THE FOUNDATION FOR PRADER-WILLI RESEARCH

ADVANCING TREATMENTS FOR PWS

It Takes a Village

We know the challenges our loved ones will continue to face if we do not find treatments for PWS. For many of us who live with PWS every day, the status quo is simply not an option. **Our loved ones with PWS are relying on us.** This is why we pour our blood, sweat, and tears into raising funds for research: it's our only path to a better future. Our donor network of family and friends are imperative to our success and we thank you for your continued support!

Your FPWR dollars:

- ☑ Identify potential new treatments
- ☑ Expand the drug development pipeline
- ☑ Advance genetic therapies
- ☑ Improve clinical care
- ☑ Establish and maintain research tools
- ☑ Advocate for new treatments

FOUNDATION FOR PRADER-WILLI RESEARCH

To learn more about your impact contact:
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ADVANCING TREATMENTS FOR PWS

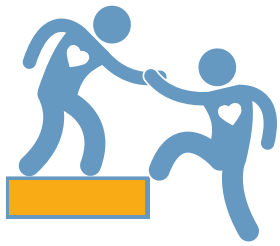
Why do we need your help to fulfill our mission?

WE HAVE MADE TREMENDOUS PROGRESS BUT WE CAN'T SLOW OUR EFFORTS UNTIL TREATMENTS ARE FOUND.

Community support has been vital in developing our drug pipeline. More than one dozen drugs are now in various stages of development. Our team dedicates hundreds of hours each year to supporting drug development and we need your continued support so that we may aid in clinical trial design and work to develop trials that have the best chances for success.

WE NEED TREATMENTS NOW!

On average it takes 12 years to bring a new drug to market, but people with PWS and their families need treatments NOW! We are working to find near-term solutions that will alleviate symptoms of PWS and improve quality of life. FPWR dedicates funds each year to support projects that will improve the treatment of behavioral, scoliosis, seizures, and GI motility, mental health, and more.



GENETIC THERAPY COULD TRANSFORM PWS AS WE KNOW IT

Can you imagine an independent life for our loved ones with PWS? We can. A successful genetic therapy approach could simultaneously improve many aspects of PWS, but much still needs to be learned, and tested, to determine what benefits this approach will have for individuals with PWS BEFORE we can advance this treatment into clinical trials.

FAILURE IS NOT AN OPTION

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