The Children’s Tumor Foundation supports individuals of all ages

The Children’s Tumor Foundation (CTF) is a non-profit organization dedicated to improving the health and well-being of children and adults affected by neurofibromatosis (NF), including neurofibromatosis type 1 (NF1), neurofibromatosis type 2 (NF2), and schwannomatosis.

Our Mission
Drive research, expand knowledge, and advance care for the NF community.

Our Vision
End NF.

We are called the Children’s Tumor Foundation because NF is most often diagnosed in childhood, however NF is a lifelong condition, and CTF is committed to finding treatments for all who live with NF, young and old.

There are many ways that CTF supports adults with NF:

• Organizes a social gathering for adults with NF at the annual NF Family Forum
• Hosts an “Adults with NF” Facebook group, moderated by a trained CTF volunteer
• Highlights stories of adults with NF in our “This is NF” photo series
• Answers questions from adults in our “Ask Kate!” video series
• Holds a camp for young adults through 25 years of age
• Distributes and updates CTF publications
• Provides a searchable feature on our NF Clinic listing to easily find existing NF adult health care providers
• Creates events and programs for parents of children with NF
• Offers courses in becoming a Patient Advocate
• Promotes awareness of the NF1 Adult Care guidelines
• Supports health care professionals in staying current with the latest in clinical research and patient care at our yearly NF Conference
• Funds adult-focused NF research

Please visit www.ctf.org for more information about CTF and the above programs and opportunities.