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

Date of Report (Date of earliest event reported) April 7, 2025

SONOMA PHARMACEUTICALS, INC.

(Exact name of registrant as specified in its charter)

Delaware (State or other jurisdiction of incorporation) **001-33216** (Commission File Number) 68-0423298 (IRS Employer Identification No.)

5445 Conestoga Court, Suite 150 Boulder, CO 80301

(Address of principal executive offices) (Zip Code)

(800) 759-9305

(Registrant's telephone number, including area code)

Not applicable.

(Former name or former address, if changed since last report)

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions (see General Instruction A.2. below):

□ Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)

□ Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)

□ Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))

□ Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

Securities registered pursuant to Section 12(b) of the Act:

Title of each class	Trading symbol(s)	Name of each exchange on which registered
Common Stock	SNOA	The Nasdaq Stock Market LLC

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (17 CFR §230.405) or Rule 12b-2 of the Securities Exchange Act of 1934 (17 CFR §240.12b-2). Emerging growth company \Box

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act.

Item 8.01 Other Events.

Attached is the updated investor presentation of Sonoma Pharmaceuticals, Inc. and its affiliates (the "Company"). The presentation materials are filed hereto as Exhibit 99.1.

Except for historical information herein, matters set forth in this report are forward-looking within the meaning of the "safe harbor" provisions of the Private Securities Litigation Reform Act of 1995, including statements about the commercial and technology progress and future financial performance of the Company.

These forward-looking statements are identified by the use of words such as "will," "develop," "project," "expect," and "expand," among others. Forward-looking statements in this report are subject to certain risks and uncertainties inherent in the Company's business that could cause actual results to vary, including such risks that regulatory clinical and guideline developments may change, scientific data may not be sufficient to meet regulatory standards or receipt of required regulatory clearances or approvals, clinical results may not be replicated in actual patient settings, protection offered by the Company's patents and patent applications may be challenged, invalidated or circumvented by its competitors, the available market for the Company's products will not be as large as expected, the Company's products will not be able to penetrate one or more targeted markets, revenues will not be sufficient to meet the Company's cash needs, fund further development, as well as uncertainties relative to fluctuations in foreign currency exchange rates, global economic conditions, prospective tariffs or changes to trade policies, varying product formulations and a multitude of diverse regulatory and marketing requirements in different countries and municipalities, and other risks detailed from time to time in the Company's filings with the Securities and Exchange Commission. The Company disclaims any obligation to update these forward-looking statements, except as required by law.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits.

E-1.11.14

Exhibit	
Number	Description
99.1	Investor Presentation as of April 7, 2025

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.

SONOMA PHARMACEUTICALS, INC.

Date: April 7, 2025

By:/s/ Amy TromblyName:Amy TromblyTitle:Chief Executive Officer

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Legal Disclaimers

INVESTOR PRESENTATION

This communication is for informational purposes only. The information contained herein does not purport to be all-inclusive. The data contained herein is derived from various internal and external sources. No representation is made as to the reasonableness of the assumptions made within or the accuracy or completeness of any information contained herein. Any data on past performance is no indication as to future performance. Sonoma Pharmaceuticals, Inc. and its subsidiaries ("Sonoma" or, the "Company") assumes no obligation to update the information in this communication. This presentation is not an offer to buy or the solicitation of an offer to sell Sonoma securities.

FORWARD-LOOKING STATEMENTS

Except for historical information herein, matters set forth in this presentation are forward-looking within the meaning of the "safe harbor" provisions of the Private Securities Litigation Reform Act of 1995, including statements about the commercial and technology progress and future financial performance of Sonoma Pharmaceuticals, Inc. and its subsidiaries.

These forward-looking statements are identified by the use of words such as "believe," "achieve," and "expect," among others. Forward-looking statements in this presentation are subject to certain risks and uncertainties inherent in the Company's business that could cause actual results to vary, including such risks that regulatory clinical and guideline developments may change, scientific data may not be sufficient to meet regulatory standards or receipt of required regulatory clearances or approvals, clinical results may not be replicated in actual patient settings, the Company's products will not have sufficient capital to implement its business plan, invalidated or circumvented by its competitors, the available market for the Company's products will not be sufficient to fund further development and clinical studies, uncertainties relative to fluctuations in foreign currency exchange rates, global economic conditions, prospective tariffs or changes to trade policies, as well as uncertainties relative to varying product formulations and a multitude of diverse regulatory and marketing requirements in different countries and municipalities, and other risks detailed from time to time in the Company's filings with the Securites and Exchange Commission. The Company disclaims any obligation to update these forward-looking statements, except as required by law.

TRADEMARKS AND INTELLECTUAL PROPERTY

All trademarks, service marks, and trade names of the Company and its subsidiaries or affiliates used herein are trademarks, service marks, or registered trademarks of the Company as noted herein. Any other product, company names, or logos mentioned herein are the trademarks and/or intellectual property of their respective owners.



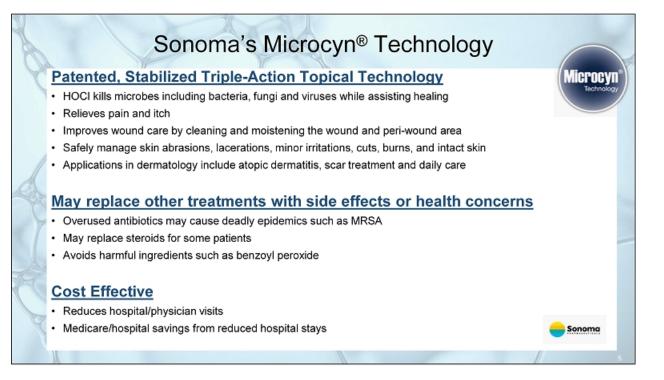
About Sonoma

Sonoma Pharmaceuticals is a global healthcare leader for developing and producing stabilized hypochlorous acid (HOCI) products for a wide range of applications, including wound, eye, oral and nasal care, dermatological conditions, podiatry, animal health care and non-toxic disinfectants. Sonoma's products are clinically proven to reduce itch, pain, scarring, and irritation safely and without damaging healthy tissue. In-vitro and clinical studies of HOCI show it to safely manage skin abrasions, lacerations, minor irritations, cuts, and intact skin.



Sonoma's products are sold either directly or via partners in 55 countries worldwide. Sonoma actively seeks new distribution partners.

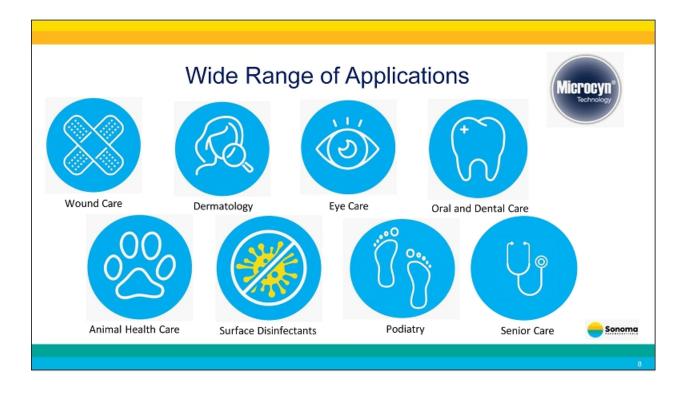






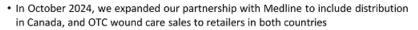
Over 100 research articles and case and clinical studies showcasing both the efficacy and safety of our Microcyn[®] technology

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Our Wound Care Business is Growing in the U.S. and Internationally

- · Celebrating 20 years of selling wound care
- In August 2024, we added Medline Industries, LP as a U.S. distributor for our wound care products – Medline is one of the largest medical supply distributors in the United States



- We sell wound care across Europe and added a new distributor for wound care in Ukraine in 2024
- Expanding Product Lines: including new options for our Microcyn[®] Negative Pressure Wound Therapy Solution product line.
- Product innovation: In June 2023, we announced a new application of our Microcyn[®] technology for intraoperative pulse lavage irrigation treatment.



Industry-Leading Safe and Effective Dermatology Products

- · Products to relieve symptoms of atopic dermatitis, reduce scarring, and for daily skin care
- · Line of office dispense products targeting growing med spa market
- · Eczema seal of approval



- All natural ingredients
- No harmful chemicals
- OTC, office dispense, and Rx products available
- · Innovating packaging available including high end and cost competitive options
- Continual Product Innovation: Introduced LumacynTM Clarifying Mist in 2024, a daily toner formulated to reduce redness and manage blemishes by reducing infections.











Growth Strategy

Continue to introduce new high margin products

 Launch new applications of Sonoma's Microcyn® technology for consumer-focused products, including in the dermatology and aesthetic space

Expand Rx and OTC reach in U.S.

- Increase sales of dermatology products through the direct dispense model that generates Sonoma's highest margins
- Increase direct-to-consumer marketing in niche markets
- Seek retail partners for OTC products
- · Expand eye care marketing for Rx and OTC

Manufacturing capabilities can support significant future growth, leading to improved margins overall

- Efficient, flexible and FDA-regulated manufacturing facility currently operating at only 30% capacity
- · Seek new manufacturing customers to utilize overflow space

Invest in R&D to expand commercialization opportunities

- · Seek additional regulatory clearances for expanded indications
- Introduce new technology to new markets
- Fully commercialize robust pipeline of new products via directto-consumer sales or distribution partnerships

Sonoma



Recent Business Developments

Introduction of new products using our Microcyn® technology

- In December 2024, we announced the relaunch of our prescription eye care product, Acuicyn[®], our prescription dermatology
 products Celacyn[®], Levicyn[®] and Epicyn[®], and our over-the-counter Lasercyn[®] Dermal Spray and Lasercyn Gel.
- In September 2024, we announced the consumer-friendly redesign of Ocucyn[®] Eyelid & Eyelash Cleanser.
- In April 2024, we announced expansion of our Microcyn[®] Negative Pressure Wound Therapy Solution product line, now available in 250mL, 450mL and 990mL sizes to meet the diverse needs of healthcare professionals and patients.
- In January 2024, we launched Lumacyn™ Clarifying Mist, a new direct-to-consumer skincare product in the United States. Lumacyn is an all-natural daily toner formulated with Microcyn technology to soothe the skin and relieve irritation.
- In June 2023, we announced a new application of our Microcyn technology for intraoperative pulse lavage irrigation treatment, which can replace commonly used IV bags in a variety of surgical procedures.
- Our MicrocynAH[®] products are now available through Pets at Home, with over 450 stores across the UK, through our partner Compana Pet Brands
 - In May 2024, we announced expansion of our MicrocynAH animal health care products in the Menards[®] chain of home improvement stores in the United States, through Compana Pet Brands.
- In April 2023, we launched Podiacyn[™] Advanced Everyday Foot Care direct to consumers for over-the-counter use in the United States.

Recent Business Developments

Expanding commercialization opportunities by investing in R&D

- We successfully transitioned all of our commercialized products in Europe, including eye care, wound care, scar gel, acne products and atopic dermatitis products to the new European Union (EU) Medical Device Regulation (MDR).
- Our manufacturing facility and five of our products are successfully registered with the Medicines & Healthcare products Regulatory Agency (MHRA) in the United Kingdom, including our wound irrigation solution, scar management products, wound hydrogel, and skin exfoliant.
- In 2024 we received two new 510(k) clearances from the FDA, including specific over-the-counter indications for the face, eyelid
 and eyelashes, as well as improved biocompatibility for our Microcyn-based solution and hydrogel.
- In March 2023, we announced new EPA claims for Nanocyn® Hospital-Grade Disinfectant for effective use against MRSA, Salmonella, Norovirus, Poliovirus, and as a fungicide. Nanocyn was previously approved for use against COVID-19 and emerging pathogens including Ebola virus, Mpox, and SARS-CoV-2. Nanocyn also received the esteemed Green Seal[®] Certification. In August 2024, the Australian TGA approved new claims for use against C. auris and C. diff.
- Reliefacyn® Advanced Itch-Burn-Rash-Pain Relief Hydrogel was awarded the NATIONAL ECZEMA Association SEAL OF ACCEPTANCE
- A recent publication in the journal Neurourology and Urodynamics highlighted the potential for Microdox® in the management of urinary tract infections, or UTIs, in children with neurogenic or non-neurogenic bladder dysfunction.¹
- 1. Singh G-K, Deshpande A, Schlegel G, Starkey M, Taghavi K. The rationale for bladder washouts in children with neurogenic bladder. Neurourol Urodyn. 2024;1-6. doi:10.1002/nau.25450

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Recent Business Developments

Continuously expanding our distributor network

- In January 2025, we partnered with WellSpring Pharmaceutical Corporation for the sale of our Microcyn technology-based products to large retailers in the United States.
 - In March 2025, we expanded our agreement with WellSpring to include additional consumer focused products.
- In August 2024, we announced a new distribution agreement with Medline Industries, LP for the marketing and distribution of our wound care products in the United States.
 - In October 2024, we expanded our agreement with Medline for marketing and distribution of our wound care products in Canada, and the sale of OTC wound care products to retailers in both countries.
- · In July 2024, we announced a new distribution agreement with Smart Healthcare Company for wound care in Ukraine.
- In April 2023, our partner Te Arai BioFarma Limited launched BabySoothe for diaper rash applications in Taiwan.
 - In April 2024, Te Arai launched Microdacyn for wound care in Taiwan.
- Our partner Microderm Technologies recently launched Dermodacyn for wound care applications in Thailand.
- Our partner Brill Pharma SL is now selling Sonoma's eye care products in Italy, Spain, Portugal and Germany.
- In January 2023, we entered into a distribution agreement with Daewoong Pharmaceutical Co., Ltd., one of the largest ph companies in South Korea, for marketing and distribution of Primocyn™ Skin Solution products

Amy Trombly CEO

Amy Trombly is our Chief Executive Officer and also serves on our Board of Directors. She counseled public companies for over two decades in corporate and securities law and mergers and acquisitions, including as the owner and manager of Trombly Business Law, PC. In her earlier career, Ms. Trombly was a Vice President at State Street Bank and Special Counsel at the U.S. Securities and Exchange Commission. Ms. Trombly is a member of the bar in Massachusetts and Colorado.

Management

Bruce Thornton COO

Bruce Thornton has served as our Chief Operating Officer, Vice President of Global Operations, and US General Manager since 2004, and currently as Executive Vice President and Chief Operating Officer. He served as Vice President of Operations for Jomed (formerly EndoSonic Corp.) from January 1999 to September 2003, and as Vice President of Manufacturing for Volcano Therapeutics, an international medical device company, following its acquisition of Jomed, until March 2004, Mr. Thornton received a B.S. in Aeronautical Science from Embry-Riddle Aeronautical University and an M.B.A. from National University. He also has served in the US Army.

Jerry Dvonch CFO

Mr. Dvonch serves as our Chief Financial Officer. He joined us from the SpineCenter Atlanta where he was the controller and Senior Vice President of Finance and Accounting since March 2017. From March 2016 to April 2016 he was a consultant controller for DS Healthcare Group, Inc. Prior to that he was the director for external reporting and director of finance of NeoGenomics Laboratories from July 2005 to July 2015. He has over 10 years of experience with SEC reporting. Mr. Dvonch is a licensed Certified Public Accountant in New York. He holds a Master of Business Administration in Finance from the University of of Rochester and a Bachelor of Business Administration in Accounting from Niagara University.



