## **Testimonials**

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I return to Mexico with valuable learnings, new allies, and a renewed commitment to continue working in support of individuals with PWS. This experience marks a turning point in our actions and dedication. Without doubt, the outcomes of this conference will directly benefit our community in Mexico.



Josefina Pérez Roderiguez, Fundación María Jose, Mexico.

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The International Prader-Willi Conference is not just a platform for exchange but also a bridge connecting the global PWS community. This experience deepened my understanding of the importance of "unity, collaboration, and shared progress." In the future, China's PWS organization will continue upholding this philosophy, bringing more hope to patients and their families.



Lin Xiaojing, PWSA China.



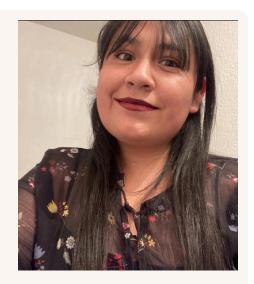
I was honored to speak about the Sub-Saharan African experience and engage in meaningful discussions with delegates about the way forward, including potential areas for collaboration. I am sincerely grateful to the Norway Prader-Willi Syndrome Organization and IPWSO for the travel fellowship that made my participation possible.



Dr Menbere Kahssay, Paediatric Endocrinologist, Kenya and Ethiopia.



Thank you IPWSO, for believing in the power of education, connection and opportunity. You have opened the door to an experience that will leave a lasting impact on my work and my heart. I look forward to reuniting at the next conference, equipped with even more tools, knowledge and enthusiasm to support people with PWS and their families. With heartfelt gratitude.



Claudia Rubio, Psychologist, Mexico.



A particular session on the 2<sup>nd</sup> day of the family program "PWS Around the World " Prof Tony Holland, Immediate Past President of IPWSO, inspired me greatly about the work that has been done in making sure that many countries are covered and efforts at reaching the unreached. It opened my eyes to the enormous work that still needs to be done especially concerning case finding in SubSaharan Africa (except South Africa) and optimal or near-optimal care with children with PWS and family.



Dr Elizabeth Eberechi Oyenusi, President of the African Society for Paediatric and Adolescent Endocrinology, Nigeria.







