Mydrymouth.com Provides Resources For Women With Sjögren's Syndrome

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mydrymouth.com

(NAPSA)—Do you have persistent dry mouth? You may be one of an estimated four million people, 90 percent of whom are women, suffering from a little-known autoimmune disease called Sjögren's syndrome (SS). People with SS can experience symptoms that mimic other conditions, which makes diagnosis difficult—often taking up to 10 years.

Now there is a new Web site, www.mydrymouth.com, designed to educate you about SS. The Web site offers tools to help you recognize symptoms, tips to manage the disease, as well as information on potential treatment options.

Web site features include:

• Living with the Dry Mouth Symptoms of Sjögren's Syndrome offers tips for relief and information on how to recognize symptoms.

- Is Dryness a Problem for You? includes a brief questionnaire that can be completed and shared with your health care provider.
- Resources provides an additional outlet to visit to learn more about SS.
- Join the Moisture Network allows visitors to sign up for supplementary information.

"The Sjögren's Syndrome Foundation's mission is to raise public awareness," says Steven Taylor, Chief Executive Officer, Sjögren's Syndrome Foundation. "That's why we are so pleased that mydrymouth.com gives people a great way to get the facts about this condition and hopefully, get diagnosed sooner."

Visit www.mydrymouth.com today. The Web site is sponsored by Daiichi Sankyo, Inc.

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Note to Editors: This article was funded and reviewed by Daiichi Sankyo, Inc.