



CURED

Campaign Urging Research for Eosinophilic Disease

My name is Abby. I am 4 years old. I love Disney movies, riding my bike, swimming, and being a big sister. I also have Eosinophilic Esophagitis (an "EOS" disease). I was recently diagnosed with "E E" and am still learning about how it will affect me as I grow up. Right now when I eat I vomit so I choose not to eat much. I take medications daily and I am on a very restricted diet. I have already undergone several medical exams and procedures. I will continue to have them until there are better treatments or a cure. I have a long road ahead of me and need all the support I can get. This is just Abby's story. Come join us at the walk to meet others who suffer from EOS diseases.

Eosinophilic Esophagitis is a very complicated and complex disease that depletes the body of iron and essential nutrients necessary for proper development and physical growth. In addition, the symptoms associated with this disease are painful cramping, bleeding, vomiting, polyps, acid reflux, ulcers, asthma, and low immune system. Each case varies from person to person. Some children must rely on a simplified liquid formula or a feeding tube for their nutrition on a daily basis. IMAGINE A CHILD LIVING THEIR DAYS LIKE THIS . . .

Come join us for **Kid's Walk for CURED** at
Rogers-Lakewood Park
Family Fun Day
May 19, 2012 starting at 8:00 am
100% of funds donated will go directly to research.

Questions? Please contact Amber Carper at amber4cured@yahoo.com
Visit our facebook page www.facebook.com/Abby4Cured
For more information about EE, please visit www.curedfoundation.org

*** Moonwalk * Face Painting * KIDS Auction * Live Music ***