

BISOUS FOR LÉO BENEFITS THE INADCURE FOUNDATION, THE ONLY US-BASED NON-PROFIT WORKING ON TREATMENT AND CURE DEVELOPMENT FOR INFANTILE NEUROAXONAL DYSTROPHY (INAD).

INFANTILE NEUROAXONAL DYSTROPHY (INAD)

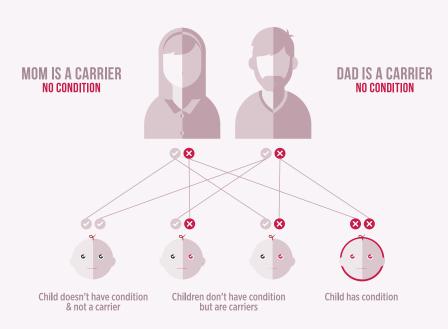
OFTEN REFERRED TO AS PARKINSON'S MIXED WITH ALZHEIMER'S FOR KIDS.

INAD IS AN EXTREMELY RARE, INHERITED DEGENERATIVE DISORDER

OF THE NERVOUS SYSTEM.

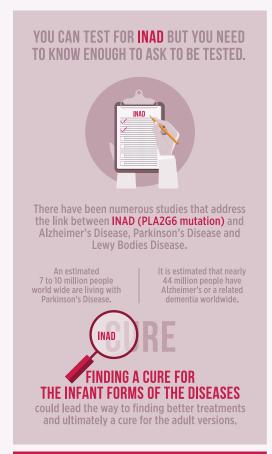


IT IS AN AUTOSOMAL RECESSIVE, MEANING KIDS INHERIT The Mutated genes from Both of their "Carrier" parents



Children typically develop "normally" until they're between 6 months and three years old, and then they start to lose acquired skills. They will lose mental and cognitive abilities, as seen in Alzheimer's, and physical abilities, as seen in Parkinson's, resulting in death likely before they reach their 10th birthday.





AWARENESS IS KEY

In September 2018, Bisous For Léo launched an Instagram account@BisousForLeo encouraging its followers to help

KISS INAD GOODBYE of not just for Léo but for all children affected by the disease.

People from all around the world have been posting their kisses (or Bisous as they say in French) and tagging Bisous For Léo.



Gal Gadot, Sebastian Stan, Zoe Saldana, Eva Longoria, Olivia Munn, Isla Fisher and so many others have all posted in solidarity with Léo. We'd love for you to post your kisses too!

FOR MORE INFORMATION OR TO MAKE A DONATION VISIT WWW.BISOUSFORLEO.ORG

THANK YOU FOR HELPING US KISS INAD GOODBYE!