Mission Statement

Narcolepsy Network is dedicated to improving the lives of men, women, and children with narcolepsy. The organization’s goals include increasing public awareness to foster early diagnoses, advocating for all people with narcolepsy (PWNs), promoting and supporting narcolepsy and related research, and providing education and resources both to people living with narcolepsy and the public at large.

A Brief History of Narcolepsy Network

Narcolepsy Network emerged from the vision of 13 women from self-help groups across the United States —Mary Babcock (NE), Ruth Justice Nebus (NJ), Norma J. Potter (FL), Nicole L. Kephart (OR), Ann Matthes (MA), Eve Davis (NH), Sue Brockway Carella (CA), Niss H. Ryan (NY), Clair S. Sassin (DC), Jan Wright (CA), Violet Baumann, Blanche Baechalin, and Lorraine Baird (NJ).

In 1984, most of these women met for the first time in Akron, Ohio. All were members of the American Narcolepsy Association (ANA), a group that no longer exists today. They originally convened as a task force charged with resolving ANA member issues, but realized those issues were intractable, and instead formed a new organization.

They elected Mary Babcock provisional President, decided on the name “Network”, and began outlining the new organization's programs. Demonstrating the power of REM sleep, committee member Jan Wright awoke from a short nap inspired with the new group's motto: “CARE,” standing for COMMUNICATION, ADVOCACY, RESEARCH, and EDUCATION.

In 1986, the team met in Shrewsbury, NJ, completed the new organization's Articles of Incorporation and Bylaws, appointed Ruth Justice Nebus as its incorporator, and elected Niss H. Ryan, Sue Brockway Carella, Jan Wright, Norma J. Potter, and Ruth Justice Nebus as its founders. They incorporated the organization that spring, and later held the first Narcolepsy Network Annual Meeting and election of officers in May. Approximately 75 members attended that first meeting.

From its first year, Narcolepsy Network opened communications with the National Institutes of Health, the few sleep disorders centers then in existence, and other key health and science leaders. Understanding and supporting relevant research, advocating for PWNs, and building the organization were foundational goals.

By 1988, Narcolepsy Network's New York City conference drew 200 attendees. There, Dr. John Holloman received the first William C. Dement Research Award, and members staged a “Nap-In” in Central Park, which gained national attention and coverage on CNN. Members crisscrossed the country in the “Wake Up America” caravan as part of a concerted effort to raise both public and Congressional awareness about sleep disorders. This 12,000-mile journey bore fruit in 1993 when President Clinton signed The National Sleep Disorders Research Advisory Board into law.

By 1996, after ten years as an all-volunteer organization, Narcolepsy Network hired professional staff, including its first Executive Director. This allowed for further advocacy, organizational development, and physician outreach at such venues as the American Academy of Neurology and the National Science Foundation. Membership drives, health provider education, and fundraising efforts also became better coordinated.

What does the future hold for Narcolepsy Network? Since our founding, we have seen major advances in research and treatments. Technology has vastly increased access to information on symptoms, diagnosis, treatments, research, resources, and more. Our membership has grown significantly from its original 79. As newly diagnosed join our organization, they bring with them the energy and creativity to advance Narcolepsy Network and its goals into the future.
As we all know, 2020 was a tough year. It was full of challenges that many of us never expected to encounter. The impact of the COVID-19 pandemic has been felt in many ways. It’s disrupted our daily lives, as well as our emotional, mental, and physical well-being. Before the pandemic took hold in the United States, Narcolepsy Network was able to hold two in-person events. The first was a One-Day Seminar in Florida on February 22nd, where we presented a mix of educational and clinical updates. In March, we brought 23 narcolepsy advocates to Capitol Hill as part of Rare Disease Week. Almost immediately following this event, life in the U.S. took a sharp turn as the first series of lockdowns went into effect in major cities.

When the pandemic hit, we realized that the most impactful thing we could do was keep our community connected together. People with narcolepsy need support from each other, and when we collectively lost the ability to do so it made a difficult situation even harder. In response to this new situation, Narcolepsy Network quickly pivoted its focus to building an online experience for our members.

First we created a series of Virtual Support Groups, which were so successful that we will continue facilitating them long after the pandemic is over. One might consider this a small silver lining—we all learned how to connect together in new virtual ways. Doing so has allowed us to make new friends in ways that were never before possible.

Next, we transformed our in-person events into a series of Virtual Patient Education Days, Virtual Mini-Conferences, and introduced a new program for youth called CONNECT. The 2nd annual World Narcolepsy Day (which was already a virtual, distributed event) was a smashing success given the circumstances. Ultimately as a community we all learned that we can support, educate, and advocate for one another in brand new ways.

Despite the significant challenges of 2020, we continued to deliver on our commitment to support, educate, and advocate for this community. As Narcolepsy Network moves into new, uncharted territory, we look to the future with fresh eyes around how we can best serve this community.

—Keith Harper, Board President
Going back as far as kindergarten, Brandon suffered from migraines. He knew it wasn’t normal to be in pain all the time. He was also very tired, but a sleep disorder wasn’t on anyone’s radar. Even though he had some of the best access to doctors in different specialties, first getting a diagnosis for migraines was difficult, but that still didn’t account for how tired he always felt. His narcolepsy diagnosis would later be confirmed with a sleep study.

Brandon received his narcolepsy diagnosis at age 12. It took many years to find the proper diagnosis and a treatment plan that worked for him. He adds, “I really loved school and I still do…but I was taken away from those avenues because I had to be out of school.” Between the tiredness, the headaches, and the other health issues, it was debilitating. He wanted to keep up with his love of sports, but couldn’t because he lacked the stamina to compete. His teachers didn’t understand – they thought he just wasn’t bright, or that he might have a learning disability. Even though Brandon was able to find an outlet in theater, performing, and signing as a way to cope with his exhaustion and pain, he was only able to go to school on a part time basis, and as a result, he became isolated from his peers.

Fortunately, Brandon’s parents advocated for him, as well as certain teachers and coaches, and even doctors, so by his sophomore year in high school, he finally figured out what worked for him – he knew how to pursue a demanding academic schedule, manage his creative outlets, all while dealing with his narcolepsy symptoms. Two years ago, he was able to go back to school full time. He also explains that he did ask for accommodations and these helped with being successful in school. “I definitely did seek out accommodations…and they have helped immensely. Things would have been much harder if I had to go to school without them.”

Even though going through his narcolepsy journey to diagnosis was long, he still views himself as one of the lucky ones who was able to find some sort of relief through medications and this allowed him to fulfill his dreams and keep on dreaming. He adds, “It may be cliché, but I [got involved with] Narcolepsy Network to improve lives. If people with narcolepsy and rare diseases don’t have to go through the challenges I did, I feel the sense of responsibility to [help others].” In 2019, Brandon still wanted to help others so he applied to be part of the 2019 class of Youth Ambassadors. He also found solace in the other Youth Ambassadors he met. He says “I felt a connection to my peers through hearing their experiences. I knew I wasn’t alone.”

As a Youth Ambassador, Brandon has been able to raise awareness of sleep disorders locally and nationally at schools and conferences, spoke with doctors who want to learn more about what it’s like to live with narcolepsy, met other kids and teens who may just be starting on their journey, and attended Rare Disease Week on Capitol Hill to advocate for rare diseases. He hopes that by sharing his story, he can help someone else get an accurate diagnosis. He goes on to say, “By sharing my experiences with an often-misunderstood sleep disorder, I know I can help others recognize the symptoms of narcolepsy. I can also help teachers and others [recognize] the psychosocial impact of narcolepsy’s symptoms.” He was also involved with the CONNECT program that helped bring together kids with narcolepsy and their parents in an online format. He wanted to get involved with this program because, as he says, “I would have wanted someone who had narcolepsy to be there for me.” Additionally, he learned so much from the kids. He explains, “I recognized myself in these kids [and I told them] a lot of overcoming narcolepsy is taking the belief over the anxiety of telling people you have it and being comfortable with letting them know so you can succeed in life.”

When asked what he would say to other kids who may feel overwhelmed with their disorder he responded with, “They are tremendously brave to be going through this because they are part of a rare population. They have a friend in me and in Narcolepsy Network and whenever they feel scared or alone, they have someone to tell them to never give up. If one dream doesn’t work, create another dream.”

Brandon’s narcolepsy has allowed him to see the world through a different lens. With his experience in the rare disease community, he hopes that someday, he will be able to manage a major bio-pharmaceutical company and has aspirations in patent law.

“They have a friend in me and in Narcolepsy Network and whenever they feel scared or alone, they have someone to tell them to never give up. If one dream doesn’t work, create another dream.”
A Reason to Give: Christen

Christen was diagnosed with narcolepsy in the 5th grade. She was fortunate to be diagnosed early and was able to avoid the typical stereotypes that come along with the pre-diagnosis phase, including being called lazy, a label that Christen acknowledges is detrimental.

For many people with narcolepsy, a quick diagnosis usually doesn’t happen in a narcolepsy journey, but it just so happened that Christen had to undergo brain surgery and felt very tired post-surgery. She thought this was from her recovery, but would also experience cataplexy, a symptom usually only associated with narcolepsy. A doctor recognized her symptoms and sent her to a sleep lab to undergo further studies. In addition to cataplexy, Christen was also experiencing excessive daytime sleepiness, hallucinations, and would rely on naps to help her get her through the day. Additionally, she was also diagnosed with epilepsy and had to navigate what regimen would help her not only stay awake during the day, but also would keep her seizure free.

Luckily, Christen’s parents were advocates for her during her schooling and helped her get what she needed to succeed. She was determined to get her college degree and she succeeded. She added, “School may have taken me longer, but I was determined to get my degree.”

Fast forward to 2013, Christen attended an epilepsy support group. There she found a sense of community. She explains, “I really enjoyed connecting with others; getting peer support that I couldn’t find with my family.” Christen added, “Even though I have an extremely supportive and loving family, they don’t know what it’s like to have a seizure.” This meeting inspired her to want to find the same sense of togetherness for the narcolepsy community. She saw how beneficial a support group could be, so she decided to form her own narcolepsy support group.

Through attending the epilepsy support group, Christen realized that one of the ways to have a successful support group is to be a good leader and she was determined to be one. She came up with guidelines for the group and would show up to meetings regardless if no one else came. She also manages the group’s meetup.com page, Facebook page, and personally sends out a welcome email to individuals just going the group. She personally reaches out to new group members because, she says, “about half of them have never met anyone with narcolepsy. Before COVID, I would meet new group members in-person, but now it’s online. This is just another way to connect with a person who understands what the other person is going through.”

While Christen doesn’t push people one way or another with how they should be treating their disorder, she does bring up topics for the group to discuss. They may include: finding a sleep doctor, and not just seeing a primary care doctor for their sleep issues, dietary changes and if a gluten free diet may help, medications and how they can affect the overall quality of a person’s well-being, exercise, and nap schedules. Christen also tries to keep things upbeat at meetings. She usually starts off by asking people to voice something positive that happened during the day. She adds, “This isn’t a bunch of people complaining. It’s a bunch of people relating.”

Christen’s group usually gathers about 7-12 people and also invites people with idiopathic hypersomnia to join as well. Meetings alternate between Saturdays and weekdays so people can join without worrying about work conflicts. Some meetings have set topics, like a specific IH meeting or a friends and family meeting, while others are more organic to what will be discussed. The group also invites speakers such as doctors to talk about medications and clinical trials, a lawyer to speak about disability, and even informational discussions about mindfulness.

Sometimes Christen asks the group to bring tips that help individuals live their day-to-day life. She says, “I learn from the group too! For example, timing a nap is beneficial. Before I would sleep as long as I wanted, but it’s helpful to limit my nap time.”

Ultimately, for Christen, being a leader is part of her life’s purpose. She explains, “If I can help just one person, it’s all worth it.”

For Christen’s next mission, she hopes to talk with teachers or other school professionals who can reach students while they may be in their pre-diagnosis phase.

“If I can help just one person, it’s all worth it.”
2020 Highlights

One-Day Seminar in St. Petersburg, FL

We had our first 2020 in-person event in St. Petersburg, FL on February 22, 2020. We purposely kept this seminar small and intimate compared to our annual in-person conference. A week before the event, we had a surge in registrations and were at capacity days before the seminar. Dr. Bobbi Hopkins, medical director of John Hopkins at All Children’s Sleep Center, was the keynote speaker. There was also a psychosocial presentation by Dr. William Frye, a narcolepsy treatment update, a presentation by Youth Ambassador Dayyan Lee, and pharmaceutical updates.

Rare Disease Week

At the end of February, 900 rare disease advocates traveled from across the country to join forces during Rare Disease Week on Capitol Hill, which was held on February 22-27. Advocates participated in 393 meetings with their Members of Congress. In 2020, Narcolepsy Network sent 23 advocates to Washington, DC to be a voice for the narcolepsy community. Every year, the Rare Disease Legislative Advocates (RDLA), in coordination with the Everylife Foundation for Rare Diseases, hosts a free week-long event that brings together rare disease members to be educated on legislative issues, meet other advocates, and share perspectives with legislators.

One narcolepsy advocate, Bianca Lara, said this about her experience, “It’s a rare opportunity to dialogue directly with legislators who are creating legislation that will impact us all and be a part of that process through feedback and lobbying sessions. Individually we are few, but together we are many. United, we are a strong voice of advocacy.”

Suddenly Sleepy Saturday

We once again asked the narcolepsy community to submit proclamations to their governor, mayor, or local official declaring Suddenly Sleepy Saturday (Narcolepsy Awareness Day) on March 7, 2020. Supporters in 23 US states requested proclamations. Governors in 19 states approved the request and issued proclamations, so did mayors in 11 US cities.
2020 Highlights

National Virtual Support Group Meetings

As states began to shut down due to COVID-19, we had to cancel in-person gatherings, including support group meetings. Just because we had to cancel these meetings, it did not mean that people did not need our support. On the contrary, in these times of uncertainty, it was even more important that we came together. Since we could not meet in-person, we started to host Virtual Support Group Meetings via Zoom. We hosted our first virtual support group on March 26, 2020 and have hosted a virtual group meeting nearly every Thursday evening since. Many who were unable to attend in-person meetings due to time or distance constraints were able to attend a weekly meeting and make connections. One attendee said, “The support group reminds me that I am not alone, while also affording me the knowledge and awareness that I need in order to be mindful of each PWN’s unique experience.”

Virtual Patient Education Days, CONNECT Program & Virtual Mini-Conference Series

Even though we could not gather in-person, we still wanted to provide virtual programming to educate and connect the narcolepsy community.

We kicked off our Virtual Patient Education Day series on May 30. In total, we held three webinars in this series. Topics included psychosocial effects on narcolepsy, youth with narcolepsy, advocacy, and telling your story.

Another virtual program was the CONNECT program, which invited children, ages 8-13, who have been diagnosed with narcolepsy and their parent to join us online for a time of support, caring, and fun.

Lastly, the Virtual Mini-Conferences presented topics that normally would have been presented at our annual conference over the course of several webinars. These topics included applying for disability, workplace accommodations, and medication updates.

World Narcolepsy Day

We partnered with 26 national and international organizations to celebrate World Narcolepsy Day on September 22. We were able to share content from the narcolepsy community all day long including 19 stories, 1 poem, and two videos featuring an interview with Ginger Zee, PWN and Chief Meteorologist at ABC News.
Thank You 2020 Donors!

Narcolepsy Network thanks the following donors, whose generous gifts made it possible for us to provide information and support to thousands of people living with narcolepsy and their families. We also thank those not listed who volunteered their time and energy to help further Narcolepsy Network’s mission.

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Covington, KY  
**Seattle Children’s Sleep Disorder Center**  
Bellevue, Washington  
**South Florida Sleep Diagnostic Center**  
Miami, FL  
**University of Michigan Sleep Disorders Center**  
Ann Arbor, MI

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Lakeland, FL

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University of Michigan Sleep Disorders Center  
Ann Arbor, MI  
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St. Francis Sleep, Allergy & Lung Institute  
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Medical University of SC  
Charleston, SC

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**Narcolepsy Network Leadership**

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<th>Name</th>
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<tbody>
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<td>St. Petersburg, FL</td>
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<td>Cynthia Zorn, Vice President</td>
<td>San Diego, CA</td>
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<td>Mee Warren, Treasurer</td>
<td>New York, NY</td>
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<td>Stephanie Fischer, Secretary</td>
<td>Bensalem, PA</td>
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<td>Keith Harper</td>
<td>New York, NY</td>
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<td>Sharon O’Shaughnessy, MA, SLP</td>
<td>Downington, PA</td>
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<tr>
<td>Paul Reynolds</td>
<td>New York, NY</td>
</tr>
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<td>Rajeev Sachdeva</td>
<td>East Windsor, NJ</td>
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<tr>
<td>Carol Arnette</td>
<td>Williamsburg, VA</td>
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<tr>
<td>Emily Kreuziger</td>
<td>Juneau, WI</td>
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<tr>
<td>Niss Ryan (Trustee Emerita)</td>
<td>Rochester, NY</td>
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<tr>
<td>Emmanuel Mignot, MD, PhD Chair</td>
<td>Stanford Center for Narcolepsy Palo Alto, CA</td>
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<td>Stephen A. Amira, PhD</td>
<td>Brookline, MA</td>
</tr>
<tr>
<td>Meeta Goswami, MPH, PhD</td>
<td>Narcolepsy Institute Bronx, NY</td>
</tr>
<tr>
<td>Lois Krahm, MD</td>
<td>Mayo Clinic Scottsdale, AZ</td>
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<tr>
<td>J. Gila Lindsley, PhD</td>
<td>Lexington, MA</td>
</tr>
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<td>Northwestern University Feinberg School of Medicine Chicago, IL</td>
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<tr>
<td>Quentin Regestein, MD</td>
<td>Brigham &amp; Women’s Hospital Boston, MA</td>
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<tr>
<td>Chad M. Ruoff, MD</td>
<td>Mayo Clinic, Scottsdale AZ</td>
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<tr>
<td>David B. Rye, MD, PhD</td>
<td>Emory University School of Medicine Atlanta, GA</td>
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<tr>
<td>Thomas E. Scammell, MD</td>
<td>Beth Israel Deaconness Medical Center Boston, MA</td>
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<tr>
<td><strong>Lawrence Scrima, PhD</strong></td>
<td>Sleep-Alertness Disorders Center, Inc. Aurora, CO</td>
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<tr>
<td>Jerome Siegel, PhD</td>
<td>UCLA Neurobiology Research Sepulveda, CA</td>
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<tr>
<td>Michael J. Thorpy, MD</td>
<td>Sleep-Wake Disorders Center Bronx, NY</td>
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<td>Agnes Kenny, MD</td>
<td>Peru, IN</td>
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<tr>
<td>Suresh Kotagal, MD</td>
<td>Rochester, MN</td>
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<tr>
<td>Suzanne Moore-Darms</td>
<td>Marlton, NJ</td>
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<tr>
<td>Michael Eig, Esq</td>
<td>Chevy Chase, MD</td>
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**Mali Einen**  
Menlo Park, CA

**Michele Profeta**  
Atlanta, GA

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Board of Directors videoconference calls were held in:

- January
- February
- March
- April
- May
- June
- July
- August
- September
- November
- December
This past year was unprecedented to say the least. We experienced shutdowns, virtual school, and quarantines. While we were saddened to have to cancel our in-person events, we were excited to present educational and informational programs in a new format through our online patient days and our virtual mini-conferences.

As we all tried to adjust to everything happening around us, more people felt isolated and alone. That meant making connections and having a sense of community was still needed. We saw this need and held virtual support groups, virtual meetings for teens and young adults, and virtual gatherings for kids and their parents.

We thank everyone for your continued support. You allowed us to reach more people in new ways as we all navigated this new normal.

Warm Regards,

Paul Reynolds  
Treasurer  
Narcolepsy Network Board of Directors

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**Statement of Financial Position - As of December 31**

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<td>Unrestricted</td>
<td>$422,881</td>
<td>$402,134</td>
</tr>
<tr>
<td>Temporarily restricted</td>
<td>258,336</td>
<td>96,192</td>
</tr>
<tr>
<td><strong>Total net assets</strong></td>
<td>$681,217</td>
<td>$498,326</td>
</tr>
</tbody>
</table>

| **Total liabilities and net assets** | $703,473 | $534,942 |

**Statement of Activities - Years ended December 31**

<table>
<thead>
<tr>
<th></th>
<th>2020</th>
<th>2019</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Income</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Corporate Grants</td>
<td>$303,995</td>
<td>$277,500</td>
</tr>
<tr>
<td>Conference</td>
<td>1,375</td>
<td>72,908</td>
</tr>
<tr>
<td>Contributions</td>
<td>111,679</td>
<td>197,121</td>
</tr>
<tr>
<td>Membership Dues</td>
<td>36,105</td>
<td>36,734</td>
</tr>
<tr>
<td>Other</td>
<td>34,773</td>
<td>19,266</td>
</tr>
<tr>
<td><strong>Total Income</strong></td>
<td>$603,927</td>
<td>$487,927</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th></th>
<th>2020</th>
<th>2019</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Expenses</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Programs</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Conference and Seminars</td>
<td>$57,932</td>
<td>$205,781</td>
</tr>
<tr>
<td>Public Information</td>
<td>91,223</td>
<td>80,432</td>
</tr>
<tr>
<td>Membership</td>
<td>67,614</td>
<td>63,131</td>
</tr>
<tr>
<td><strong>Total Program Expense</strong></td>
<td>$216,769</td>
<td>$349,344</td>
</tr>
<tr>
<td>General and administrative</td>
<td>61,118</td>
<td>65,081</td>
</tr>
<tr>
<td>Fundraising</td>
<td>27,149</td>
<td>32,918</td>
</tr>
<tr>
<td><strong>Total Expenses</strong></td>
<td>$305,036</td>
<td>$447,343</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th></th>
<th>2020</th>
<th>2019</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Net increase (decrease) in net assets</strong></td>
<td>$271,158</td>
<td>$156,186</td>
</tr>
</tbody>
</table>

**Five Year Revenue & Expense Comparison**

<table>
<thead>
<tr>
<th>Year</th>
<th>Income</th>
<th>Expense</th>
</tr>
</thead>
<tbody>
<tr>
<td>2015</td>
<td>$310,175</td>
<td>$362,000</td>
</tr>
<tr>
<td>2016</td>
<td>$335,527</td>
<td>$438,000</td>
</tr>
<tr>
<td>2017</td>
<td>$403,000</td>
<td>$396,000</td>
</tr>
<tr>
<td>2018</td>
<td>$421,500</td>
<td>$423,000</td>
</tr>
<tr>
<td>2019</td>
<td>$482,763</td>
<td>$487,927</td>
</tr>
</tbody>
</table>

**2020 Revenue Categories**

- National Conference 1%
- Contributions 23%
- Corporate Grants 62%
- Membership Dues 7%
- Other 7%

**2020 Expense Categories**

- Conference & Seminars 19%
- Public Information 30%
- Membership 22%
- General & Administrative 20%
- Fundraising 9%
Donating to Narcolepsy Network

We express our sincere appreciation to all those who support Narcolepsy Network’s mission. To make a donation, visit www.narcolepsynetwork.org. Please consult your lawyer for additional information on planned giving through a will, trust, or estate. Narcolepsy Network is a tax-exempt 501(c)(3) nonprofit organization. Your contributions are tax-deductible to the extent permitted by law. Consult your tax advisor for details.

Other Ways to Support Us

Shop Til You Drop While Raising Funds!

If you do a lot of your shopping online, you can shop and help out Narcolepsy Network at the same time. Just remember to go to www.smile.amazon.com for your online shopping. When a purchase is made through the Amazon Smile website, the company donates a portion of the purchase price of items you buy to an organization of your choice, like Narcolepsy Network. It’s just like shopping on the regular Amazon site, but we benefit. If you are new to Amazon Smile, the website will ask you to pick an organization when you first create your account and you can select “Narcolepsy Network” at that point. For returning users, you can select Narcolepsy Network by going to Your Account and selecting Change Your Charity.

Create a Fundraising Campaign on Facebook

Are you thinking about doing something special for your birthday or other special event? How about create a fundraising campaign on Facebook where you can ask your friends to support your cause? To start a fundraiser, click Fundraisers in the left menu of your News Feed, then click Raise Money. Select Narcolepsy Network as your charitable organization, choose a cover photo, and fill in the details. Then click Create. That’s it!

Narcolepsy Network Staff

Eveline Honig
Executive Director

Christine Hackenbruck
Executive Assistant

Kristin Fisher
Communications Coordinator

Michelle Doherty
Accountant