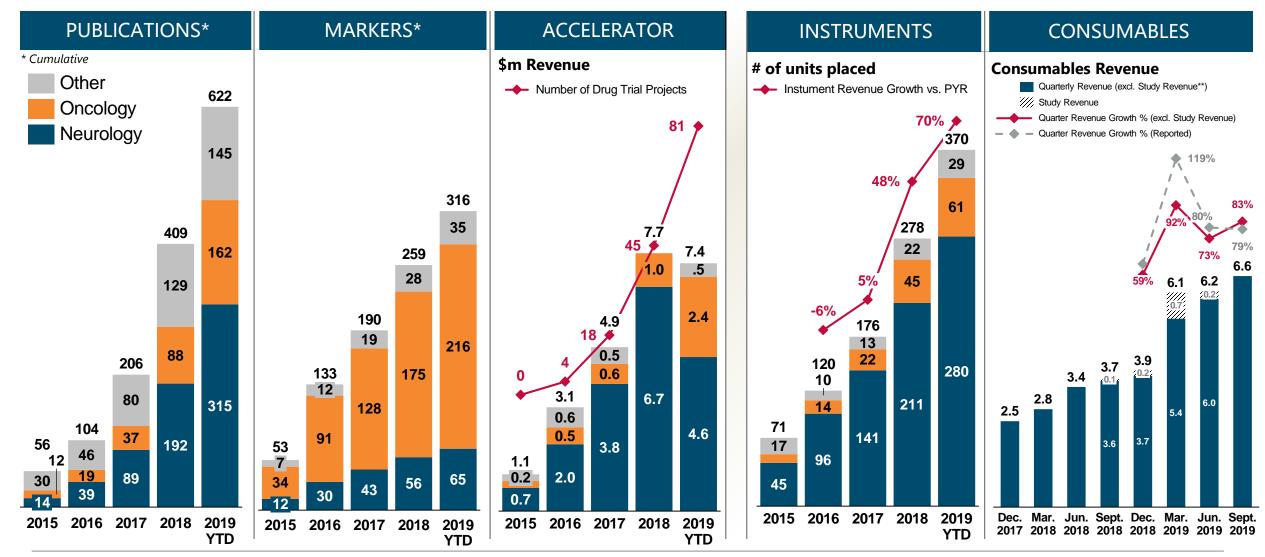
Q3 YTD 2019 Growth Stratification





Scientific Research is Driving Brand Awareness, Performance and Utilization





** Study Revenue includes \$1.2m of Consumables Revenue recognized between Q3 2018 and Q2 2019 in connection with orders from one customer for use in a large clinical study.

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Quanterix Product Offerings



	Instruments		Assay kits	Services
HD-X	SR-X	Quanterix SP-X	Plate Bead	Accelerator
Floor-standing integrated Benchtop semi- automated assay prep Assay prep and detection Major performance improvements over HD-1		Simoa planar assay Benchtop semi- automated assay prep Multiplex capabilities	300+ assays Homebrew kits Singleplex and Multiplex	Contract research & testing Custom assay development & reagent production CLIA and LDT capabilities
550+ publications				gs Call Q3 2019 November 6, 2019 10

Simoa HD-X Analyzer ahead of schedule Delivering disruption that you can count on





Unparalleled ultra-sensitivity, leveraging years of experience

Productivity improvements, greater flexibility and temperature control

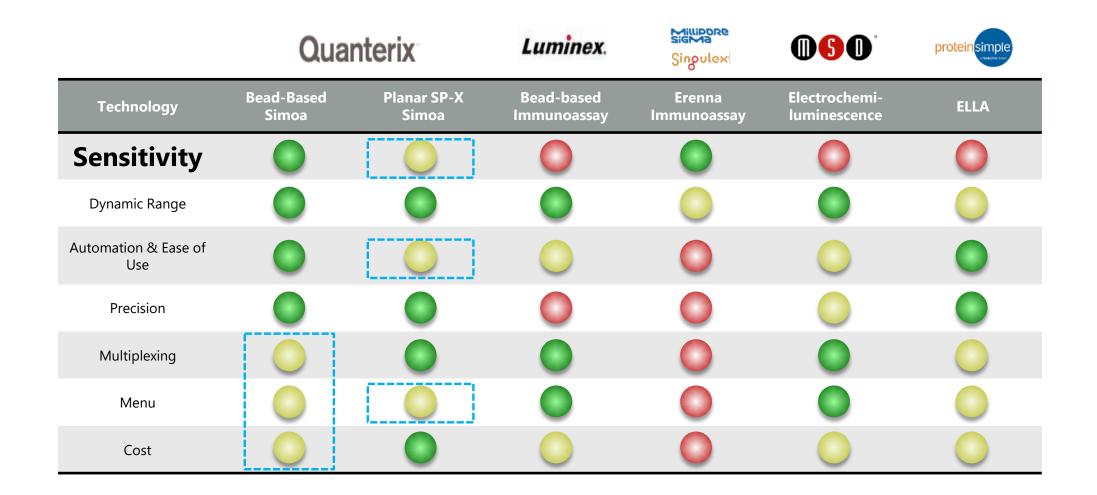
Best-in-class assay performance across a broad assay menu

Regulatory Compliance – Enables 21CFR Part 11 compliance

- Began shipping ahead of schedule in Q3
- Trade-in program exceeding expectations
- Early access customer results very promising

Competitive Landscape





2019 Goals Large strides in Q3 towards securing our 2019 objectives



		STRATEGY		
Neurology <10% Penetrated	Oncology 3x Neuro	Strategy Dx via Biopharma	Financials	Technology
		LDT/IVD partnerships	LT Growth: 40%	
High double digit growth with high utilization	Penetrate	50 phase I, II, III trials	Gross Margin: +300 bps	100x sensitivity increase by YE 2021
Add 25 assays	Immuno therapy focus	M&A to accelerate strategy	Instrument Growth: 25%+	~)
• Launched HD-X one quarter	• Launched SPX Q2'19	Clinically validate Nf-L for	• YTD Revenue Growth +53%	• 40X defined
ahead of schedule	 Strengthening our position in Oncology; PPH Onco track 	 2nd'ary endpoint & DP Acquired Uman; Licensed Siemens 	 Non-GAAP* GM 50.7%, +570 bps vs. 45.0% last year; GAAP GM 48.9% vs. 47.6% last year 	 Prototype developed
7				7

* Non-GAAP adjustments exclude (i) the impact of \$1.3M in revenue recognized in Q3 2018 in connection with the termination of a license agreement; and (ii) acquisition-related purchase accounting adjustments. Reconciliation of these non-GAAP measures to their most directly comparable GAAP financial measures is set forth in the Appendix to this presentation. Figures may not foot due to rounding.

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We are Addressing a Significant Unmet Need in Drug Development

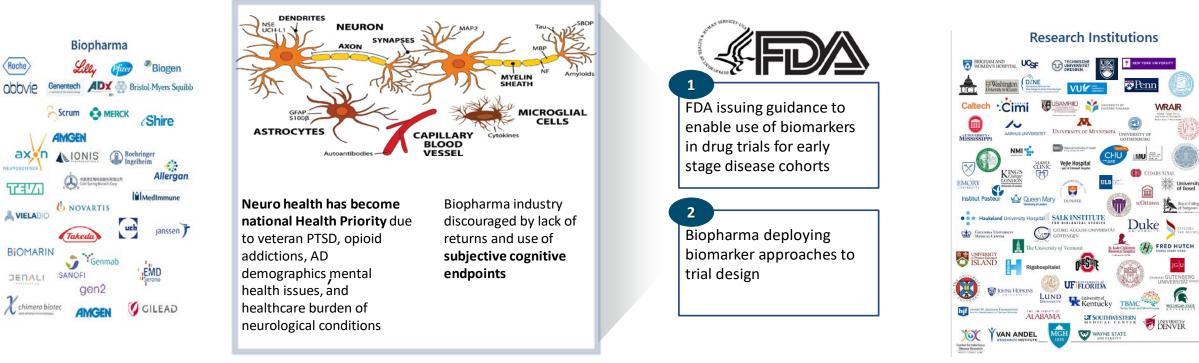


DRUG PERFORMANCE	PROBABILI	TY OF DRUG APPROVAL	VALIDATION OF SIMOA IMPACT		
TOXICITY Adverse drug events are a substantial cause of Death in USA	300% increase if biomarkers are used		Simoa'S at CROs Trials at Quanterix 38		
EFFICACY	Hematology	26.1%			
Depression 62%	Infectious disease	19.1%	2 0		
Schizophrenia 60%	Metabolic	15.3%	2015 2018 2015 2018		
Cardiac arrhythmia 60%	Gastroenterolgy	15.1%	650 clinical trials		
Asthma 60%	Autoimmune	11.1% 8.4 % Goes to	with Simoa at single CRO		
Diabetes 57%	All indications	^{9.6%} 25%	400 PHASE I		
Osteoporosis 48%	Neurology	8.4% Approval			
Hepatitis C 47%	Cardiovascular	after PHASE I	200 PHASE II		
AD 30%	Oncology 5.	1%	50 PHASE III		
Cancer 25%		ty of phase III approval Phase 1 approval	MYRIAD • RBM.		

Source: Bio Industry Analysis; Clinical Development Success Rates (June 2016)

FDA Supporting Biomarker Development



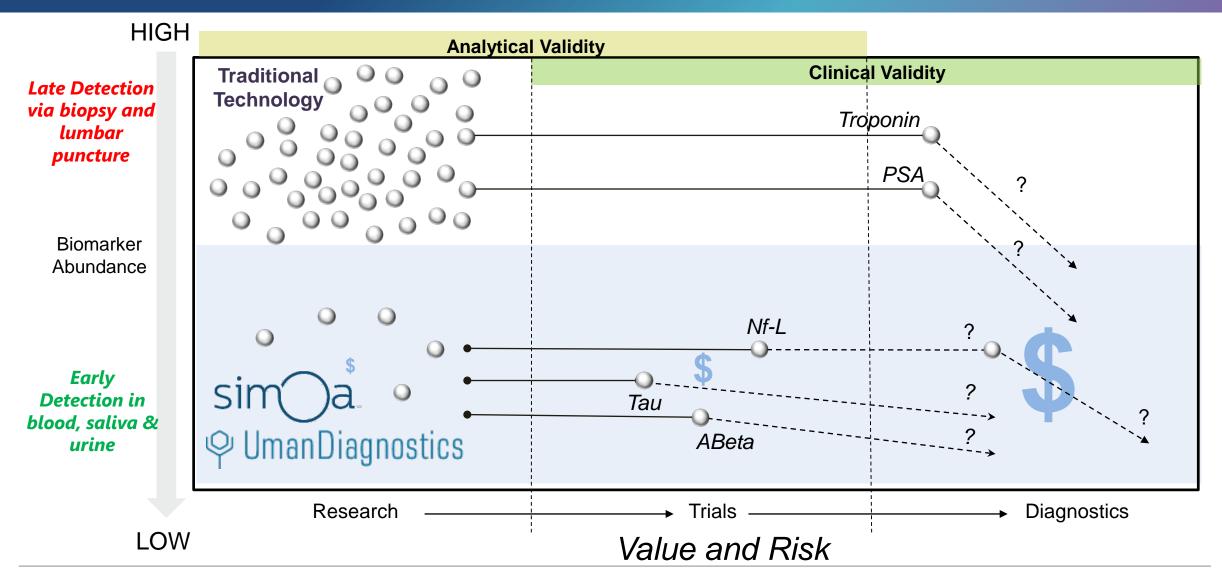


Source: Health Advances analysis, Quanterix materials.



Strategic Roadmap Analytical Validity to Clinical Validity







PEER-REVIEWED PUBLICATIONS ON Nf-L NFL PUBLICATIONS Simoa NfL 160 46 active Key meeting Abstracts and presentations 25 60 **Our Focus** clinical trials 2015 140 2016 50 using Nf-L 20 2017 PubMed Citations 40 2018 15 30 20 10 10 5 AAN 2018 AAN 2019 **ECTRIMS ECTRIMS** 2018 2019 0 60 5 Phase III trails across >1600 MS patients Parkinsons FTD ALS Alzheimers TBI MS demonstrate clinical utility of Simoa NfL STUDIES CONFIRM NFL CLINICAL UTILITY: 40 Disease activity monitoring Drug efficacy monitoring 20 Relapse/severity prognostic 0 5 2008 2010 Publication Year 2012 2014 2016 2000 2002 2004 2006 2018 Majority of published data obtained with Simoa NfL Source: Health Advances analysis, PubMed.

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Serum Nf-L Powering Major Drug Trials Advancing toward the Clinic

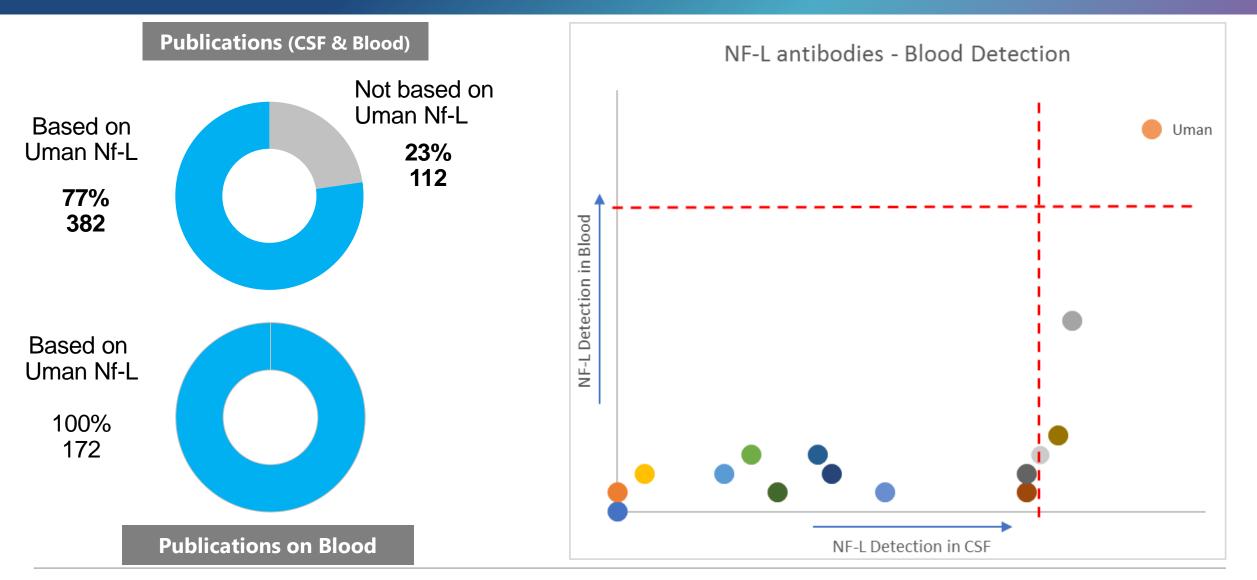




Powering Drug Trials	U NOVARTIS Arzerra () ofatumumab	 <u>Phase III ASCLEPIOS I and II trials</u>: Relapsing MS patients had reduced disease progression Nf-L measurements used as secondary endpoint, reduction after 3 months
Comparative Effectiveness of Therapeutics	Roche OCREVUS [®] ocrelizumab #####	 <u>Retrospective data from Phase III OPERA I, OPERA II, AND ORATORIO trials</u>: MS patients treated saw Nf-L levels lowered to healthy donor levels. Ocrevus: \$1.75B sales, highest MS drug share in US of 40%
Revealing Biology of Disease	JAMA Neurology	 Understanding of genetics of MS severity Identifying additional biomarkers in MS sNf-L measured shown to be increased 6 years before onset of MS
Advancing Toward Clinic	IAMA Neurology Biogen SIEMENS Healthineers	 Nf-L showing extensive evidence for future IVD assay

Uman Nf-L Antibody is Unparalleled





Quanterix to Offer Nf-L IVD Assay in Partnership with Siemens





- Non-exclusive license to Siemens to develop NF-L IVD assay on Siemens platform
- Siemens Nf-L assay combined with Quanterix/Uman's Nf-L antibody specificity provides clinically relevant sensitivity in blood/serum for Multiple Sclerosis monitoring applications
- 1M patients with MS in US, should be tested twice annually similar to MRI, TAM of 2M tests annually just for MS monitoring
- Provides pathway for Uman NF-L monetization in IVD with immediate global scale – leverage Siemens installed base globally
- Highly favorable economics (details undisclosed)
- Demand already exists for MS treatment monitoring application but volume could be significantly higher if AD, TBI, etc. clinically proven in future
- Maintains Quanterix optionality for further value creation through either diagnostic multiplex panels/algorithms and ultra-sensitive clinical applications

SIEMENS HEALTHINEERS ENTERS INTO LICENSE AND SUPPLY ARRANGEMENT WITH QUANTERIX FOR ACCESS TO NEUROFILAMENT LIGHT (NF-L) ANTIBODIES TO DEVELOP NF-L ASSAYS

BILLERICA, Mass.--(BUSINESS WIRE)--Nov. 4, 2019-- Quanterix Corporation (NASDAQ: QTRX), a company digitizing biomarker analysis to advance the science of precision health, today announced it has entered into a licensing and supply arrangement with Siemens Healthineers for access to Quanterix' proprietary NF-L antibodies, which were recently acquired from UmanDiagnostics. Access to the Nf-L antibodies will allow Siemens Healthineers to begin developing blood-based NF-L clinical tests for future commercialization. Financial terms were not disclosed.

The agreement entered into with Siemens Healthineers marks yet another significant milestone in Quanterix' mission to provide early detection of disease, measurements for treatment efficacy, and disease progression for neurological disorders. It comes on the heels of Quanterix' recent acquisition of privately held UmanDiagnostics, which has become the provider of choice of Nf-L antibodies for biopharmaceutical and diagnostic applications. This acquisition allows Quanterix to supply researchers globally with the "best-in-class" Simoa Nf-L assays, while continuing to innovate new digital biomarkers to advance the field of research in diagnostics for neurological disorders. It also positions Quanterix to capitalize on the growth fueled by the momentum in Nf-L as a tremendously promising brain biomarker in research and clinical applications. Data presented at the recent European Committee for Treatment and Research in Multiple Sclerosis (ECTRIMS) utilizing a sensitive Siemens Healthineers research Nf-L assay using Quanterix antibodies showed a high correlation with the established Simoa gold standard for measuring Nf-L in blood.

"We are pleased to have the opportunity to work with as formidable a partner as Siemens Healthineers to accelerate the availability of an Nf-L test for patients around the world," said Kevin Hrusovsky, Chief Executive Officer, President and Chairman of Quanterix. "This agreement demonstrates a clear path for our research customers working with Simoa technology to take groundbreaking advances like the Nf-L test to the clinic. We believe this will give our researchers even more confidence working with Simoa in research, while leveraging Siemens Healthineers' global footprint to allow Quanterix to share meaningfully in the large IVD (*in vitro diagnostics*) market for blood-based neurology testing."

"Expanding precision medicine leading to new and more accurate ways to improve patient outcomes is a key value of Siemens Healthineers," confirmed David Stein, Head of Global Strategy, Siemens Healthineers. "We believe Nf-L is an important blood-based neurological biomarker that, in the future, could positively impact patient outcomes by potentially aiding in earlier diagnoses, intervention, and management of patients with a broad array of neurological conditions, such as Multiple Sclerosis (MS), Alzheimer's disease, Parkinson's disease, and ALS. A blood-based Nf-L assay is highly compatible with our global diagnostic solutions, and we are pleased to be working with Quanterix for access to its unique set of Uman Nf-L antibodies for the development of high performance clinical Nf-L assays," Stein explained.

To learn more about Quanterix' Nf-L assay, click here.

About Quanterix

Quanterix is a company that's digitizing biomarker analysis with the goal of advancing the science of precision health. The company's digital health solution, Simoa, has the potential to change the way in which healthcare is provided today by giving researchers the ability to closely examine the continuum from health to disease. Quanterix' technology is designed to enable much earlier disease detection, better prognoses and enhanced treatment methods to improve the quality of life and longevity of the population for generations to come. The technology is currently being used for research applications in several therapeutic areas, including oncology, neurology, cardiology, inflammation and infectious disease. The company was established in 2007 and is located in Billerica, Massachusetts.

Forward-Looking Statements

This press release contains forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995. Words such as "may," "will," "expect," "plan," "anticipate," "estimate," "intend" and similar expressions (as well as other words or expressions referencing future events, conditions or circumstances) are intended to identify forward-looking statements. Forward-looking statements in this news release are based on Quanterix' expectations and assumptions as of the date of this press release. Each of these forward-looking statements involves risks and uncertainties. Factors that may cause Quanterix' actual results to differ from those expressed or implied in the forward-looking statements in this press release are discussed in Quanterix' filings with the U.S. Securities and Exchange Commission, including the "Risk Factors" sections contained therein. Except as required by law, Quanterix assumes no obligation to update any forward-looking statements contained herein to reflect any change in expectations, even as new information becomes available.

Digital Biomarkers Disruption Paradigm Alzheimer's Disease Opportunity – Biogen Resurrects Aducanumab





Alzheimer's disease not diagnosed until symptoms



Imaging expensive and often not covered



Therapies for later stage disease have limited effectiveness



Detect or screen in annual blood test







Therapy delivered sooner with less dosing / toxicity. Blood test monitors progression



Aducanumab from Biogen

Powering Precision Health - Summit 2019

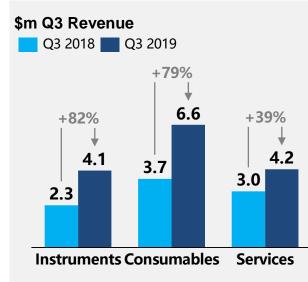
2019 Spons				Barcelona,
Quanterix The Science of Precision Health				November, 19
Abbott		2018	2019	
SVBLEERINK	Speakers	18	42	
IP.Morgan	Total Attendees	<100	Expecting 250	
COWEN	Focus	Neuro Only	Neuro & Onco concurrent tracks	
Biogen . P/W/P	Leadership	Quanterix Only	10 Member External PPH Advisory Board	

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Quanterix

Q3 2019 & YTD Sept. 2019 Financials



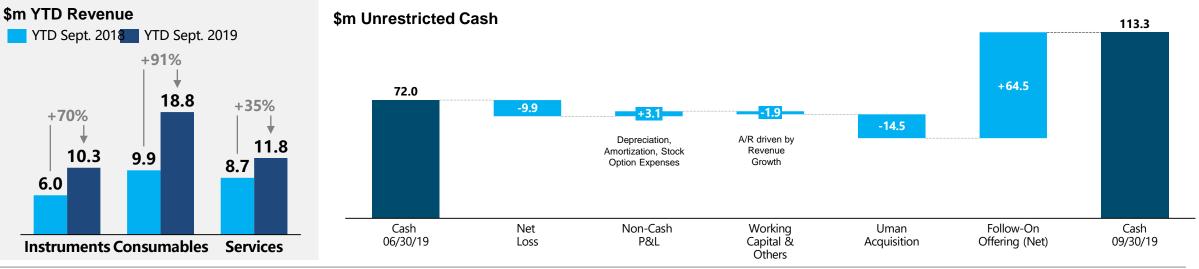


+70%

6.0

10.3

in \$m	GAAP			Non-GAAP*				
	Q3	Q3	YTD Sept.	YTD Sept.	Q3	Q3	YTD Sept.	YTD Sept.
	2019	2018	2019	2018	2019	2018	2019	2018
Total Revenue	14.9	10.6	40.8	26.8	14.9	9.2	40.8	25.4
Growth vs. PYR	41%		53%		62%		61%	
Gross Profit	7.0	5.6	20.0	12.7	7.7	4.3	20.7	11.4
Gross Margin %	47.1%	52.8%	48.9%	47.6%	51.8%	46.2%	50.7%	45.0%
Operating Expenses	17.3	13.3	50.1	34.9	17.3	13.3	50.1	34.9
Loss from Operations	-10.2	-7.7	-30.1	-22.1	-9.5	-9.0	-29.4	-23.4



* Non-GAAP item. Reconciliations are included in the Appendix to this presentation and in our Q3 2019 press release.

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Poised to Disrupt Healthcare and Create Significant Value



	Differentiator	Value
1 Category-defining; Unrivaled Sensitivity / Technology	Best in Class	Disrupt
Methodical market penetration strategy to reward investors	\$3B to \$40B	New Answers
3 DNA – RNA - Protein; Better linked to Disease / Health		Holy Grail
Quanterix 4 Validation: 19/20 top pharma, PPH, 800+ trials	600+ pubs All Areas	Proven
6 Growth & Value; Razor – razor blade, \$150M invested	Product Launches	Rapid Growth
6 Low Risk / Solid Return + Uber Return Prospect		Retail
7 Track Record for Commercializing Disruption		Lynchpin

Quanterix







(\$million)	Q3 2019	Q3 2018	YonY Change
Revenue	\$14.9	\$10.6	41%
Gross Profit	\$7.0	\$5.6	
Gross Margin (%)	47.1%	52.8%	- 570 bps
Operating Expenses	\$17.3	13.3	
Loss from	-\$10.2	-\$7.7	
Operations			
(\$million)	Q3 2019 YTD	Q3 2018 YTD	YonY Change
Revenue	\$40.8	\$26.8	53%
Gross Profit	\$20.0	\$12.7	
	40.00/	47 00/	100 hrs

Gross Profit	\$20.0	\$12.7	
Gross Margin (%)	48.9%	47.6%	+ 130 bps
Operating	\$50.1	\$34.9	
Expenses			
Loss from	-\$30.1	-\$22.1	
Operations			

Reconciliation of non-GAAP Financials in thousands (unaudited)



	2019 2018 Three months ended September 30,		2019 2018 Nine months ender September 30	
Total revenue Revenue from termination of license agreement (Note 1)	\$14,944 —	\$10,591 (1,342)	\$40,816 —	\$26,755 (1,342)
Non-GAAP total revenue	\$14,944	\$9,249	\$40,816	\$25,413
Gross profit Revenue from termination of license agreement (Note 1)	\$7,033 —	\$5,595 (1,342)	\$19,969 —	\$12,739 (1,342)
Acquisition-related purchase accounting charges (Note 2)	711	16	711	43
Non-GAAP gross profit	\$7,744	\$4,270	\$20,680	\$11,441
GAAP gross margin	47.1%	52.8%	48.9%	47.6%
Non-GAAP gross margin	51.8%	46.2%	50.7%	45.0%

Note 1: During the three-month period ended September 30, 2018, we recognized \$1.3 million in collaboration and license revenue in connection with the termination of our license agreement with bioMerieux SA.

Note 2: During each of the three and the nine months ended September 30, 2019, we incurred \$328 thousand of acquisitionrelated amortization of inventory valuation and \$383 thousand of acquisition-related amortization of intangible assets adjustments in connection with our acquisition of UmanDiagnosotics.During the three and nine months ended September 30, 2018, we incurred \$16 thousand and \$43 thousand, respectively, of acquisition-related amortization of inventory valuation adjustments in connection with our acquisition of Aushon Biosystems .