Sponsor's Name, Phone	Amount Collected	
PLEDGES		
1		
2		
3		
4		
5		
6		
7		
8		
9		
10		
Make checks payable to : Sanfilippo Research Foundation. TOTAL AMOUNT		

LAPS FOR LUCAS

LAPS FOR LUCAS ANNUAL WALK /RUN EVENT WHEN: SUNDAY, MAY 18TH, 2014 REGISTRATION 2 PM WALK / RUN 3 PM

SILENT AUCTION / RAFFLE WHERE: COLLEGE COMMUNITY CAMPUS PRAIRIE POINT ACADEMY 401 76TH AVE SW CEDAR RAPIDS, IA WHAT: FUNDRAISER FOR THE SANFILIPPO RESEARCH FOUNDATION





Lucas Montgomery Age 16, Sanfilippo B



The Sanfilippo Research Foundation, Inc. AKA- Bens Dream, is a public 501(c)(3) non-profit organization. Its mission is to raise awareness of Sanfilippo Syndrome and sponsor research toward a cure.

MARCH 2014, THE NIH VOTED UNANIMOUSLY FOR GENE THERAPY FOR SANFILIPPO TYPE B TO MOVE FORWARD TO THE NEXT STEP OF CLINICAL TRIAL. HOWEVER, MILLIONS OF DOLLARS WILL BE NEEDED TO MAKE THIS HAPPEN. LET'S MAKE IT HAPPEN. **Sanfilippo Syndrome** is a rare and catastrophic genetic disorder that takes its name from Dr. Sylvester Sanfilippo, one of the doctors who first described the condition in 1963. Sanfilippo Syndrome is a mucopolysaccharide disorder and is also known respectively as MPS III. It falls within a broader group of genetic disorders known as Lysosomal Storage Diseases. The consequences vary greatly among the subgroups, ranging from a mild impact with a normal life expectancy to a severe and terminal outcome

Children afflicted with Sanfilippo Syndrome are missing an essential enzyme that breaks down strings of a complex body sugar called heparan sulfate. The partially broken-down sugar, or mucopolysaccharide, accumulates in the brain and the body's cells and tissue causing progressive damage. The storage process affects children's appearances, bodily functions and development. Sanfilippo children tend to look alike and have similar health problems.

A Sanfilippo child appears normal at birth and develops within the range of normal for the first year or two, but as more and more cells become damaged symptoms begin to appear. Eventually, the build-up of muccopolysaccharides will cause hyperactivity, sleep disorders, loss of speech, mental retardation, seizures, loss of mobility, dementia, vegetative state and finally death. Life expectancy for a child with Sanfilippo Syndrome is between 12 to 20 years.

REGISTRATION FORM 5K1MILE		
NAME:	T-shirt Size:	
DONATION:	PLEDGES:	
Address:		
City:	ST:Zip:	
E-mail address:		
MAKE CHECKS PAYABLE TO : SANFILIPPO		
RESEARCH FOUNDATION DONATIONS AND PLEDGES OF \$100 OR MORE QUALIFY FOR LAPS FOR LUCAS TEES		
SIZES : ADULT 2XL,XL,L,M,S YOUTH M PLEASE EMAIL TSHIRT SI.ZES AND PREREGISTRATION TO		
DT WAT 1°', 2014		
(ASK YOUR EMPLOYER ABOUT A MATCHING GIFT PROGRAM and ATTACH		
FORM)		
MAIL DONATIONS OR BRING REGISTRATION THE DAY OF THE EVENT		

2014 LAPS FOR LUCAS 3507 REMINGTON ST SW CEDAR RAPIDS IA 52404