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THURSDAY, JULY 13th | 6 PM to 8 PM

4130 THISTLEWOOD ROAD

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20% of the proceeds benefit Cure SPG47

KENDRA SCOTT

This is Molly and she is 3 years old. In December 2015, she was diagnosed with a neurological brain disorder called Spastic Paraplegia-Type 47 (SPG47). This is an extremely rare, newly recognized genetic disorder resembling cerebral palsy. Today, there are only 15 known patients worldwide that suffer with SPG47. Doctors believe there are many more undiagnosed or misdiagnosed with cerebral palsy.

SPG47 is present at birth and results in cognitive impairments, developmental delays, speech deficits, microcephaly, and severe, progressive spasticity and muscle weakness in the lower limbs that make movement difficult. It may eventually lead to paralysis.

Cure SPG47 aims to improve the quality of life for children impacted by SPG47 by accelerating the research for a cure and treatment. Cure SPG47 is a 501(c)3 and donations go directly to researching SPG47 and working towards a cure.

For more information about Cure SPG47 or to donate, please visit www.curespg47.org

Cure SPG47

For a Future as Bright as Their Smiles