



Kenan Witczak with friend Marissa Anderson

Share your name & email address, when you donate through the text-to give campaign, for a chance to win an Apple iPod touch.

*Valid 9/1/18-9/30/18

SEPTEMBER IS LEUKODYSTROPHY AWARENESS MONTH

Kenan and Krabbe disease

Kenan was diagnosed with Krabbe disease, also known as Globoid Cell Leukodystrophy, at 8 months of age. His parents, Natasha Spenser and Dann Witczak, learned quickly this disease would prevent Kenan from experiencing a full and boisterous life. Each day, Kenan was faced with increasingly profound and debilitating side effects due to the disease; something no parent should have to accept. Shortly after diagnosis, Kenan lost the ability to smile, hold his head, and grab brightly colored toys. As things progressed over his 7-year lifespan he lost the ability to swallow and required oxygen to keep him comfortable until the day he died~May 31st, 2018

With early diagnosis, state of the art resources, and access to comprehensive care, patients and caregivers can arm themselves with the information and support they need to make the journey slightly less stressful until a cure is found! Please consider a donation to help make the life of a Krabbe patient better.

Text-to-Give

9/1/18-9/30/18

Krabbe disease (KD) is a rare neurological disorder.

Any gift is powerful & WILL make the life of a Krabbe patient better!

Patients affected by KD live a shortened lifespan

Text KrabbeConnect to 707070 to make a donation today!

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KRABBECONNECT

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