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# Sonoma Pharmaceuticals to Exhibit at 2025 American Academy of Dermatology Annual Meeting in Orlando

**BOULDER, CO / [ACCESS Newswire](#) / February 25, 2025** / Sonoma Pharmaceuticals, Inc. (Nasdaq:SNOA), a global healthcare leader developing and producing patented Microcyn<sup>®</sup> technology based stabilized hypochlorous acid (HOCl) products for a wide range of applications, including wound care, eye, oral and nasal care, dermatological conditions, podiatry, and animal health care, today announced that it will be exhibiting at the 2025 American Academy of Dermatology (AAD) Annual Meeting in Orlando, Florida, March 7-11, 2025.

The AAD annual meeting is one of the leading dermatology conferences in the U.S., with over 350 exhibitors and close to 20,000 attendees, including more than 10,000 medical personnel, and more than 300 educational sessions. Founded in 1938 and headquartered in Rosemont, Illinois, AAD is committed to advancing the diagnosis and medical, surgical, and cosmetic treatment of the skin, hair, and nails, advocating high standards in clinical practice, education, and research in dermatology, and supporting and enhancing patient care for a lifetime of healthier skin, hair, and nails. Its annual meeting provides an ideal opportunity to connect with peers and industry leaders while learning about the latest advancements in dermatology.

Sonoma Pharmaceuticals is aiming to expand its distributor network and to showcase its line of prescription and OTC dermatology products. Interested parties are invited to visit Sonoma Pharmaceuticals' booth 1054 to learn about its patented, innovative, safe, and effective products for skin care, scar management and atopic dermatitis. For more information, please visit <https://sonomapharma.com>.

"We are excited to attend AAD again this year and to present our portfolio of dermatology products at this highly regarded event," said Amy Trombly, CEO of Sonoma. "We look forward to bridging new connections for the continued distribution of our products and the increased availability of our safe and effective technology in the U.S."

## **About Sonoma Pharmaceuticals, Inc.**

Sonoma Pharmaceuticals is a global healthcare leader for developing and producing stabilized hypochlorous acid (HOCl) 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 HOCl 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 and the company actively seeks new distribution partners. The company's principal office is in Boulder, Colorado, with manufacturing operations in Guadalajara, Mexico. European marketing and sales are headquartered in Roermond, Netherlands. More information can be found at [www.sonomapharma.com](http://www.sonomapharma.com). For partnership opportunities, please contact [busdev@sonomapharma.com](mailto:busdev@sonomapharma.com).

## **Forward-Looking Statements**

Except for historical information herein, matters set forth in this press release 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 (the "company"). These forward-looking statements are identified by the use of words such as "continue," "develop," "anticipate," "expect" and "expand," among others. Forward-looking statements in this press release 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 the COVID-19 pandemic and economic development, 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.

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