

Submitter: Cassandra Brown
On Behalf Of: Ezra R
Committee: Senate Committee On Human Services
Measure: SB646

Hi, I'm writing on behalf of my son Ezra who was born with a complex heart anatomy and a rare genetic condition called Cornelia DeLange Syndrome. We are in support of SB 646. My son is 3 and takes several medications, has breathing treatments, therapies, and constant appointments. Like many families with disabled children, we either sacrifice our disabled child with a non-parent caregiver, increasing the odds of abuse and neglect or we sacrifice our income so our child gets the absolute best care they can receive. My son is my absolute world, so of course, I chose him. We must pass SB 646 so families can continue to care for and enrich their disabled kids lives without living in poverty. We must support families with disabled children. This is the safest and most cost effective choice for the state financially and ethically.

If you'd like to read more about my rare superstar, feel free to follow the link.

https://issuu.com/cdlsusa/docs/ro-t3-2022?fbclid=IwAR35H_c0mQJEGIJss96_OzejftgOIYtCe96azxum_S7VUU-3fNOHeVnmW9Q&mibextid=Zxz2cZ