

This holiday season, give the gift of hope to families affected by MPS. When our journey began in 1984, children with MPS faced a devastating reality: many went undiagnosed, and those who were diagnosed had heartbreakingly short life expectancies. Thanks to a community of supporters like you, we've made incredible strides, bringing improved diagnoses, treatments, and support to countless families. Yet, our work is far from complete—there is still no cure, and countless families continue to face the difficult realities of living with MPS.

Every dollar you contribute to our Annual Fundraiser gives Canadian families facing MPS the precious gift of hope by:

- Supporting families as they navigate complex medical journeys
- Providing resources for earlier diagnoses and better treatment
- Offering hope and strength to those confronting these rare, challenging disorders.

www.mpssociety.ca/donate

This season, your generosity can bring a brighter future to families affected by MPS.

Thank you for your support.

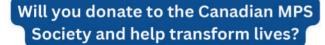
Yes!! I want to support families living with MPS & related lysosomal storage diseases and help advance research for a cure. \$25 \$50 \$100 \$250 I prefer to give \$	Canadian MPS Society for Mucopolysaccharide & Related Diseases You will receive a charitable tax receipt for the full amount of your donation.
Please charge my: Visa Mastercard Amex	I do not wish to have my name appear in your recognition materials.
CARD NUMBER EXPIRY	I wish to subscribe to your monthly e-newsletters
CARDHOLDER NAME CVV	Email address:
SIGNATURE NAME OF DONOR: ADDRESS:	If you no longer wish to be contacted by the Society and wish to have your name removed from our lists, please email us on info@mpssociety.ca
CITY: POSTAL CODE: TELEPHONE: EMAIL:	#218-2055 Commercial Drive, Vancouver, B.C. V5N OC7 Phone Toll Free: 1-(800)-667-1846 or: 1-(604) 924-5130 Registered Charity #12903 0409 RR0001



YEARS STRONG

1984-2024

www.mpssociety.ca



"Thanks For Giving Me a New Life!"

Meet Abia Sheraz, a young girl with MPS IVA from Karachi, Pakistan. Abia's family spent years worrying about accessing medical treatment which was not available in Pakistan. When in July 2022 they reached out to the Canadian MPS Society in desperation, they found hope. With quick responses and unwavering support, the Society guided the family through every step of their journey. In September 2024, Abia finally arrived to Canada, and is on her way to accessing vital treatments and supports. Her story exemplifies the life-changing impact of your donations.

Keeping Emmett Safe

"We received a Wonderfold Wagon for Emmett, through the Financial Assistance funding. This was so helpful as these wagons are so expensive and we were looking for a safe alternative to our stroller to walk my daughter to and from school. Emmett loves to wait on the playground for his big sister and it was such a great way to keep him safe on the walk as it has seat belts and could fit my daughter as well. Thank you!!"

- Steph, mother of Emmett (MPS IIIA)

Mobility is a Game Changer

"Simon is so grateful to the MPS Society for helping pitch in for the cost of his snazzy new wheelchair that is designed to last! He just recovered from a double hip osteotomy and is using a mix of push bike, wheelchair and walker to get around. Having high quality mobility equipment is a game changer for this active and energetic kid!" - Becs, mother of Simon (MPS IVA)





Give the Gift of Hope.

Together, we can create a brighter future where no child's life is limited by MPS.