



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

JOIN US

Is Your Life Touched by FTD?

Join AFTD Support Services Manager, Bridget Moran-McCabe, for an informal gathering in Wichita to meet others in your community whose lives have been affected by FTD.

- **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.

Wednesday, November 1, 2017
4:30 p.m. – 6 p.m.

Old Chicago Pizza and Taproom
(The Thirty-Eight Room)
7626 E. Kellogg Dr.
Wichita, KS 67207

We look forward to seeing you soon!

An RSVP by October 29th would be appreciated. To RSVP, or for any questions, please e-mail Bridget Moran-McCabe at bmoran@theaftd.org.

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