



Forward-Looking Statements & Non-GAAP Financial Measures

This presentation contains forward-looking statements within the meaning of the U.S. 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 presentation are based on Quanterix's expectations and assumptions as of the date of this presentation. Each of these forward-looking statements involves risks and uncertainties. Factors that may cause Quanterix's actual results to differ from those expressed or implied in the forward-looking statements in this presentation are discussed in Quanterix's 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.

To supplement Quanterix's financial statements presented on a GAAP basis, Quanterix has provided certain non-GAAP financial measures. Management uses these non-GAAP measures to evaluate our operating performance in a manner that allows for meaningful period-to-period comparison and analysis of trends in our business and as a factor in assessing progress against the Restructuring Plan. Management believes that presentation of these non-GAAP measures provides useful information to investors in assessing our operating performance within our industry and in order to allow comparability to the presentation of other companies in our industry. 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 herein and in the associated earnings press release.



Corporate Transformation: Progress on Assay Redevelopment Roadmap



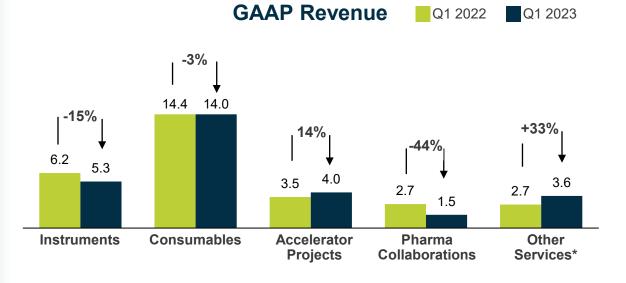


Q1 2023 Results vs PYQ1

(in millions)

	Q1 G	SAAP	Q1 Non-GAAP					
	2022	2023	2022	2023				
Revenue	29.6	28.5	29.6	28.5				
Gross Margin \$	14.6	16.9	12.8	15.1				
Gross Margin %	49.3%	59.5%	43.2%	53.1%				
Operating Expense	32.7	26.3	31.0	24.5				
Operating Loss	-18.2	-9.4	-18.2	-9.4				
Cash Usage	-22.1	-9.1	-22.1	-9.1				

	INSTRUMENTS	CONSUMABLES	ACCELERATOR	Pharma Collaborations	OTHER SERVICES*
Revenue Mix	19%	49%	14%	5%	13%

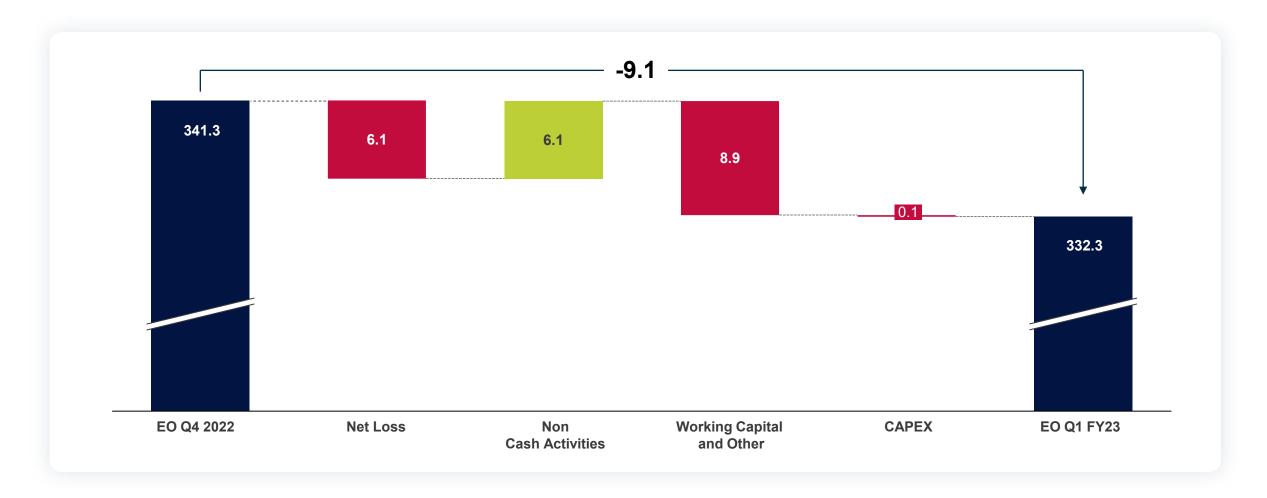


^{*}Includes Collaboration revenue



Q1 2023 Cash & Cash Burn

(in millions)



Note: Ending total cash \$332.3M; unrestricted cash balance \$329.4M and restricted cash balance \$2.9M



Scientific Validation Driving Adoption







2023 Guidance

We have made a modest, positive adjustment to our full year 2023 Revenue and Gross Margin % guidance

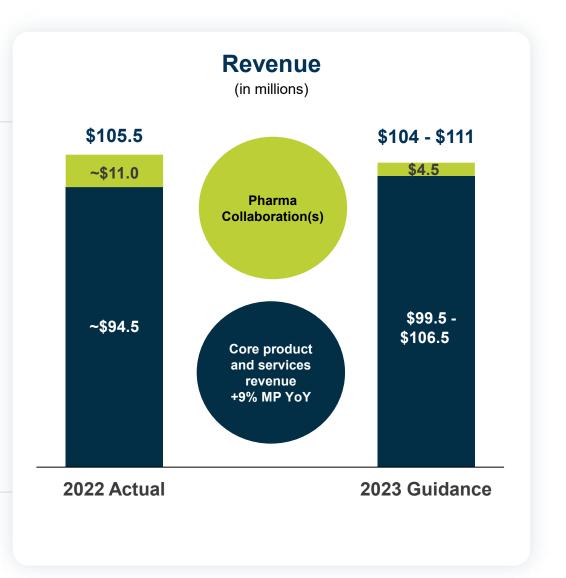
Gross Margin %

Expect FY 2023 GAAP gross margin in the high 40s and non-GAAP gross margin in the mid 40s

Cash Burn

Cash burn guidance remains unchanged. Expect cash burn to improve by approximately 10% in 2023 versus 2022

Cash flow positive at \$170-\$190M in revenue





FDA Accelerated Approval of Tofersen

On the basis of Neurofilament light as a secondary end point

Accelerated approval of Tofersen to treat genetic form of ALS

VALOR and OLE Tofersen trials showed a substantial reduction (40-50%) in sNfL levels, indicative of reduction in neuronal injury

FDA approval was based in part on sNfL as a secondary endpoint validating this biomarker's importance as a measure of neurodegeneration

Relevance for sNfL as a biomarker for neurological disorders

Approval suggests all ALS and Neuro research and drug development should be monitoring this biomarker

Potentially all clinical trials for drugs impacting neurodegeneration should be using sNfL as a biomarker

FDA Accelerated Approval of Tofersen Highlights Importance of Blood Neurofilament Light Chain as Surrogate Endpoint in Neurology **Therapeutic Trials** Blood-based measurement of neurofilament light chain (NfL) provides compelling evidence to support an accelerated drug approval by the FDA. The decision has positive implications for NfL blood testing as a key tool in neuro-drug development. BILLERICA, Mass.--(BUSINESS WIRE)--Quanterix Corporation (NASDAQ: QTRX), a company fueling scientific discovery and breakthrough diagnostics through ultrasensitive biomarker detection, today announced that blood-based NfL measurements provided compelling support for the FDA's accelerated approval of tofersen for treatment of superoxide dismutase 1 amyotrophic lateral sclerosis (SOD1-ALS), a devastating rare genetic form of ALS. This is the first known case in which a blood biomarker was successfully used as a surrogate endpoint for a neurology therapeutic trial to gain accelerated approval, highlighting the potentia for other therapeutic trial designs to benefit from including blood NfL measurements. EMBRACING A BIOMARKER Industry-led clinical trials that test drugs for neurodegenerative diseases have increasingly used the biomarker neurofilament light chain (NFL) as a measure of efficacy. NFL levels in the blood and cerebrospinal fluid are thought to correlate with neuron damage and death. Multiple sclerosis Alzheimer's disease Amyotrophic lateral sclerosis Huntington's disease Frontotemporal dementia Parkinson's disease Other 2011 2012 2013 2014 2015 2016 2017 2018 2019 2020 2021 2022 2023





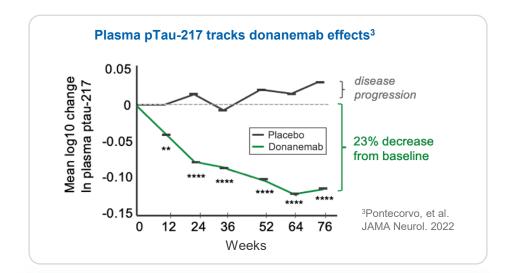
pTau-217 emerges as the top blood-based biomarker for Alzheimer's

IVD Test

Growing consensus among the scientific community pTau-217 is so accurate it has the potential to replace PET for Dx our results suggest that plasma pTau-217. may have strong enough performance to be used in combination with clinical evaluation for the diagnosis of AD" - Prof. Henrik Zetterberg, 2023 1 These data indicate that pTau-217 can detect disease before clinical signs..." - Prof. Oskar Hansson, 2023 ² ¹Therriault, et al. Alz & Dementia 2023 ²Jonaitis, et al. Brain Comm 2023 2023 2024 2025 2026 QTRX pTau-217

CLIA LDT

Expanding integration into therapeutic trial designs for enrollment & monitoring therapeutic response



Quanterix Announces New Agreements With Lilly To Advance Alzheimer's Disease Diagnosis And Treatment

License agreement provides Quanterix access to Lilly's **P-tau217** antibody technology, creating pathways for plasma-based biomarkers for used in Alzheimer's disease

March 01. 2022 07:30 AM Eastern Standard Time



roadmap

RUO Product

Quanterix Corporation

Reconciliation of GAAP Financial Measures to Non-GAAP Financial Measures (Unaudited and in thousands, except percentages)

	Three Months Ended March 31,			Three Months Ended December 31, September 30,		Three Months Ended June 30,		Year Ended December 31,						
	202		2022		2022		2022		2022		2022		2021	
GAAP gross profit	\$	16,926	\$	14,559	\$	12,592	\$	10,944	\$	8,711	\$	46,806	\$	61,728
Shipping and handling costs (1)		(1,829)		(1,781)		(1,926)	_	(1,636)		(1,868)		(7,206)		(6,892)
Non-GAAP gross profit	\$	15,097	\$	12,778	\$	10,666	\$	9,308	\$	6,843	\$	39,600	\$	54,836
GAAP Revenue		28,456		29,552		25,824		26,646		23,500		105,522		110,556
GAAP Gross margin (GAAP gross profit as % of revenue)		59.5%		49.3%		48.8%		41.1%		37.1%		44.4%		55.8%
Non-GAAP gross margin (non-GAAP gross profit as % of revenue)		53.1%		43.2%		41.3%		34.9%		29.1%		37.5%		49.6%
GAAP total operating expenses	\$	26,346	\$	32,746	\$	34,547	\$	47,547	\$	33,670	\$	148,510	\$	120,314
Shipping and handling costs (1)		(1,829)		(1,781)		(1,926)		(1,636)		(1,868)		(7,206)		(6,892)
Non-GAAP total operating costs	\$	24,517	\$	30,965	\$	32,621	\$	45,911	\$	31,802	\$	141,304	\$	113,422
GAAP loss from operations	\$	(9,420)	\$	(18,187)	\$	(21,955)	\$	(36,603)	\$	(24,959)	\$	(101,704)	\$	(58,586)
Non-GAAP loss from operations	\$	(9,420)	\$	(18,187)	\$	(21,955)	\$	(36,603)	\$	(24,959)	\$	(101,704)	\$	(58,586)

⁽¹⁾ Shipping and handling costs, which include freight and other activities costs associated with product shipments, net of charges passed on to the customer, are captured within operating expenses in our consolidated statements of operations. During the three months ended March 31, 2023 and 2022, we incurred \$1.8 million and \$1.8 million, respectively, of shipping and handling costs recorded within operating expenses. During the three months ended December 31, 2022, we incurred \$1.9 million of shipping and handling costs within operating expenses. During the three months ended September 30, 2022, we incurred \$1.9 million of shipping and handling costs within operating expenses. During the three months ended June 30, 2022, we incurred \$1.9 million of shipping and handling costs within operating expenses.





Quanterix

Leader in proteomics, positioning ourselves to accelerate faster

- Most Sensitive Protein
 Measurements in Blood
 Strong IP protection
- Translational Leader
 FDA BTD pTau181, NfL
 LDT implementation
 Pharma / KOL partnerships
- Sample to Answer Platform Rapid TAT; global footprint
- Continued Innovation
 Simoa in every lab, menu expansion, pushing limits of sensitivity further
- Strong Financials
 With commercial scale