



Jonny Lee Miller meets Jonah's Just Begun

Jonny plays Sherlock in the new hit show Elementary. Jonny first learned about Jonah's Just Begun, when Jeremy Weishaar a camera man on Elementary asked him to support a local fundraiser. As Jonny learned more and more about the horrific situation that Jeremy and his family faced he was compelled to do something. Running his first ultra-marathon (50 Miles) in May 2013, raising \$120,000 for medical research. Jonny did not stop there, he followed up with a second ultra-marathon and on July 5th 2014, he will run 50 miles through the streets of NYC.

To support Jonny and his compassion to help save the lives of children go to: <http://bit.ly/NYCultra4JJB>

Jonny's compassion and commitment to helping JJB fund a treatment for Sanfilippo has gone beyond running marathons. Jonny has leant his celebrity to appeal to the masses by appearing on the View and CNN's Dr. Sanjay Gupta. Jonny also accompanied Jonah's Mom, Jill Wood to D.C. where they spoke to our countries legislators on behalf of the Rare Disease Congressional Caucus. Most recently Jonny and Kristine Johnson, anchor person for CBS 2 hosted JJB's first NYC annual "Rare Occurrence Reception."

Stay tuned and in the know by following: JLM @jonnyjlm and JJB @JJB4CURE on twitter.



If a glass of lemonade could save a child—would you buy one? Of course, you would!

That is what 6 year old Grey, “PURPLE LEMONADE -a -STAND for Sanfilippo Syndrome” founder, knew when she decided to host her first lemonade stand. Determined to fund a cure for the ultra-rare and terminal disorder Sanfilippo Syndrome, which her sister Blair, suffers from. Grey delivered flyers, hung posters, baked goodies and made PURPLE lemonade. F.Y.I. Purple is the most rare naturally occurring color and is the designated color for Sanfilippo Syndrome.

Grey’s stand was a huge success & today “Purple Lemonade A Stand for Sanfilippo Syndrome” is a worldwide effort!

Several Sanfilippo medical research foundations such as: Jonah’s Just Begun, Ben’s Dream and The National MPS Society have joined together to raise funds for breakthrough gene therapy research. Once only a possibility, gene therapy for Sanfilippo Syndrome is NOW on the horizon. but it needs funding— it needs YOU to buy a glass of lemonade! Be it in person at a “PURPLE LEMONADE - a-STAND for Sanfilippo Syndrome” or virtually through our crowdrise Purple Lemonade Team.

Find us here: <https://www.crowdrise.com/purplelemonade4jbb> #purplelemonade #JJB4CURE #MPSIII

