

# TitanTough21 Foundation

Join the Journey...

Dear Li-Fraumeni families,

My name is James D. Everett and I'm the Chairman of the Board of the **TitanTough21 Foundation**, and also a father of two children with Li-Fraumeni syndrome.

One year ago, the TitanTough21 Foundation was formed. We are very excited to team up with other great charities that are working hard to directly fund LFS research and help LFS families that are in need of paying outstanding medical bills.

It has been a very good year for TitanTough21. We've made great progress putting all the pieces together and holding some great events. We are ready to go worldwide and help find a cure through our fundraising efforts.

I am reaching out to you because TitanTough21 will be having an official **launch event** on Saturday, October, 17<sup>th</sup> 2015 at 11:00am MST in Salt Lake City, Utah. We are pleased to announce that we will be having a live **Webcast** that will let you know what we're all about. We'll be having **Dr. Joshua Schiffman** as our keynote speaker. He will be sharing the latest medical research from the Schiffman Laboratories located inside the Huntsman Cancer Institute. In addition, you will get to meet and hear from **Reagan Everett**, the founder of the **TitanTough21 Foundation**, as well as our board members.

TitanTough21 can directly help your family find hope in what can be very troubling times as you deal with LFS first hand.

Just before the broadcast time please go to the TitanTough21 website [www.titantough21.org](http://www.titantough21.org) and click on the **"Live Broadcast"** button to watch the stream.

I'll be sharing with you how we can raise **"Thousands of dollars"** for LFS research.

The only way that a cure will happen is by having every LFS family come together. I look forward to presenting TitanTough21 to you and your family.

Much love and hope,

James D. Everett

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