ANNUAL REP@RT

The mission of the Sjögren's Society of Canada

To provide support and education for diagnosed and undiagnosed Sjögren's patients, to increase professional and public awareness, and to promote and fund Sjögren's related research.

IMPACT SNAPSHOT

138

508

9306

New members joined the SjSC in 2022

2022 Town Hall/National Patient Conference attendees

Visitors to our new website in 2022

Sjögren's Society

of Canada

A year of making strides for Sjögren's

Reviewing the SjSC's accomplishments in the past year, we are very proud of what we have achieved to support and educate Sjögren's patients in Canada through our Support Groups, Town Halls, National Patient Conference, and newsletters. We are particularly excited about the launch of our brand-new website. Members have access to updated disease information and can now manage all interactions with the SjSC through the website. Our continued focus on spreading the word about Sjögren's through our walks, brochures, website, and the Canadian Sjögren's Awareness social media campaign serve to educate the Canadian public about our unknown disease. The SjSC has emerged from the pandemic stronger than ever with solid increases in membership and fundraising, more in-person events, and plans to resume our research grant program.

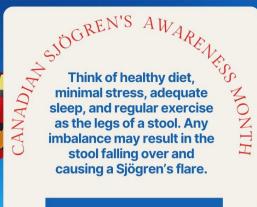
OBJECTIVES IMPACT

Launch New SjSC Website	 Increased capacity and functionality Up-to-date Sjögren's content Event Calendar with full schedule of upcoming events Easy membership registration process
Increase Awareness	 French Sjögren's brochures now available (by mail & online) Canadian Sjögren's Awareness Month held in July 2022 Daily Sjögren's 'tips' & info shared through social media
Expand Walks for Sjögren's	 New walks in Brockville, ON & London, ON 135 Donors/80+ Participants All walks were held in person in 2022 Calgary & Saskatchewan interested in walks
Enhance Fundraising Efforts	 Events include The Great Canadian Giving Challenge, Walks for Sjögren's, Giving Tuesday, Year End Giving Campaign 665 Individual Donors in 2022
Research	 Committed to 5% of all donations be allocated to research Over \$9,000 raised in 2022 (6.08%) Plans to resume our Summership Research Grants for medical and dental students in 2023
Advocacy	 Letter campaign and protest conducted for approval of Cevimeline/Sholyne in Canada Although the campaign was unsuccessful, Health Canada acknowledged that Sjögren's is a serious disease

TOTAL FUNDS RAISED

\$126,766.77





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WWW.SJOGRENSCANADA.ORG



'est-ce que c'est le Sjögre

s yeux secs, la bouche sèche, la fatigue s douleurs articulaires sont les symptôm aractéristiques de l'une des maladies aut nunes les plus répandues qui touche envi des Canadiens, dont 90% sont des femn



Sjögren's Society of Canada 1-888-558-0950 www.sjogrenscanada.org







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We couldn't have done it without your generous support. "I am so thankful for this organization! I have learned so much in 3 yrs, have found a better rheumatologist who understands the myriad manifestations of Sjögren's, and the Conferences are a wealth of information that has helped me in so many ways, including the knowledge that my symptoms are not in my head, and that there are so many other people out there who are in a similar life situation. Many, many thanks!" AM

"Many thanks, to you and the SjSC for all the great work in organizing and presenting the annual conference and being a great resource for all of us affected by Sjögren's." HY

"You are not alone, and it's okay to not be okay. There's a whole community of Sjögren's Warriors out there waiting to support you along your journey. The people I have met through the SjSC have not only become my friends but also have become part of my support system." CM

THANK YOU

to all our donors and volunteers.

Special thanks go to Lauren Vording (Managing Director), Dr. Leslie Laing (President), Mary McNeil (Vice-President), Marc Foisy (Treasurer), Christin McGinn, Ellen Wang & Julie Yoshikuni (Board of Directors), our Medical Advisory Board and all of our incredible Support Group Leaders.



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