





Jaxson



- Starts and ends at Indian Motorcycle of El Cajon.
- ® Registration starts at 9am; Bikes out by 10am. (Complimentary donuts from Mary's Donuts!)
- \$20 per player, \$10 each additional /card or riders.
- © End of Ride Celebration at 1pm which will include results from Winning Poker Hand, 50/50, our Raffle Drawings, AND Auction Items

Food will be graciously provided.

~ Please check with Jaxson's Train of Hope on Facebook
for any updates or ride changes ~

https://www.facebook.com/groups/jaxsonstrainofhope/
or: https://www.jaxsonstrainofhope.net

Jaxson Karp has a rare disease called Tay-Sachs.

It is a genetic neurological disorder that is fatal and there is no cure <u>yet</u>.

It would be an honor if you would join us to help raise money towards a cure and celebrate his <u>10th</u> birthday in the meantime!







National Tay-Sachs & Allied Diseases 501c (3) www.ntsad.org

If you would like to be involved with drawing prizes, sponsorship, volunteering, or assisting with our ride, please reach out to thekarps@jaxsonstrainofhope.net or call 619 916-9226





Jaxson & Christine











Starts at 9:00 am

What is NTSAD?

NATIONAL TAY-SACHS & ALLIED DISEASES Association How do they help Tay-Sachs patients?

Anyone can be a carrier of Tay-Sachs. The carrier rate for the general population is 1/250. Supporting families is the center of everything NTSAD does. There is no cure at this time, but there is hope for a gene therapy treatment in the near future that can possibly stop the progression of the disease. Without treatment Jaxson will suffer from muscle weakness, lack of coordination, swallowing difficulty, seizures, loss of ability to communicate, see, walk, and eat by mouth. This disease progresses at it own pace; there is no time line for life expectancy.

National Tay-Sachs & Allied Diseases Association (NTSAD) is one of the oldest non-profit organizations dedicated to helping families.

- Connects families through their Family Services program including annual family conferences, fundraisers, and awareness materials;
- Funds research on all levels and facilitates the steps toward clinical trials;
- Educates families and the public to promote building healthy families though screening.

Each year NTSAD supports nearly 750 families by providing services, resources, and compassionate support to the affected families of Tay-Sachs, Canavan, GM1, and Sandhoff diseases.

THANK YOU ALL FOR YOUR SUPPORT!
The Karp Family & National Tay-Sachs & Allied Diseases



