No one should face MS alone. The National MS Society is here so that no one has to.

8.48 MILLION people engage with the Society
FY23 target: 8.6 million people

299,212 people connected with the Society for the first time
FY23 target: 405,000

62.6%
572,032 of the estimated 914,000 people with MS in the United States are connected to their National MS Society
FY23 target: 62.4%

We bring our entire MS community together so everyone feels supported by their National MS Society.

The third Black MS Experience Summit took place in June, continuing to deepen connections with and support the Black community living with MS.

- Registration for the summit increased by 148% over last year
- 97% of attendees feel the Society is a trusted source of support
- 85% feel more connected to the Black MS community

“I walked away from this summit with the inspiration to keep going. When I was first diagnosed with MS as a black man, I felt alone. This summit allowed people to connect with others to feel empowered and no longer invisible.”
— Akeem R. A. McMichael

The National MS Society works to increase access to personalized, affordable, high-quality healthcare and MS medications.

With the U.S. Department of Health & Human Services’ Office of Minority Health, the Society provided two Ask an MS Expert programs during Minority Health Month, which were viewed more than 2,400 times.

People relying on power wheelchairs have more access to seat elevation devices to enable safe transfers. The life-changing decision for the Centers for Medicare & Medicaid Services to cover the cost came following evidence on medical necessity provided by the Society.

CMS also revised their Medicare Drug Price Negotiation Program guidance document for 2026, accepting two changes proposed by the Society: establishing listening sessions to inform the negotiations process and implementing patient protections to ensure access to all drugs selected for negotiation.

The Society’s MS clinical fellowship program trained 29 neurologists or physiatrists to increase access to high-quality MS care. An estimated 4,000 people with MS will benefit from specialized MS care.

The Society’s National Medical Advisory Committee released, “What to Expect from Rehabilitation Care: A Guide for People with MS,” which explains the roles of the rehab team and the importance of early treatment, and outlines what to expect from a rehab evaluation. The guide has been translated into Spanish, and was shared with 2,100 healthcare providers.

Our summer issue of Momentum highlighted the voices and experiences of transgender people living with MS.
We are bringing the world together to cure MS for every single person — as fast as possible.

The first-ever Pathways to Cures Global Summit was held in May, convening nearly 200 participants from 15 countries, including leadership from MS advocacy organizations, researchers, doctors, government funders, pharmaceutical companies, supporters and people living with MS. As a result of this meeting, the Pathways to Cures Roadmap will be revised to include the most recent scientific knowledge, and a scientific strategy group will create an MS research landscape analysis to identify opportunities for further global collaboration.

“Bringing [a group like] this together in one meeting has never been done before. It is hugely ambitious, but without ambition, you don’t achieve. This is a really great step forward.”
— Alan Thompson, MD, Summit Co-Chair

Reaching a world free of MS will take all of us — our time, our energy, our resources. Together is the only way forward.

FY23 Net Revenue

<table>
<thead>
<tr>
<th>FY23 Target:</th>
<th>$165.6M</th>
</tr>
</thead>
<tbody>
<tr>
<td>FY23 Year-to-Date:</td>
<td>$118.9M*</td>
</tr>
<tr>
<td>Q3 $42.5M</td>
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</tr>
<tr>
<td>Q2 $36.5M</td>
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<tr>
<td>Q1 $39.9M</td>
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FY23 Gross Revenue to Date*

<table>
<thead>
<tr>
<th>Bike Revenue</th>
<th>Walk Revenue</th>
<th>Other Events**</th>
<th>IG &amp; Direct Marketing</th>
<th>Bequests</th>
<th>Corporate Giving &amp; Central Sponsorships</th>
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<tbody>
<tr>
<td>$53.0</td>
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<td>$8.5</td>
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</table>

*Excludes Other Income

**Includes Leadership Events, Finish MS, DIY Fundraising MS, Climb to the Top MS, Challenge Walk MS and other events

A team of Society funded researchers published an update on the **prevalence of MS in the U.S.** and recently released a report that confirms anyone can develop MS, regardless of age, race, ethnicity or sex assigned at birth.

* Green light represents excellent progress toward implementation milestones