

Niagen.

BIOSCIENCE

Earnings Presentation

Second Quarter 2026



Nasdaq: NAGE | August 4, 2026

Safe Harbor Statement

This presentation and other written or oral statements made from time to time by representatives of Niagen Bioscience contain “forward-looking statements” within the meaning of Section 27A of the Securities Act of 1933, as amended, and Section 21E of the Securities Exchange Act of 1934, as amended. Forward-looking statements reflect the current view about future events. Statements that are not historical in nature, such as 2026 financial outlook, and which may be identified by the use of words like “expects,” “anticipates,” “intends,” “estimates,” “plans,” “potential,” “possible,” “probable,” “believes,” “seeks,” “may,” “will,” “should,” “could,” “predicts,” “projects,” “continue,” “would” or the negative of these terms and other words of similar meaning, are forward-looking statements. Such statements include, but are not limited to, statements contained in this presentation relating to our expected sales, cash flows, planned investments, and financial performance, business, business strategy, expansion, growth, key drivers (including cost savings and increased investments), products and services we offer and their impact on our performance or products and services we may offer in the future and the timing of their development, sales and marketing strategy and capital outlook. Forward-looking statements are based on management’s current expectations and assumptions regarding our business, the economy and other future conditions and are subject to inherent risks, uncertainties and changes of circumstances that are difficult to predict and may cause actual results to differ materially from those contemplated or expressed. We caution you therefore against relying on any of these forward-looking statements. These risks and uncertainties include those risk factors discussed in Part I, “Item 1A. Risk Factors” of our most recent Annual Report on Form 10-K and Quarterly Report on Form 10-Q filed with the Securities and Exchange Commission (the “Commission”), and in subsequent filings with the Commission. Any forward-looking statements are qualified in their entirety by reference to the factors discussed in these filings with the Commission. Should one or more of these risks or uncertainties materialize, or should the underlying assumptions prove incorrect, actual results may differ significantly from those anticipated, believed, estimated, expected, intended or planned.

Important factors that could cause actual results to differ materially from those in the forward looking statements include but are not limited to: our relationships with major customers; a decline in general economic conditions nationally and internationally; the market and size of the vitamin, mineral and dietary supplement market and the intravenous market; decreased demand for our products and services; market acceptance of our products; interruptions in our ability to sell products through third-party online marketplaces; the ability to protect our intellectual property rights; impact of any litigation or infringement actions brought against us; competition from other providers and products, including other supplements such as NMN; risks in product development; our ability to develop pharmaceutical business; the Company's therapeutic pipeline, including NB4168, preclinical and IND-enabling studies, and regulatory designations and development plans; inability to raise capital to fund continuing operations or new product development; changes in government regulation or regulatory priorities of government officials; the ability to complete customer transactions and capital raising transactions; inflationary conditions and adverse economic conditions; our history of operating losses; the growth and profitability of our product sales; our ability to maintain and grow sales, marketing and distribution capabilities; changing consumer perceptions of our products; our reliance on a single or limited number of third-party suppliers; risks of conducting business in China; unanticipated developments in and risks related to the Company's ability to secure adequate quantities of pharmaceutical-grade Niagen in a timely manner; the Company's ability to obtain appropriate contracts and arrangements with U.S. FDA-registered 503B outsourcing facilities required to compound and distribute pharmaceutical-grade Niagen to clinics; the Company's ability to remain on the U.S. FDA Bulk Drug Substances Nominated for Use in Compounding Under Section 503B of the Federal Food, Drug, and Cosmetic Act Category 1 list; the Company's ability to maintain and enforce the Company's existing intellectual property and obtain new patents; whether the potential benefits of NRC can be further supported; further research and development and the results of clinical trials possibly being unsuccessful or insufficient to meet applicable regulatory standards or warrant continued development; the ability to enroll sufficient numbers of subjects in clinical trials; determinations made by the FDA and other governmental authorities, including with respect to products seeking to compete in our market; mislabeling or other misleading marketing practices by competitors; economic and market instability, including as a result of tariffs or trade conflicts; and the risks and uncertainties associated with our business and financial condition in general.

Factors or events that could cause our actual results to differ may emerge from time to time, and it is not possible for us to predict all of them. We cannot guarantee future results, levels of activity, performance or achievements. Except as required by applicable law, including the securities laws of the United States, we do not intend to update any of the forward-looking statements to conform these statements to actual results.

About Non-GAAP Financial Measures

Niagen Bioscience’s non-GAAP financial measure, Adjusted EBITDA, is defined as net income before interest, provision for income taxes, depreciation, amortization, non-cash share-based compensation costs, severance and restructuring expense and other infrequent items, including gains recognized related to the sale of an operating segment and a royalty settlement. Niagen Bioscience used this non-GAAP measures when evaluating its financial results as well as for internal resource management, planning and forecasting purposes. This non-GAAP measure should not be viewed in isolation from or as a substitute for Niagen Bioscience’s financial results in accordance with GAAP. Reconciliation of this non-GAAP measure to the most directly comparable GAAP measure is attached to this presentation.

FDA Disclaimer

Statements made in this presentation have not been evaluated by the Food and Drug Administration. Niagen Bioscience products are not intended to diagnose, treat, cure, or prevent any disease. The statements in this presentation are for investor relations and educational purposes only and not intended for consumers or vendors.

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Rob Fried
Chief Executive Officer



Ozan Pamir
Chief Financial Officer



Andrew Shao
*SVP, Global Regulatory
& Scientific Affairs*



Carlos Lopez
SVP, General Counsel



Michiko Kelley
Chief Marketing Officer



Second Quarter 2026 Highlights

\$29.8 million net sales

64.8% gross profit margin

\$1.0 million net income

Diluted EPS of **\$0.01**

\$3.0 million Adjusted EBITDA⁽¹⁾

\$66.7 million cash, no outstanding borrowings

(1) See slide 12 for the non-GAAP reconciliation

World's Leading NAD+ Platform

CORE BUSINESS



INJECTABLE & TELEHEALTH



SKINCARE



PHARMACEUTICAL



Ingredient partners

Executing Long-Term Strategic Growth Initiatives

Key strategic initiatives during Q2 2026.

Consumer Products

- Direct-to-consumer sales on the Company's website grew by 23% year-over-year.
- Completed and sold out of the pilot launch of the first skincare product, Nanocloud™.

Niagen Plus

- Launched clinician-directed telehealth platform under Niagen Plus, expanding the Niagen Plus clinic channel beyond in-person settings and introducing a direct-to-patient access model.
- Received LegitScript certification for Niagen Plus, allowing the company to advertise Niagen Plus at-home injection kits
- Added Olympia Pharmaceuticals as a 503B compound pharmacy partner.

Therapeutic Development

- Introduced a proprietary lead candidate, NB4168, to address rare genetic diseases and aging related disorders, initially targeting Ataxia-Telangiectasia.

Scientific Leadership

- Rebranded the Niagen Research Program and appointed new scientific leadership to support the Company's expanding external research strategy.

Q2 2026: Investing for Long-Term Growth While Maintaining Financial Discipline

Net Sales

QTD Q2 2026

\$29.8M

Tru Niagen **\$24.2M**

Gross Margin

QTD Q2 2026

64.8%

Net Income

QTD Q2 2026

\$1.0M

Adj. EBITDA⁽¹⁾ **\$3.0M**

Cash Generation

YTD Q2 2026

\$1.6M

Cash from operations

Cash & Liquidity

Quarter-End Cash: **\$66.7M**

Borrowings: **None**

Strong balance sheet supports our investment in long-term growth.

2026 Financial Priorities



E-commerce to grow 10-15%, **remaining consumer** business in line with expectations



Cash from **core business** to fund the strategic optionality of pharma, skincare and telehealth



Deploy capital toward initiatives with long-term growth potential



Maintain **financial flexibility** through disciplined capital allocation

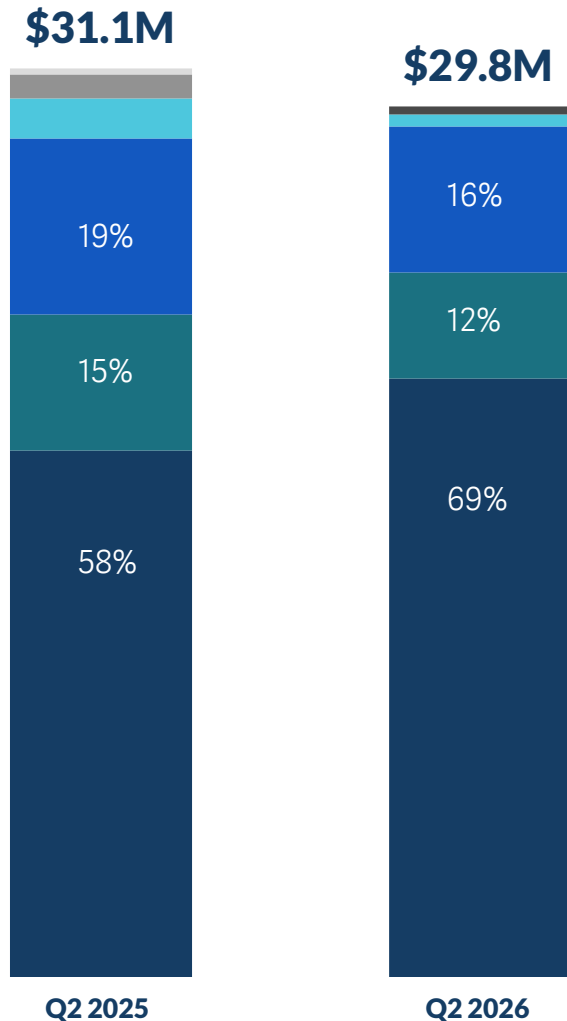
(1) See slide 12 for the non-GAAP reconciliation.

Financial Highlights

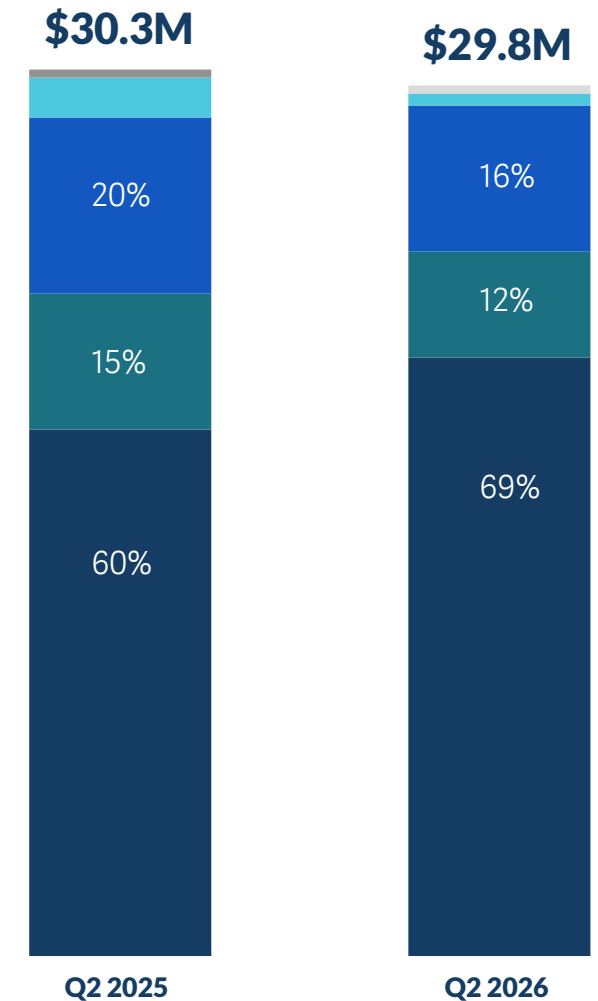


Q2 2026 Net Sales: **\$29.8M**

Total Company:



Total Company, excluding divested segment:



Tru Niagen



E-Commerce

Watson's & Other B2B

Niagen Ingredients



Food-grade Niagen®

Pharmaceutical-grade Niagen®



Other Ingredients

Analytical Reference Standards & Services⁽¹⁾



Corporate and Other⁽²⁾

Q2 2026

\$24.2M

Q2 2025

\$22.7M

\$20.5M

\$18.1M

\$3.6M

\$4.6M

\$5.4M

\$7.4M

\$4.9M

\$6.0M

\$0.4M

\$1.4M

\$0.0M

\$0.2M

\$0.0M

\$0.8M

\$0.2M

\$0.0M

(1) In February 2026, the Company divested its Analytical Reference Standards and Services operating segment.

(2) Beginning March 2026 includes net sales from the transition services agreement (TSA) related to the disposition of the operating segment. TSA revenue is expected to continue only through the agreement's anticipated completion in Q3 2026.

Note: Amounts presented in millions may not tie or sum to reported amounts in our 10-Q due to rounding.

2025 – 2026 Net Sales Summary

(\$ in millions)

	2026		2025				
	Q1	Q2	Q1	Q2	Q3	Q4	FY
Ecommerce	19.2	20.5	16.8	18.1	19.0	20.2	74.1
Watson's & Other B2B	3.2	3.6	4.7	4.6	7.0	7.3	23.6
Total Tru Niagen	22.4	24.2	21.5	22.7	26.0	27.5	97.7
Food-grade Niagen	7.3	4.9	7.0	6.0	6.4	4.7	24.1
Pharmaceutical-grade Niagen	0.9	0.4	1.0	1.4	0.5	0.9	3.8
Total Niagen Ingredient	8.2	5.4	8.0	7.4	6.9	5.6	27.9
Niagen Related Revenues	30.6	29.5	29.5	30.1	32.9	33.1	125.6
Other Ingredients	0.4	0.0	0.2	0.2	0.3	0.0	0.7
Analytical Reference Standards & Services ⁽¹⁾	0.4	0.0	0.8	0.8	0.8	0.7	3.1
Corporate and Other ⁽²⁾	0.1	0.2	0.0	0.0	0.0	0.0	0.0
Total Net Sales	31.5	29.8	30.5	31.1	34.0	33.8	129.4
Total Net Sales, Excluding Divested Segment	31.1	29.8	29.7	30.3	33.2	33.1	126.3
Tru Niagen as % of Total Net Sales	71 %	81 %	71 %	73 %	77 %	81 %	75 %
Niagen Related Revenues as % of Total Net Sales	97 %	99 %	97 %	97 %	97 %	98 %	97 %
YoY Growth Rate - Net Sales							
Total Company	3 %	(4)%	38 %	37 %	33 %	16 %	30 %
Niagen Related	4 %	(2)%	37 %	38 %	33 %	18 %	31 %
Total Tru Niagen	4 %	6 %	24 %	22 %	44 %	21 %	27 %

(1) In February 2026, the Company divested its Analytical Reference Standards and Services operating segment.

(2) Beginning March 2026 includes net sales from the transition services agreement (TSA) related to the disposition of the operating segment. TSA revenue is expected to continue only through the agreement's anticipated completion in Q3 2026.

Note: Amounts presented in millions may not tie or sum to reported amounts in our 10-Q due to rounding.

Key P&L Metrics

(\$ in thousands)	Three Months Ended June 30,		Six Months Ended June 30,	
	2026	2025	2026	2025
Net Sales	29,785	31,117	61,259	61,598
Gross Profit % of Net Sales	19,314 64.8%	20,226 65.0%	39,290 64.1%	39,557 64.2%
Sales and Marketing % of Net Sales	10,130 34.0%	8,207 26.4%	19,805 32.3%	16,324 26.5%
Research and Development	1,513	1,567	2,994	2,825
General and Administrative ⁽¹⁾	6,971	7,267	14,215	12,451
Operating Income	700	3,185	2,276	7,957
Nonoperating - Gain on Sale of Operating Segment ⁽²⁾	(3)	—	4,781	—
Net Income	963	3,609	7,281	8,672
Adjusted EBITDA ⁽³⁾	3,037	5,049	6,872	9,943

(1) G&A during the six months ended June 30, 2025 was reduced by a \$1.3 million recovery of prior credit losses related to the Elysium Health legal settlement.

(2) Reflects the gain on sale related to the divestiture of the analytical reference standards and services operating segment in February 2026.

(3) See slide 12 for the non-GAAP reconciliation.

Adjusted EBITDA Summary

Reconciliation of Non-GAAP Financial Measures

(In thousands)

	Three Months Ended				FY 2025	Three Months Ended	
	Q1 2025	Q2 2025	Q3 2025	Q4 2025		Q1 2026	Q2 2026
Net income, as reported	\$ 5,063	\$ 3,609	\$ 4,578	\$ 4,132	\$ 17,382	\$ 6,318	\$ 963
<i>Adjustments</i>							
Interest income, net	(459)	(552)	(564)	(552)	(2,127)	(375)	(372)
Provision for income taxes	168	128	222	292	810	417	106
Depreciation	158	158	157	139	612	116	117
Amortization of intangibles	37	38	38	60	173	175	174
Noncash lease expense	173	159	164	169	665	173	179
Share-based compensation	1,075	1,488	1,756	1,748	6,067	1,716	1,723
Severance and restructuring	4	21	10	53	88	79	144
Gain on settlement of royalty obligation (1)	—	—	—	(1,983)	(1,983)	—	—
Recovery of credit losses related to legal settlement (2)	(1,325)	—	—	—	(1,325)	—	—
Gain on sale of operating segment (3)	—	—	—	—	—	(4,784)	3
Adjusted EBITDA	\$ 4,894	\$ 5,049	\$ 6,361	\$ 4,058	\$ 20,362	\$ 3,835	\$ 3,037

Adjusted EBITDA remained positive at \$3.0 million in Q2 2026, reflecting focus on balancing profitability with investments designed to support long-term shareholder value.

(1) Gain recognized related to the settlement of royalty obligations from an agreement with Queen's University Belfast.

(2) The recovery of credit losses stems from the 2024 legal settlement with Elysium Health, LLC, paid in two installments, reversing a bad debt write-off from 2019.

(3) Gain related to the sale of the Company's Analytical Reference Standards and Services operating segment.

Quarterly Balance Sheet Highlights

(in thousands)	6/30/25	9/30/25	12/31/25	3/31/26	6/30/26	Key Drivers Since Year-End (Q4 2025 vs Q2 2026)
Cash	\$60,474	\$64,290	\$64,788	\$66,549	\$66,740	Up \$2.0 million primarily driven by positive operating results and proceeds from the sale of the non-core business, partially offset by share repurchases
Inventory	14,406	18,791	20,424	24,016	20,640	Remained well controlled as the Company balanced inventory with expected demand
Trade Receivables	9,656	8,506	9,741	13,068	8,580	Down \$1.2 million, primarily reflecting normal collection timing
Accrued Liabilities	7,381	8,700	7,722	7,621	3,841	Down \$3.9 million primarily due to payment of the liability settlement in Q1 2026 and normal changes in accrued compensation and other operating accruals
Accounts Payable	13,680	12,742	10,796	13,277	9,439	Down \$1.4 million reflecting lower inventory purchases and normal disbursement timing
Equity	\$64,195	\$70,676	\$76,533	\$82,330	\$82,528	Up \$6.0 million driven by earnings, share-based compensation and stock option exercises, partially offset by share repurchases

Financial strength supports disciplined investment, strategic flexibility, and long-term shareholder value creation.

Cash Flow Highlights

(in thousands)	Three Months Ended				FY 2025	Three Months Ended		Six Months Ended June 30, 2026
	3/31/25	6/30/25	9/30/25	12/31/25		3/31/26	6/30/26	
Net Income	\$5,063	\$3,609	\$4,578	\$4,132	\$17,382	\$6,318	\$963	\$7,281
Working Capital Changes	2,681	(4,399)	(3,004)	(3,669)	(8,391)	(5,017)	(334)	(5,351)
Cash Provided by / (Used for) Operations	7,883	1,250	3,692	679	13,504	(1,194)	2,772	1,578
Cash Provided by / (Used for) Investing	(32)	(135)	(24)	(101)	(292)	5,245	(92)	5,153
Cash Provided by / (Used for) Financing	3,105 ⁽¹⁾	3,743 ⁽²⁾	148 ⁽³⁾	(80) ⁽⁴⁾	6,916 ⁽⁵⁾	(2,290) ⁽⁶⁾	(2,489) ⁽⁷⁾	(4,779) ⁽⁸⁾
Net Increase in Cash	\$10,956	\$4,858	\$3,816	\$498	\$20,128	\$1,761	\$191	\$1,952
Ending Cash Balance	\$55,616	\$60,474	\$64,290	\$64,788	\$64,788	\$66,549	\$66,740	\$66,740

Positive operating cash flow and **\$66.7M** cash balance and no borrowings.

1. Includes \$3.1 million in proceeds from the exercise of stock options.
2. Includes \$3.7 million in proceeds from the exercise of stock options.
3. Includes \$0.2 million in proceeds from the exercise of stock options.
4. Includes \$0.2 million in proceeds from the exercise of stock options and (\$0.3) million in repurchase of common stock.
5. Includes \$7.2 million in proceeds from the exercise of stock options and (\$0.3) million in repurchase of common stock.

6. Includes \$0.1 million in proceeds from the exercise of stock options and (\$2.4) million in repurchase of common stock.
7. Includes \$0.2 million in proceeds from the exercise of stock options, \$0.1 million in proceeds from ESPP, and (\$2.8) million in repurchase of common stock.
8. Includes \$0.3 million in proceeds from the exercise of stock options, \$0.1 million in proceeds from ESPP, and (\$5.1) million in repurchase of common stock.

The Science

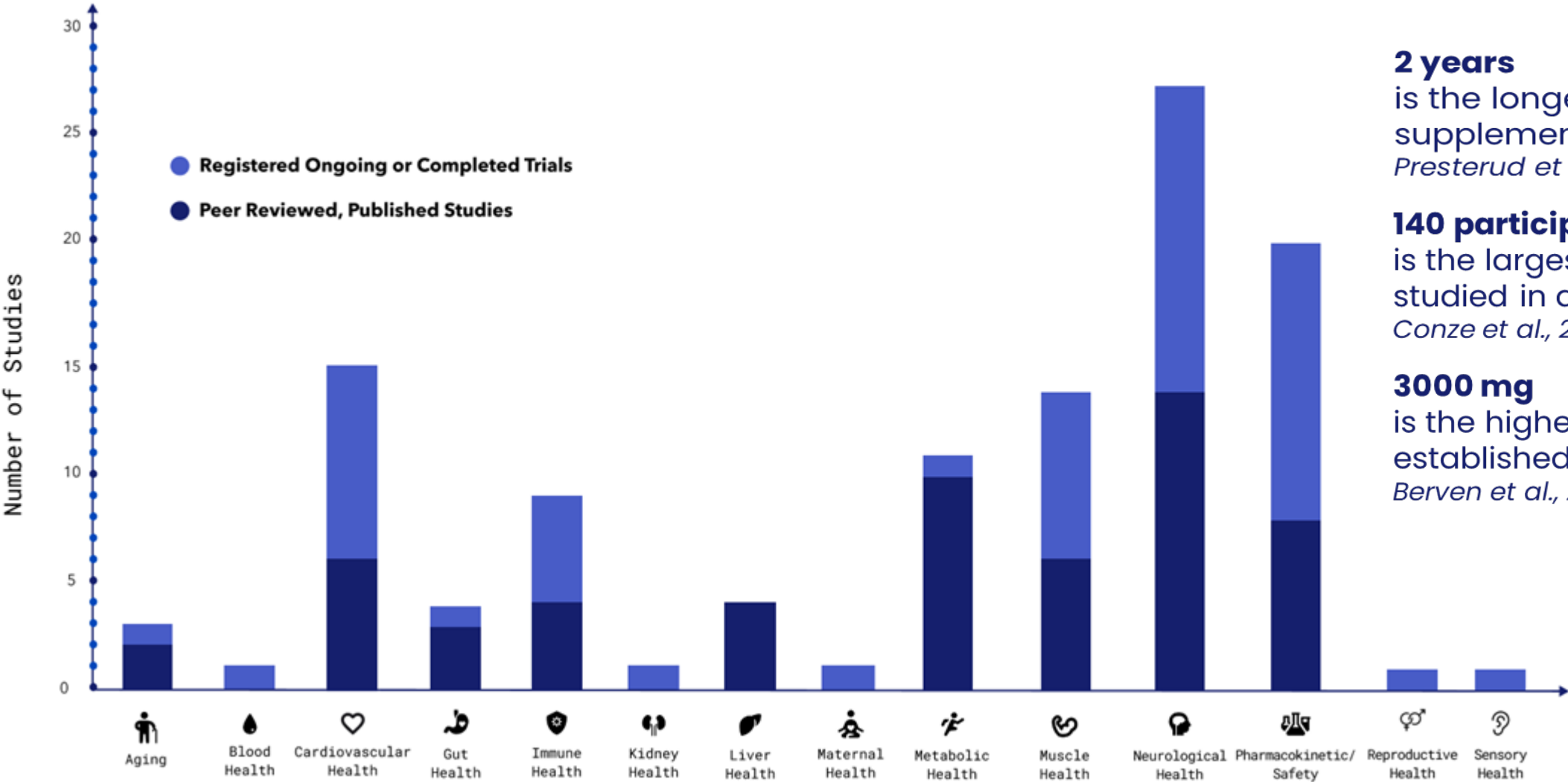


Scientific Advisory Board

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Alfred E Mann Family Foundation Chair, Department of Diabetes & Cancer Metabolism City of Hope World's Foremost Authority on NAD+ Metabolism	Professor of Structural Biology Stanford University Nobel Prize Winner, Chemistry, 2006	Kennedy Professor of Neurology Harvard University Leading Alzheimer's Researcher, TIME 100 Most Influential 2015	Chairman of Food, Nutrition, & Health University of California, Davis Leader in Food, Nutrition, & Wellness Innovation	Dean & Professor of Gerontology, Medicine and Biological Sciences University of Southern California Aging expert with pioneering research and discoveries on mitochondria and novel microproteins	Associate Professor of Molecular Medicine Scripps Research Institute Renowned Breast Cancer Researcher focused on NAD+ supplementation	Professor in Genome Instability and Neurodegeneration, Department of Cellular and Molecular Medicine, University of Copenhagen. One of the world's most published researchers on aging and neurodegenerative disease

Clinical Studies on Oral Niagen® in Multiple Health Areas

Registered Ongoing or Completed Niagen® Trials and Peer Reviewed Clinical Studies



2 years
is the longest duration of supplementation
Presterud et al., 2023

140 participants
is the largest population studied in a clinical trial
Conze et al., 2019

3000 mg
is the highest dose with established safety
Berven et al., 2023

Note: Highlighted achievements in duration, participation, and dosage only consider peer-reviewed, published studies. Status of clinical studies presented as of July 23, 2026.



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