Sjogren’s World is...

...an internet community built for people who have Sjogren's Syndrome to meet and share their experiences and knowledge in order to improve quality of life and help thwart the loneliness and isolation that often accompanies living with this *rare* disease. Sjogren's World offers various ways to meet with others, including discussion forums and live chats as well as reliable medical articles and links.

For your use:

Read about [Who We Are](#). There you can also find information about Sjogren's World and also meet the Sjogren's World Staff.

**Sjogren's World Forums:** If you are seeking "someone like you," have general comments and concerns, or are asking for help, our multiple message boards with over 1,300 registered users and thousands of visitors are for you! Do you have a personal tip that you would like to share with others? How about some good news to share? A question about Sjogren's? Leave a message and before you know it, there will be comments from our users.

Visit our [Live Chat instructions](#) for information on where to go and how to chat live. If you already know, [click here to sign in and chat](#)! (Thurs Evenings)

**Neurological Connections** is the place to read articles and abstracts about the neurological problems associated with Sjögren's Syndrome.

**Useful Links** on the Forum is the place to find a collection of links to sites offering useful information regarding Sjogren's Syndrome, its diagnosis, treatment, and research. Please feel free to share any links you find that you feel will be helpful.

There is also a [Sjogren's World Book Club](#) board, where you can read and/or comment about Sjogren's related books.

All of us at Sjogren’s World Staff look forward to meeting you!

[Disclaimer](#)