CHOC Foundation

CHRONICLE

NEWS AND INFORMATION FOR SUPPORTERS AND FRIENDS

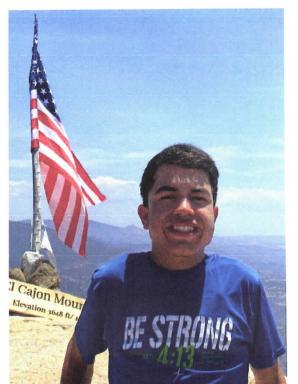
CALEB'S STORY OF COURAGE, DETERMINATION AND GIVING BACK

When he was three years old, Caleb was diagnosed by a CHOC genetic specialist with what's known as MPS II. This extremely rare and devastating genetic disease affects nearly every organ in the body, along with the joints and spine.

"Within a few days," his dad, Rory, recalls, "we were in CHOC's metabolic clinic and meeting Dr. Raymond Wang, a metabolic disorders specialist. When we walked into the patient room, we weren't just another number. We had all these experts in one room, but they also had really big hearts."

Over the past 13 years, the family has formed a special bond with Dr. Wang and each of the many doctors on Caleb's care team.

Caleb has had numerous procedures at CHOC, including hernia repair, adenoids and tonsils removal, two double-knee surgeries and, most recently, double abdominal surgery. There have also been many MRIs, ultrasounds, appointments and physical therapy sessions. Caleb stresses how our devoted doctors and nurses are "very compassionate and understanding, whatever the situation."



Persevering through the ups and downs

Despite his many health challenges—and enduring a four-hour infusion treatment every week—Caleb's determined to live his best life possible. His mom Marie confides, "Suffering produced his character. When I think of Caleb today, I think of strength and courage."

For example, an avid runner since he was seven, this high school junior didn't let his double-knee surgeries keep him down. His resilience has shined through. Amazingly, he currently competes on his high school's track and field team.

Just as amazing is the deeply appreciated commitment Caleb and his family are making to CHOC's lifesaving work.

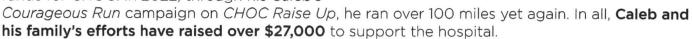


"For all that we had been blessed with by CHOC, we wanted to give back and help support other families."

-Marie, Caleb's mom

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From 2014 to 2019, they participated in *CHOC Walk in the Park*. In the following two years, Caleb ran over 100 miles each year to raise funds for CHOC. In 2022, through his *Caleb's*



Rory explains, "As a patient, you're on the receiving end. We wanted to be on the giving end. We give back because we want Caleb to be a part of helping to find a cure and helping other families."

Caleb, who plans to attend college and study American History, says, "I just feel like I wanted to do something to give back to CHOC and help them in what they're pursuing for children like me who are in the very difficult situation of fighting a rare disease."

To find out more about how Caleb created his own fundraiser to help the children at CHOC, please visit: *raiseup.choc.org*





CHOCHeroes of Hope

BE A HERO EVERY MONTH

When you join our monthly giving group, **CHOC Heroes of Hope**, you'll help us respond to urgent patient needs as soon as they arise. This is an especially meaningful way for you to make an ongoing difference in the lives of the children and families we serve.

Please contact Chloe Addleman at *chloe.addleman@choc.org* or (714) 509-4301 for details about how to become a member of **CHOC Heroes of Hope**.



Nurturing, advancing and protecting the health and well-being of children since 1964