

Parents left in the cold when it comes to kids with autism

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First-line health professionals must vastly improve their communication

and engagement with parents if they are to help address the growing prevalence of autism among children, say researchers from the University of South Australia.

Undertaking a [meta-synthesis of 22 international studies](#), researchers consolidated the voices of 1178 [parents](#) advocating for their [children](#) with autism, finding that parents feel ignored and dismissed by medical practitioners as they navigate initial concerns for their child, further investigations, and finally, a formal diagnosis of autism.

Researchers say that medical practitioners need to adopt a family-focused approach to ensure that parents' concerns, perspectives and observations are taken seriously so that their child has appropriate and timely access to early intervention services.

Autism spectrum disorder (ASD) is a persistent developmental disorder characterized by social difficulties, restricted or repetitive patterns of behavior, and impaired communication skills. The symptoms can range from mild to severe, with early signs often evident from early childhood.

Autism is one of the most prevalent developmental conditions among children, with one in 70 people in Australia on the spectrum, an estimated 40 percent increase over the past four years. Internationally, statistics are higher with one in 59 children on the spectrum.

UniSA lead researcher, Dr. Kobie Boshoff, says the parent advocacy role is critical and must be taken more seriously by medical practitioners.

"Parents are natural advocates for their child, making them an invaluable source of information when it comes to complex diagnoses for invisible disabilities like autism," Dr. Boshoff says.

"Yet parents are increasingly finding the diagnosis process overly

stressful and complicated.

"In this study, parents commonly reported their concerns for their child were not being heard or taken seriously by medical professionals. They said they felt confused, stressed and frustrated at the lack of support and understanding.

"They also reported lengthy delays in receiving a diagnosis for their child, as well as a variety of unsatisfactory explanations as alternatives to autism. As access to early intervention services is essential for improving the development outcomes of children with [autism](#), this too is unacceptable."

Dr. Boshoff says first-line [medical professionals](#) and service providers must recognize both the role of parents as advocates for their [child](#), and the importance of the parent-practitioner role, which can significantly impact future relationships with other professionals.

She says to build trust [medical practitioners](#) must reassess the way they talk and engage with parents.

"First line health professionals and diagnostic services must ensure emotional support is provided to parents throughout the diagnosis process, engaging parents as partners and taking their concerns seriously," Dr. Boshoff says.

"Autism spectrum disorder is a lifelong developmental condition. A positive experience in the early stages of [diagnosis](#) can deliver better relationships with future professionals, and most importantly, secure better outcomes for the children."

More information: Kobie Boshoff et al. A meta-synthesis of how parents of children with autism describe their experience of advocating

for their children during the process of diagnosis, *Health & Social Care in the Community* (2018). [DOI: 10.1111/hsc.12691](https://doi.org/10.1111/hsc.12691)

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