



The Association for
Frontotemporal Degeneration
Opening the gateway to help and a cure

JOIN US

Is Your Life Touched by FTD?

Join one of AFTD's Middle Atlantic Regional Coordinator Volunteers, Corey Esannason, and AFTD Volunteer Elizabeth Soriero for an informal gathering to meet others in your community whose lives have been affected by FTD.

Saturday, October 14, 2017
1:00 p.m. – 3:00 p.m.

Lawrence Branch Library
2751 Brunswick Pike
Lawrenceville, NJ 08648

<https://www.mcl.org/branches/direct.html>

- **Connect** with others in the area whose lives have been affected by FTD.
- **Learn** more about AFTD events, programs and resources.
- **Discover** how you can get involved.
- **Share** time with others in your community.

We look forward to seeing you soon!

An RSVP by October 11th would be appreciated. To RSVP, or for any questions, please contact Corey Esannason at cesannason.aftd@gmail.com.

WHAT IS FRONTOTEMPORAL DEGENERATION?

Frontotemporal degeneration is a rare brain disease that strikes people in the prime of their life, gradually destroying their personality, ability to speak, make sound decisions, move and/or behave within social norms. Clinical diagnoses may include: behavioral variant FTD, primary progressive aphasia, progressive supranuclear palsy, corticobasal degeneration, and FTD with motor neuron disease.

WHAT IS THE ASSOCIATION FOR FRONTOTEMPORAL DEGENERATION?

The Association for Frontotemporal Degeneration (AFTD) is a non-profit organization whose mission is to improve the quality of life of people affected by FTD and drive research to a cure. www.theaftd.org