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### **Kat Buckley**

*Brian Buckley, father*

We are incredibly lucky to have a strong support system—friends, family, and coworkers that continuously show up. Most importantly, Dad was still the book loving, life of the party, family man that we all knew and loved. He never complained, and he never gave up. *#buckleystrong*



### **Kennedy Watson**

*Wendy Gunderson, mother*

We have had many ups and downs, but her strength always shines through. She has always been a super mom and still hasn't changed. She continues to teach me to be patient and kind to others because you never know what someone is going through. We find hope in knowing one day there will be a cure, maybe not in my mom's lifetime but someday, and we will know we were a part of that.



### **Brooke Stobbe**

*Geena Stobbe, mother*

Her impact as a nurse, support staff, and advocate at work is too extensive to report. Only one word comes close: Hero. It's our responsibility to take what she has taught us and carry it into our own lives and adopt it as our own. She is alive in all of us, and we must all live the way she demonstrated. Geena spent the last five years fighting PSP with the utmost grace, poise, and joy.

BECAUSE **HOPE**  
**MATTERS**

## WHAT YOU DO MATTERS

Dear Jeffrey,

It has been a challenging year for all of us, and there are still many hurdles that lie ahead before normal life can resume. But for patients and families suffering from progressive supranuclear palsy (PSP) and related prime of life neurodegenerative diseases, life will never return to normal. These diseases are quickly debilitating, require 24/7 care that usually causes financial hardship, and always lead to the disruption of family equilibrium. PSP is largely untreatable and currently incurable, but CurePSP is working hard to change that. Thanks to your support, we are making progress.

*In this fight, hope matters. In this brochure, we present three stories of hope.*

Kat Buckley is grateful for her support network—friends, family, and coworkers—that help her in her role as caregiver to her father, Brian. Kennedy Watson takes comfort in the strength her mother, Wendy, showed in her fight against PSP. Brooke Stobbe is inspired by the lessons in perseverance that her mom, Geena, taught her in her brave five-year struggle with the disease.

Now, we are asking for your support in our fight against neurodegeneration. With the resources YOU provide, CurePSP is able to support patients and families during their arduous combat with disease, educate physicians and allied healthcare professionals for quicker and more accurate diagnosis and better treatment, and fund important research that is opening doors to drug clinical trials and eventual prevention and cure.

Please help tens of thousands of people like Kat, Kennedy, and Brooke with your generous donation. I can personally assure you that it will be used wisely and well, and I thank you in advance for your support. Because of you, we have hope!

With gratitude,



**David Kemp**

PRESIDENT

## BECAUSE HOPE MATTERS: CHOOSE TO ACT

YES! I choose to stand with CurePSP to help fight for a future free of neurodegeneration.

\$50    \$100    \$500    \$1,000    \$5,000    Other \_\_\_\_\_



David Q. Public  
123 Anywhere Street  
Centralville, US 54321

My check payable to CurePSP is enclosed.

To use your credit card, please check the box and turn this card over.

**curePSP**<sup>®</sup>

UNLOCKING THE SECRETS OF BRAIN DISEASE<sup>®</sup>

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