

Fatty Oxidation Disorders
Family Support Group
www.fodsupport.org

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## FOD Family Support Group

June 4, 2018

Adventures for the Cure Attn: Patrick Blair 1221 Brandford Rd Baltimore, MD 21228 pblair12@gmail.com

Dear Patrick and Team:

On behalf of all of our Families, thank you for your very generous donation of \$2548.14 to our Research Trust Fund/LCHAD Research (via check# 1020). Your donation given on May 27, 2018 as part of your Fundraising Bike Rides, will assist us in continuing our worldwide efforts to provide emotional support and networking, practical information, and Medical Updates to inform families of new developments in screening, diagnosis, research and treatment. Your donation will also help us toward our goal of raising more funds for FOD clinical training and research endeavors, as well as offering Conferences for our FOD Families and free of charge grief support to bereaved parents and families. Thank you for thinking of our Families!

Deb Lee Gould, MEd, Director MCAD Parent and Grief Consultant Grief Support for Bereaved Parents and

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**Families** 

PO Box 54

Take care...

Deb Lee Gould, MEd

Deb Lee Gould, MEd, Director

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Fatty Oxidation Disorders: Medium Chain Acyl CoA Dehydrogenase Deficiency (MCAD); Medium Chain 3-Ketoacyl coA Thiolase Deficiency (MCKAT); Medium/Short Chain L-3 Hydroxy acyl coA Dehydrogenase Deficiency (M/SCHAD); L-3-Hydroxy Acyl CoA Dehydrogenase Deficiency (LCHAD); Carnitine Palmitoyl Transferase I and II (CPT I & II); Very Long Chain Acyl CoA Dehydrogenase Deficiency (VLCAD); Short Chain Acyl CoA Dehydrogenase Deficiency (SCAD); 3-Hydroxy Acyl CoA Dehydrogenase Deficiency (HADH, formerly called SCHAD); Short Chain 3-Ketoacyl coA Thiolase Deficiency (SKAT, 3-ketothialase); Electron Transfer Flavoprotein or Electron Transfer Flavoprotein ubiquinone-oxidoreductase Deficiency (ETF and ETFQ0, also known as GAII or MADD); Trifunctional Protein Deficiency (TFP); Carnitine Transport Defect (Primary Carnitine Deficiency); Carnitine-Acylcarnitine Translocase Deficiency; 2,4 Dienoyl-CoA Reductase Deficiency; 3-Hydroxy-3 Methylglutaryl Deficiency (HMG); Unclassified FODs

\*No goods or services were provided in exchange for this contribution

The FOD Family Support Group is an all Volunteer Family-based Group and a tax-exempt 501c3 Non-Profit Corporation

US Donations are tax-deductible

Thank you to all that have donated over the years so that we may continue to provide our support and information to Families and interested professionals around the world.

Donations help with copying and postage costs, phone calls to new FOD Families (US and abroad), website fees, Conference costs,

as well as to provide grants for future clinical and research activities by our FOD professionals. Please make <u>US checks</u> out to: FOD Family Support Group and <u>Mail to</u>: PO Box 54, Okemos, MI 48805-0054