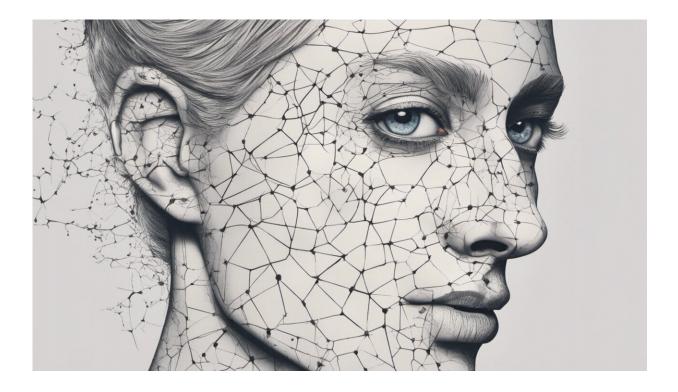


Treating suspected autism at 12 months of age improves children's language skills

July 17 2019, by Andrew Whitehouse, Kandice Varcin And Kristelle Hudry



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Therapies given to infants before they receive a diagnosis of autism may lead to important improvements in their language abilities, according to our new research published today in the journal Lancet Child and Adolescent Health.



Children with autism typically begin therapy after receiving a <u>diagnosis</u>, which usually doesn't occur until at least two years of age.

However, our new study suggests that starting therapy with 12-month-old infants who show early behavioural signs of autism may provide additional benefit.

Parents of toddlers who received six months of early therapy reported that their child understood an average of 37 more words, and spoke an average of 15 more words, than those who didn't receive the therapy.

How is autism diagnosed?

Autism is diagnosed on the basis of behavioural differences. These include difficulties in social communication and interaction, and the presence of restricted and repetitive behaviours or interests.

A diagnosis based on behaviour is an inherently subjective task. The dividing line between "typical" and "atypical" is often blurry and can lead to considerable debate.

These challenges led to the development in 2018 of new Australian guidelines for autism assessment and diagnosis, with the hope of addressing inconsistencies in autism diagnosis across Australia.

We don't tend to diagnose autism until two years of age at the very earliest. Two is the earliest age at which a diagnosis is relatively stable. In other words, when a child receives a diagnosis of autism at age two, they're also very likely to have that diagnosis if they're reassessed in later childhood. This is not necessarily the case for children younger than two.

In most health systems around the world, including in Australia, therapy for children with autism typically begins after receiving a diagnosis. This



is often a straight economic decision for a finite disability funding pool: the limited funding for therapies goes to children who need it most. And we can guarantee that children with a diagnosis of autism need these therapies.

But this model doesn't allow therapies to take advantage of the first two years of life. These are crucial years for <u>brain development</u>, and early therapies may be more effective in reducing child disability.

The theory is that if therapies are started early enough, particularly when the brain is so young, it might be possible to alter the trajectory of autism, or even prevent the development of the condition in some infants.

Our study

Our study is one of the first rigorous tests of this theory.

We recruited 103 infants from Perth and Melbourne, aged between 9 and 14 months, who were showing early behavioural signs of autism, such as not responding to their names, poor eye contact, and few social smiles.

Half the children were randomly assigned to receive an <u>intervention</u> that helps parents understand their infant's communication cues, and helps them respond in a way that promotes back-and-forth interactions.

By creating a socially rich environment around the infant that is tailored to their specific needs, we may help support the development of the neural pathways involved in language and social development.

The therapy is considered "low intensity" in the number of contact hours with a therapist (one hour per fortnight). This contrasts with "high



intensity" therapies for older children with a diagnosis of autism, which often require at least ten contact hours per week.

The other half of the group was randomly assigned to receive standard community care. This is often restricted to a parent information session, a few sessions with allied health professionals, or no intervention at all. This group served as the control group.

We then assessed the infants' development either side of a six-month therapy period.

What did we find?

The intervention group did not show a significant reduction in early autism behaviours compared with the control group. This finding is consistent with a <u>previous study</u> that found it's not easy to change the symptoms of autism.

However, after treatment, parents in the treatment group rated their infants as having better communication skills than those in the control group.

In the six-month therapy period, the treatment group improved in understanding an average of 37 more words and saying an average of 15 more words compared with the control group. Most children were not saying any words at the start of the therapy period, and so these language gains reported by parents are an important improvement.

This finding has an important caveat: parents could not be "blinded" to the therapy. It's therefore possible the finding reflects "rater bias," whereby parents whose children received the intervention hoped there was a developmental improvement, and rated their infants accordingly.



What does this mean for the NDIS?

This study is an important milestone in our understanding of how (and whether) to provide therapy to young infants who show early signs of autism.

Our findings suggest that these therapies do not reduce the core autism symptoms, but may improve language skills. These gains may be especially important for reducing long-term disability and helping each child reach their full potential.

The long-term test will be whether these improvements are still evident when the infants are older.

Few studies have found long-term developmental changes as a result of infant therapies, and we are currently in the process of reassessing the children in our study when they turn three.

If we find that pre-diagnostic <u>therapy</u> leads to longer-term developmental changes, this has important implications for health and disability services.

People on the <u>autism</u> spectrum currently represent <u>29% of all</u> <u>participants</u> enrolling in Australia's National Disability Insurance Scheme (NDIS).

A key plank of the NDIS is the Early Childhood Early Intervention pathway, which provides resources to support intervention for children up to seven years of age.

The pathway is very well conceived, but there are currently long <u>wait</u> <u>lists</u>.



The <u>federal government</u> recently <u>announced</u> a plan to resolve these delays, which indicates a recognition of the importance of early intervention for children with a confirmed diagnosis in reducing longterm disability in these children.

The follow-up of the <u>children</u> in the current study to three years of age will provide a litmus test as to whether very early intervention for <u>infants</u> should be an even higher priority.

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Provided by The Conversation

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