



# EN POINTE FOR AVA NEWSLETTER



## Meet Daphne from Arizona

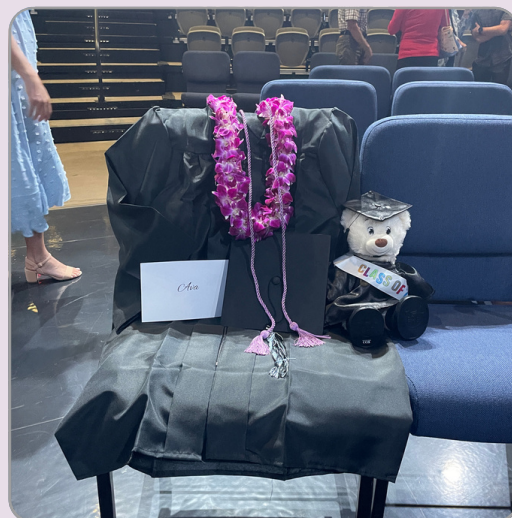
Daphne from Arizona has been receiving support from En Pointe for Ava for nearly a year! We love these success stories! Daphne, 20, travels from Arizona to Stanford Children's Hospital for her CAR-T treatment for DIPG each month. She recently wrote this: ***"I can't even begin to describe how much En Pointe for Ava has done for me and my mom. As a chronically ill daughter with a single mom, many of our challenges fall on her. Since her only job is part-time, many of our issues are financial. My social worker helped us apply for the program and the support has truly helped us stay on our feet. I am deeply grateful for this opportunity and I hope others will be able to receive the help they need."*** Daphne, we are with you through the journey.

## Upcoming Events



### THE EPFA 4<sup>TH</sup> ANNUAL GOLF TOURNAMENT OCT 24TH

Join us on Saturday,, 10/24, at Twin Oaks Golf Course for our biggest fundraiser of the year. Go to our website to sign up!



## Your help makes a difference

### BECAUSE OF YOU

We are overjoyed to share incredible news: **Meadow's tumor is no longer just stable, it's smaller!** For children battling DIPG, this is extraordinarily rare and gives all of us renewed hope for the future. **Because of your unwavering generosity,** En Pointe for Ava has been able to stand beside Meadow and her family throughout this journey for 2 years. Your support is making it possible for families to access life-saving treatments they otherwise might never reach. This victory belongs to all of us. **Together, we did this!**

### MAKING PURPOSE FROM PAIN

This year, Ava should have walked across the stage to receive her high school diploma. Milestones like these never stop reminding us of all that DIPG stole, not only from Ava, but from every child and family facing this devastating disease. These moments still weigh heavily on our hearts, but they also strengthen our resolve to honor Ava's legacy by funding hope, supporting families, and helping advance the search for a cure. Every child deserves the chance to celebrate life's milestones, and together, we will continue fighting until no family has to endure the heartbreak of DIPG.