

Resources for Families of Children with Neurofibromatosis Type 1

Neurofibromatosis Type 1 Webinar: Answering Your Important Questions



Please join Maria Acosta, M.D., a neurologist who specializes in Neurofibromatosis Type 1 (NF1), for bimonthly webinars discussing practical topics that will help you and your child!

The purpose of this webinar is to present on topics that are important to families of children with NF1. While the topics are designed around common questions asked by parents of children with NF1, the topics are also applicable to families of typically developing children and children with other disabilities. We welcome all families to join the webinar, regardless of your child's diagnosis. Dr. Acosta will host each webinar and medical experts will be invited to present on topics you care about. Each webinar will last one hour, with 40 minutes dedicated to the webinar topic and 20 minutes of question and answer with webinar participants.

If there are topics you would like to learn more about, please reach out to Kaitlyn Tiplady at kmtiplady@ childrensnational.org with your request. This webinar is designed as a practical resource for parents and children and we want to hear from you! 2018 Schedule of Topics:

January 29 - Intro to Webinar and Review of NF1 Resources with Dr. Maria Acosta

March 26 - Tips to Improve Handwriting

June 4 - Tips for Completing Homework

July 30 - Providing Social Feedback To Children

September 24 - Advocating for School Services

November 26 - Teaching Your Child to Advocate for His/Her Needs

*Schedule is subject to change based on requests from families.

For More Information Contact Kaitlyn Tiplady at kmtiplady@childrensnational.org

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