



Dear Family and Friends,

We are so excited to be a part of the National Organization for Disorders of the Corpus Callosum (NODCC)! This organization is expressly designed to help individuals living with a corpus callosum disorder. The corpus callosum is the largest connective pathway in the brain (more than 200 million nerve fibers!) and it transfers information between the hemispheres. People with corpus callosum disorders were born without this huge brain structure, with only part of it, or with severe damage to it. As you can guess, corpus callosum disorders present many challenges!

Today there are many physicians, researchers, therapists and educators dedicating their life's work to understanding disorders of the corpus callosum. Even though much of the research in this area is new, it is reassuring to see all of the hard work and progress that is being made. We've heard parents with children – all affected by the same challenges – eagerly tell their stories and share what has helped them cope on a daily basis. It is so comforting to be surrounded by a community that understands our frustrations and shares our desire to help these children in every way possible to develop and thrive.

To help the NODCC continue its great work, we hope you will consider joining us in supporting the NODCC. Monetary contributions of \$25, \$50, \$100 or any other amount will impact the lives of children, teens, adults and families affected by callosal disorders. No amount is too small to make a difference. As a 503 (c)(3) nonprofit organization, your gifts are tax deductible and can be made anytime throughout the year.

For more information on how the NODCC is enhancing lives or to make an online donation, visit www.nodcc.org. We'd love to see the NODCC become your charity of choice and thank you for facing the challenges with us.

Thank you for your wonderful support and generosity!

Best wishes,

Understanding the conditions... enhancing lives