PEOPLE WITH PASSION
AND PURPOSE

2017 Progress Report
**VISION:**

Prader-Willi Syndrome Association (USA) is a self-sustaining, internationally recognized leader, empowering those affected with Prader-Willi syndrome to enjoy a productive life in an informed and accepting community.

**OUR MISSION:**

Prader-Willi Syndrome Association (USA) is an organization of families and professionals working together to raise awareness, offer support, provide education and advocacy, and promote and fund research to enhance the quality of life of those affected by Prader-Willi syndrome.
Contents

SAVING AND TRANSFORMING LIVES TOGETHER 4
PEOPLE WITH PASSION AND PURPOSE 5
PWSA (USA): FOUNDED ON THE FIVE PILLARS OF SUPPORT 6
AWARENESS 7
FAMILY SUPPORT 8-9
RESEARCH 10-11
EDUCATION 12
ADVOCACY 13
FINANCIALS 14-15
WHO WE ARE 16-17
SPECIAL THANK YOU TO LEADER 18
OUR COMMUNITY NEEDS YOU 19
Dear Prader-Willi Syndrome Association (USA) Supporter:

Identified in 1956 by Swiss doctors A. Prader, H. Willi, and A. Labhart, Prader-Willi syndrome is a genetic birth defect that adversely affects appetite, growth, metabolism, cognitive functioning, and behavior. Individuals with PWS face life-long challenges that impact all areas of their lives, including at home and at school. Please see examples of how we worked to help alleviate these challenges on the following pages.

Prader-Willi Syndrome Association (USA) was formed in 1975 to unite parents, professionals, and other interested citizens who share a common purpose: To enhance the quality of life of those affected by Prader-Willi syndrome.

Channeling the strength, commitment, and hope of thousands of dedicated individuals, PWSA (USA) is an organization that empowers the PWS community through shared experiences, research, education, advocacy, and support. With chapters in most states, PWSA (USA) is the only national PWS support organization whose sole purpose is to assist individuals with PWS and their families every step of the way.

Our network of caregivers, medical providers, educational professionals, support staff, and families have worked together for over 40 years to support PWS families by providing information, resources, and support for school, medical, and behavioral management. PWSA (USA) is an important lifeline for individuals and families across the country; without our services, many in the PWS community would have nowhere else to turn. Thank you for your support. 

Together we are Saving and Transforming Lives.

Sincerely,

Michelle Torbert
Chairperson of the Board

Saving and Transforming Lives Together
United by passion and purpose, PWSA (USA), fellow Prader-Willi syndrome and rare genetic disease-focused researchers and organizations, volunteers, donors, and families are working together to facilitate research, spread awareness, advocate, and share knowledge for, with, and about the PWS community. It is this transformative power of collaboration that inspires our efforts and motivates us to keep pushing forward as each day brings forth new opportunities to save and transform the lives of those affected by PWS.

Since 1975, PWSA (USA) has created and expanded programs for

- Parents of newly-diagnosed children
- Deployed life-changing family support programs
- Funded cutting-edge research
- Provided training and information to school professionals, residential providers, and healthcare providers across the country

With support from the PWS community, our donors, and our volunteers, PWSA (USA) has brought hope, health, and enhanced quality of life to thousands of individuals and families, helping them thrive in the face of a rare genetic condition. People with passion and purpose working together to Save and Transform Lives…we are PWSA (USA).

Sincerely,

Steve Queior
Chief Executive Officer
PWSA (USA):
Founded on the Five Pillars of Support

PWSA (USA) is a collaboration of families, individuals, researchers, healthcare practitioners, and professionals, working together to increase awareness, provide family support, facilitate research, promote education, and advocate for the PWS community.

These five pillars serve as the foundation of who we are, what we do, and what we can accomplish working together to enhance the lives of those affected by Prader-Willi syndrome.
Awareness is the precursor to action. As such, it is imperative to gain attention for Prader-Willi syndrome and to facilitate conversations about the needs of the PWS community. Educating the public about Prader-Willi syndrome and the unique challenges faced by those affected by it, and spreading the message of what PWSA (USA) does to help individuals and families meet those challenges, are two of our most fundamental tasks.

We elevate awareness and educate through traditional media outlets, social media, group presentations, and personal interactions. Partnering with other organizations to make the most of national and global opportunities to raise awareness, such as "Rare Disease Day" is also paramount to our success.

Raising Awareness: Outreach Through Events

PWSA (USA) hosts or co-hosts several annual and biennial events to raise funds and awareness for the Prader-Willi syndrome community. Coordinated by individuals and PWSAs state chapters, events such as On the Move walks and runs provide fun and engaging opportunities to introduce others to the needs of the PWS community. The biennial PWSA (USA) National Convention is another popular event that brings medical and residential professionals, researchers and scientists, families, and individuals together to share information and support. National and local awareness and fundraising events include:

• Create Your Own FUNraisers
• On the Move run/walks
• Major fundraising events, such as the annual Clint Hurdle "Hot Stove Dinner"
• Biennial PWSA (USA) National Convention

Raising Awareness: Outreach Through Publications

Communication of professional advice, industry best practices, and more is available through PWSA (USA)'s robust catalog of publications.

Accessible online and in print, PWSA (USA) produces dozens of brochures, white papers, articles, and a bi-monthly information-packed newsletter to keep families, caregivers, and school, medical, and residential professionals up-to-date on the latest PWS information.

PWSA (USA)'s website and mobile app are other "go to" sources of PWS information available for families and professionals. In addition to blogs and articles, visitors to the site or app can find resources such as the Wyatt Special Education Advocacy Training modules and PWSA state chapter contact information.

Raising Awareness: Outreach Through Local Relationships

With 33 state chapters throughout the country, individuals and families seeking local help and support can connect with a caring group of individuals who understand the complex needs of those affected by PWS. Whether they need assistance with new diagnosis, a school crisis, or a medical referral, help is never more than a phone call away.
Family Support

The PWSA (USA) Family Support & Crisis Intervention Team includes family support counselors, a medical coordinator, new parent support coordinators, and a special education specialist. These dedicated professionals coordinate programs to help families with behavior modification, nutrition education, crisis intervention, education, advocacy, guardianship, and medical intervention. In 2017 alone, the PWSA (USA) Family Support team handled over 2,370 family crisis and medical consultations and provided staff and material support to the families of 218 newly-diagnosed children.

Family Support: Special Education Advocacy

PWSA (USA) is committed to helping individuals with PWS have a positive and successful school experience. Skilled Family Support team members are available to attend education planning meetings (either in person or by phone) and to provide information and training to school professionals working with children with PWS. PWSA (USA) also makes available the Wyatt Special Education Advocacy Training (WSEAT), a web-based course designed to equip parents and guardians with the information, skills and tools they need to be more effective special education advocates for their children. Finally, the Family Support Team produces School Times, an electronic newsletter that focuses on timely and helpful school-related topics available to families and school professionals.

Family Support: Managing the Day-to-Day

Families affected by Prader-Willi syndrome face many unique challenges, and without adequate support can quickly find themselves in crisis. Family Support team members help parents and guardians answer questions like:

- How can appropriate behavior and weight management be encouraged and supported?
- How can siblings be helped to understand and support their brother or sister with PWS?
- How can a couple cope with the stress of raising a child who has special needs?
- How do caregivers take care of themselves while caring for a child with PWS?

In addition to the many online resources available on the PWSA (USA) website, help can be found over the phone, via email, through Facebook support groups, or in-person.

Family Support: Adults with PWS

Adults with PWS require specialized care that many residential providers are not equipped to provide. Family Support team members offer professional provider training and education to ensure individuals with PWS receive the services and attention they need. In addition to spreading awareness and education through training, the Family Support team also produces Residential Times, an electronic newsletter focused on adults living with PWS, their families, and the professional caregivers who support them. Topics covered in the newsletter include:

- Do's and don'ts for improving the parent-provider relationship
- Information on the Achieving a Better Life Experience Act (ABLE)
- Advocacy network updates
Family Support: Funds Invested in Services

<table>
<thead>
<tr>
<th>Amount of budget allocated to Family Support services</th>
<th>$948,400</th>
</tr>
</thead>
<tbody>
<tr>
<td>Components including Crisis &amp; Family Support, Medical Intervention &amp; Support, New Diagnosis Support, Education, Family Advocacy, and more.</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Number of Family Support activities in 12 month period</th>
<th>2,371</th>
</tr>
</thead>
<tbody>
<tr>
<td>1. New Diagnosis Support Component</td>
<td>210 Service Activities</td>
</tr>
<tr>
<td>New Diagnosis Funding</td>
<td>$139,650</td>
</tr>
<tr>
<td>Average Invested in New Diagnosis Case</td>
<td>$665</td>
</tr>
<tr>
<td>2. All Other Family Services</td>
<td>2,161 Service Activities</td>
</tr>
<tr>
<td>All Other Family Support Funding</td>
<td>$808,750</td>
</tr>
<tr>
<td>Average Invested in Family Support Case</td>
<td>$373</td>
</tr>
</tbody>
</table>

NOTE: Overall average cost of family support activity is $400.

<table>
<thead>
<tr>
<th>Translating Donation Amounts into the Good Work They Can Fund:</th>
</tr>
</thead>
<tbody>
<tr>
<td>• A $10,000 donation can put 15 families with a newly-diagnosed child on to a path Saving and Transforming Lives of the children and their entire families.</td>
</tr>
<tr>
<td>• Monthly giving of $100 will fund the annual Family Support services for 3 families.</td>
</tr>
<tr>
<td>• $40,000 raised in the Family Support Campaign can make all the difference in the health, safety, and quality of life for 100 families this year and beyond.</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Additional PWSA (USA) Funds Invested in Research:</th>
</tr>
</thead>
<tbody>
<tr>
<td>NOTE: The average amount invested in research in 2016 and 2017 was over $454,000 annually.</td>
</tr>
<tr>
<td>This amount is separate from, and additional to, the $948,400 above.</td>
</tr>
</tbody>
</table>
Research

Since 1983, PWSA (USA) has been at the forefront of Prader-Willi syndrome scientific study. Many of the world’s most renowned PWS researchers and clinicians are on PWSA (USA)’s Scientific and Clinical Advisory boards, and the organization has invested more than $2 million to facilitate life-changing research over the past ten years. The synergy between PWSA (USA)’s advisory boards and research committee is magnified by ongoing collaboration with other PWS and rare disease researchers and organizations, and daily interaction with parents, caregivers, and individuals with PWS. Through PWSA (USA), the powerful combination of experience and information come together to inspire change and enhance the quality of life of individuals with PWS and their families.

Research: Uniting Science and Experience

The 2017 PWSA (USA) National Convention featured a special session for individuals interested in learning more about the organization’s research program. A special focus was made on familiarizing parents with PWS research with presenters sharing basic information about PWS research and encouraging questions from the audience.

Research: 2017 and Beyond

While all Prader-Willi syndrome research is important, PWSA (USA) is particularly interested in facilitating research that can more immediately enhance quality of life of individuals affected by PWS. The desire to find more and better treatment options to manage and diminish challenges inherent to Prader-Willi syndrome will guide PWSA (USA)’s research initiatives now and into the future. PWSA (USA)’s research committee and advisory boards aim to:

• Devote more resources to support development of new therapeutic interventions
• Increase efforts to collaborate with external research partners, such as pharmaceutical companies and the Foundation for Prader-Willi Research
• Evaluate the current grant-making process to ensure a treatment-based research focus
• Encourage innovative research models
• Create new opportunities for researchers and scientists to promote collaboration and information sharing
## Research: Recent PWSA (USA) Investment in Innovation

<table>
<thead>
<tr>
<th>Project Description</th>
<th>Amount</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>RDCRN</strong></td>
<td>$25,000</td>
</tr>
<tr>
<td><strong>PWS Consortium – 4 Separate Institutes</strong></td>
<td>$85,000</td>
</tr>
<tr>
<td>University of Florida</td>
<td></td>
</tr>
<tr>
<td>University of Southern Florida</td>
<td></td>
</tr>
<tr>
<td>KUMC – Butler</td>
<td></td>
</tr>
<tr>
<td>Regents of California</td>
<td></td>
</tr>
<tr>
<td><strong>“Oxytocin Study” DMCC</strong></td>
<td></td>
</tr>
<tr>
<td><strong>Oxytocin Stage Two</strong></td>
<td>$324,178</td>
</tr>
<tr>
<td>Miller, Dr. Jennifer, M.D.</td>
<td></td>
</tr>
<tr>
<td>University of Florida</td>
<td></td>
</tr>
<tr>
<td><strong>Hormone Secretion Deficits in PWS</strong></td>
<td>$240,000</td>
</tr>
<tr>
<td>Nicholls, Dr. Robert, Ph.D.</td>
<td></td>
</tr>
<tr>
<td>Children’s Hospital of Pittsburgh Foundation</td>
<td></td>
</tr>
<tr>
<td><strong>The Effect of Growth Hormone Substitution on Sleep Disordered</strong></td>
<td>$78,034</td>
</tr>
<tr>
<td><strong>Breathing in Young Children with PWS</strong></td>
<td></td>
</tr>
<tr>
<td>Choong, Dr. Catherine, M.D.</td>
<td></td>
</tr>
<tr>
<td>University of Western Australia</td>
<td></td>
</tr>
<tr>
<td><strong>Profiling of the Gut Microbiomes in Children with PWS</strong></td>
<td>$32,190</td>
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<tr>
<td>Haqq, Dr. Andrea, M.D.</td>
<td></td>
</tr>
<tr>
<td>University of Alberta</td>
<td></td>
</tr>
<tr>
<td><strong>Male Caregiver Study</strong></td>
<td>$5,000</td>
</tr>
<tr>
<td>Caldwell, Leon, Ph.D.</td>
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</tr>
<tr>
<td><strong>Stipend/Support Research Students Interested in PWS</strong></td>
<td>$6,000</td>
</tr>
<tr>
<td>Kimball, Ted (Under Direction of Dr. Merlin G. Butler, M.D., Ph.D.)</td>
<td></td>
</tr>
<tr>
<td><strong>Prevalence of Dysphagia/Reflux Contributing Factor</strong></td>
<td>$33,955</td>
</tr>
<tr>
<td>Yeung, Dr. Karla Au, M.D.</td>
<td></td>
</tr>
<tr>
<td>Johns Hopkins to Valley Children’s, Fresno, CA NOW</td>
<td></td>
</tr>
<tr>
<td><strong>Telehealth Intervention of Early Social Cognitive Process</strong></td>
<td>$59,931</td>
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<tr>
<td>Dimitropoulos, Anastasia, Ph.D.</td>
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<tr>
<td>Case Western Reserve University</td>
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<tr>
<td><strong>International Clinical Trials Consortium</strong></td>
<td>$20,000</td>
</tr>
<tr>
<td><strong>TOTAL 2016-2017 GRANT ACTIVITY</strong></td>
<td>$909,288</td>
</tr>
</tbody>
</table>
Education

Education is one of PWSA (USA)’s most comprehensive initiatives and includes informing the public as to what Prader-Willi syndrome is and the effect it has in people’s lives; familiarizing medical providers and school professionals with the needs of the PWS community; providing parents and caregivers with information to help them care for their child with PWS; and alerting legislators to the challenges faced by those affected by PWS. The white papers, brochures, presentations, videos, webinars, and online resources distributed by PWSA (USA) are life-changing, and often life-saving, sources of information for individuals and families affected by Prader-Willi syndrome.

Education: Ensuring Equal Access to All

Launched in February 2017, the online version of the Wyatt Special Education Advocacy Training series (WSEAT) is a free resource for parents and school professionals. Named in memory of David Wyatt, PWSA (USA)’s first Crisis Intervention and Family Support Counselor, the WSEAT:

• Includes 7 modules covering a range of special advocacy topics, including how to address PWS specific school challenges
• Allows viewers to watch one or all modules, focusing on the information most pertinent to them
• Includes downloadable resources for each module
• Includes free copies of From Emotions to Advocacy: The Special Education Survival Guide
• Provides recommended resources for additional learning

Education: National Convention with Multiple Conferences

The biennial PWSA (USA) National Convention is where researchers, medical professionals, educators, professional providers, and families come together with the common goal of improving the lives of individuals diagnosed with Prader-Willi syndrome. Programs are tailored to the needs and interests of professionals, families, and individuals with PWS, and planned social activities allow for networking and relationship building. Convention offerings include:

• General Conference
• Medical & Scientific Conference
• Professional Providers Conference
• Young Infant Program (YIP): 0 – 1 years old and 2 – 6 years old
• Youth & Adult Program (YAP): 7 years and up
• Sibling Program
• New Parent Mentors
• Chapter Leaders

Education: Information on the Go

Smart phone users can now carry important information and resources with them wherever they go by downloading the free PWSA (USA) app to their smart phone. Through the app, users can:

• Share important medical information with medical professionals
• Watch school videos with a child's IEP Team
• Stay up-to-date on the latest PWS research news
• Provide ER staff with key medical alerts during emergencies
• Conveniently explore family support and other available resources
Advocacy

Effective advocacy is essential to ensuring health, safety, and enhanced quality of life for those affected by Prader-Willi syndrome. As such, PWSA (USA) is committed to informing the PWS community of critical public policy issues and leveraging the power of grassroots supporters to effect change. Because Prader-Willi syndrome is considered a “rare disease,” extra effort is needed to raise awareness amongst the public and elected officials to guarantee the passage of legislation and regulations that help our community, and to defeat those that do not. PWSA (USA) also promotes individual-focused advocacy, providing training and assistance to parents and caregivers that empower them to advocate for their child or loved one with PWS.

Why It Matters

While the PWS community might not have the resources of groups with greater numbers, change is still possible. More and better education, effective communication, and strategic alliances with partners in the PWS and rare disease communities can help ensure:

• The “pre-existing condition” insurance status is maintained, and that health care remains affordable
• Individuals with PWS have access to appropriate residential and day programs to learn job skills
• Education and training is available for those affected by PWS
• Communities and schools are aware of and are equipped to work with individuals with PWS and their families
• New medicines and treatments are readily available

“Nothing so conclusively proves a man’s ability to lead others as what he does from day to day to lead himself.”

Thomas J. Watson
## Financials

**PRADER-WILLI SYNDROME ASSOCIATION (USA)**  
**STATEMENT OF ACTIVITIES**  
For The Year Ended December 31, 2016

### SUPPORT AND REVENUES

<table>
<thead>
<tr>
<th>Description</th>
<th>Unrestricted</th>
<th>Temporarily Restricted</th>
<th>Permanently Restricted</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Support</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Contributions</td>
<td>$ 945,065</td>
<td>$ 594,069</td>
<td>$ 26,050</td>
<td>$ 1,565,184</td>
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<tr>
<td>Fundraising contributions</td>
<td>386,356</td>
<td></td>
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<td>386,356</td>
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<tr>
<td>Conference income</td>
<td>1,692</td>
<td>30,000</td>
<td></td>
<td>31,692</td>
</tr>
<tr>
<td>Donated services</td>
<td>710,406</td>
<td></td>
<td></td>
<td>710,406</td>
</tr>
<tr>
<td><strong>Revenues</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Educational material sales</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Membership dues</td>
<td>18,496</td>
<td></td>
<td></td>
<td>18,496</td>
</tr>
<tr>
<td>Investment income/(loss)</td>
<td>77,795</td>
<td></td>
<td></td>
<td>77,795</td>
</tr>
<tr>
<td>Net assets released from restrictions</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Satisfaction of program restrictions</td>
<td>356,071</td>
<td>(356,071)</td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>TOTAL SUPPORT AND REVENUES</strong></td>
<td>$ 2,551,756</td>
<td>267,998</td>
<td>26,050</td>
<td>$ 2,845,804</td>
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### EXPENSES

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<tr>
<th>Description</th>
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<th>Temporarily Restricted</th>
<th>Permanently Restricted</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Program Services</strong></td>
<td></td>
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<td></td>
<td></td>
</tr>
<tr>
<td>Crisis intervention and support</td>
<td>355,715</td>
<td></td>
<td></td>
<td>355,715</td>
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<tr>
<td>Education</td>
<td>36,139</td>
<td></td>
<td></td>
<td>36,139</td>
</tr>
<tr>
<td>Medical intervention and support</td>
<td>385,958</td>
<td></td>
<td></td>
<td>385,958</td>
</tr>
<tr>
<td>New diagnosis support</td>
<td>79,206</td>
<td></td>
<td></td>
<td>79,206</td>
</tr>
<tr>
<td>Chapter, affiliated and local support</td>
<td>52,265</td>
<td></td>
<td></td>
<td>52,265</td>
</tr>
<tr>
<td>National conference</td>
<td>106,211</td>
<td></td>
<td></td>
<td>106,211</td>
</tr>
<tr>
<td>Advocacy, awareness, external collaborative relations</td>
<td>188,720</td>
<td></td>
<td></td>
<td>188,720</td>
</tr>
<tr>
<td>Research</td>
<td>577,129</td>
<td></td>
<td></td>
<td>577,129</td>
</tr>
<tr>
<td><strong>Total program services</strong></td>
<td>1,761,343</td>
<td></td>
<td></td>
<td>1,761,343</td>
</tr>
<tr>
<td><strong>Supporting services</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Administration</td>
<td>332,035</td>
<td></td>
<td></td>
<td>332,035</td>
</tr>
<tr>
<td>Fund development</td>
<td>128,821</td>
<td></td>
<td></td>
<td>128,821</td>
</tr>
<tr>
<td><strong>Total supporting services</strong></td>
<td>460,856</td>
<td></td>
<td></td>
<td>460,856</td>
</tr>
<tr>
<td><strong>TOTAL EXPENSES</strong></td>
<td>$ 2,222,199</td>
<td></td>
<td></td>
<td>$ 2,222,199</td>
</tr>
</tbody>
</table>

### CHANGE IN NET ASSETS

<table>
<thead>
<tr>
<th>Description</th>
<th>Unrestricted</th>
<th>Temporarily Restricted</th>
<th>Permanently Restricted</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>CHANGE IN NET ASSETS</strong></td>
<td>329,557</td>
<td>267,998</td>
<td>26,050</td>
<td>623,605</td>
</tr>
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</table>

### NET ASSETS AT BEGINNING OF YEAR

<table>
<thead>
<tr>
<th>Description</th>
<th>Unrestricted</th>
<th>Temporarily Restricted</th>
<th>Permanently Restricted</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>NET ASSETS AT BEGINNING OF YEAR</strong></td>
<td>535,395</td>
<td>1,070,297</td>
<td>133,884</td>
<td>1,739,576</td>
</tr>
</tbody>
</table>

### NET ASSETS AT END OF YEAR

<table>
<thead>
<tr>
<th>Description</th>
<th>Unrestricted</th>
<th>Temporarily Restricted</th>
<th>Permanently Restricted</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>NET ASSETS AT END OF YEAR</strong></td>
<td>$ 864,952</td>
<td>$ 1,338,295</td>
<td>$ 159,934</td>
<td>$ 2,363,181</td>
</tr>
</tbody>
</table>

Audit performed by Lawrence A. Kraujalis, CPA, P.A.
More than 6 of Every 7 Dollars Go to Programs & Services

2017 PWSA (USA) FUNDS

Research: Multiple Trials & Studies

Convention: Education, Family Support, & Advocacy

Advocacy & Collaboration

Fund Development & Administration

Medical

Awareness, Education, & Chapter Support

Family Support, Crisis, & New Diagnosis

The total of the investments represented on the chart is $2,251,076, and the percentage allocations are based on the annual Prader-Willi Syndrome Association (USA) audit done by an outside CPA firm. PWSA (USA) board members and staff work very hard to make sure that the great majority of your organization’s resource are used to help individuals, families, and our cause.
Who We Are:
People with Passion and Purpose

OFFICERS AND DIRECTORS
Chair - Michelle Torbert, Homestead, FL
Vice-Chair - James Koerber, Corydon, IN
Secretary - Julie Doherty, Tallahassee, FL
Treasurer - Denise Westenfield, Eden Prairie, MN
Leon D. Caldwell, Ph.D., Washington, D.C.
Mitch Cohen, Weston, CT
Thomas Conway, Albany, NY
Peter Girard, Townsend, TN
James Kane, Baltimore, MD
Steve Leightman, Cherry Hill, NJ
Rob Lutz, Bryn Mawr, PA
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