

**Sample of a fundraising appeal letter you can share on social media, email, or use for a regular mailing.**



Dear Family and Friends,

Thank you so much for all of your love, support, and prayers over the last few months/years. We are writing you to ask for your support for the neurofibromatosis cause. As you may know, our daughter Jane was diagnosed with neurofibromatosis type 1 or NF1 this past January.

Her doctors found what they consider a low-grade brain tumor on her optic nerve as a result of this disorder. Jane has been undergoing chemotherapy treatment and will continue over the course of the year in hopes to stabilize the tumor's growth to keep her eyesight intact. She also undergoes MRIs every three months to note the progression of the tumor. She is a true inspiration and shows such bravery and strength at such a young age!

Neurofibromatosis is set a genetic disorders of the nervous system that causes that growth of tumors to form on nerves anywhere in the body. NF affects 1 in 2,000 people worldwide. If can cause many symptoms and affects each person differently. Currently, there is no cure for this disorder.

As everyone with NF does, we are learning to live day by day. We are hoping for a future that will bring us new opportunities for treatment and eventually a cure for this disorder. This is where you, our family and friends come in. You play an important role in helping us to get that much closer to this goal.

In honor of Jane, we have for a team called "Join for Jane". We will be honoring her by walking for a cure in the Walk4NF event on Saturday, May 18, 2024 at the White River State Park in Indianapolis, IN. We are walking to help raise awareness about neurofibromatosis (NF) and to raise funds to support NF research.

NF Midwest is the local regional non-profit that C.A.R.E.S. to help improve the lives of people with NF. By donating to my page, YOU are helping in this mission of C.A.R.E.S.—Clinics, Awareness, Research, Education, Support.

Thank you so much for your generosity and helping us "Join for Jane" to raise NF awareness in the hopes to finding a cure! Please remember that no gift/donation is too small! Feel free to forward this to anyone who may wish to donate to the cause as well! Thank you from the bottom of our hearts.

With heartfelt gratitude,

Joe and Jean Smith