Abstract

Aim: This study examined the level of agreement between referral reason and diagnostic outcome for preschool aged children showing developmental delays and differences. In particular, the level of agreement when there was concern about autism spectrum disorder (ASD) was investigated.

Methods: Clinical records for children ($N=677$) assessed for developmental concerns at a multidisciplinary assessment clinic in Sydney, Australia, were examined. Referral reason and diagnostic outcome were compared. Referral sources and factors including age, developmental level, and language functioning, were explored for potential association with the consistency between referral reason and diagnosis. **Results:** Approximately 30% of children referred for developmental assessment with ASD concern were not diagnosed with ASD. Agreement between referral reason and diagnosis was similar regardless of referral source and child age. Of children where diagnostic outcome was discordant with referral reason, 24% had delayed language. Variability in cognitive or developmental profiles was common (64%), regardless of diagnostic outcome. **Conclusions:** Referral for ASD diagnostic assessments in the cases where ASD is not the primary cause for a child's difficulties increases pressure on assessment services, can delay access to ASD specific intervention services for children who do have ASD, and can contribute to parent anxiety unnecessarily. The current evidence that a substantial minority of children were referred for ASD diagnostic assessments when ASD was not the primary cause for the difficulties emphasises the value of adopting a developmental differences framework, focusing on dimensions of behaviour and functioning, rather than an autism-specific service structure.

Key Words: autism spectrum disorder; diagnosis

Key Points

What is already known on this topic:

- The prevalence of autism spectrum disorder (ASD) has increased over time as diagnostic criteria have broadened;
- This is associated with greater public concern and professional awareness of ASD, and with increased pressure on ASD diagnostic and intervention services.

What this paper adds:

- A substantial minority of children with ASD concern during preschool years do not meet ASD diagnostic criteria following assessment.
- There is a tendency to be overly concerned about ASD in children with language delays.

This is the author manuscript accepted for publication and has undergone full peer review but has not been through the copyediting, typesetting, pagination and proofreading process, which may lead to differences between this version and the Version of Record. Please cite this article as doi: [10.1111/jpc.13511](https://doi.org/10.1111/jpc.13511)

- A developmental differences framework, rather than autism specific service structure, may provide a more meaningful approach to conceptualising and assessing developmental differences.

Author Manuscript

There is increasing public awareness and concern about autism spectrum disorders (ASD).¹⁻³ This leads to early referral of children with atypical development or behaviour to assessment services seeking a diagnosis. Early referral can facilitate access to intervention programs, providing opportunities to promote learning and participation when appropriate.^{1, 4, 5} A recent study, however, reported that 39% of children referred to a regional ASD diagnostic clinic in the USA were not diagnosed with ASD following assessment.⁶ Thus, the heightened public awareness of ASD may also result in referrals to diagnostic centres for children who do not have ASD. An ASD diagnostic assessment for these children without ASD likely contributes to longer waiting times. This inevitably would delay diagnosis and access to ASD-specific intervention when indicated, and may also be associated with avoidable parental anxiety.

According to ICD-10⁷ and DSM-5⁸, the essential criteria for ASD are qualitative impairment in social interaction and communication, and the presence of restricted, repetitive and stereotyped patterns of behaviour, interests and activities. In the absence of biological markers for ASD, formulating a diagnosis requires comprehensive assessment of these areas, interpreted within the context of the child's medical and developmental history. Parent and teacher report is therefore important, together with direct child assessment of behaviour, language, and developmental or cognitive level.⁹⁻¹² Given the diagnostic complexity in ASD, inter-disciplinary team approaches are recommended across the UK, Europe, USA and Australia.^{9, 10, 12, 13} The diagnostic team should at least comprise a paediatrician, psychologist and/or speech pathologist, with input from occupational therapists, neurologists, and psychiatrists also recommended where possible.

XX XX XX Assessment Team (XXXXX) of Sydney Children Hospital Network (SCHN), Australia, provides diagnostic and assessment services for preschool children (aged 6 years and under) with delays in two or more unrelated domains of development. Concerns regarding ASD and/or other medical conditions may also be present. Consistent with practice guidelines, the diagnostic team consists of developmental paediatricians and psychologists. This study reviewed data for all children referred to XXXXX between 2008 and 2010 to better understand the relationships between referral sources, child characteristics, and assessment outcomes. During this period, referrals were received from parents, medical professionals (GPs, private and hospital paediatricians), nurses, allied health clinicians (occupational therapists, speech pathologists, physiotherapists, and psychologists) and non-medical professionals from the community (e.g., child care centre and preschool staff). Data for children referred with and without ASD concern were compared to examine the level of agreement between the referral concern and assessment outcome, and whether this varied according to referral

source and/or child characteristics (e.g., child age, language functioning, and cognitive or developmental ability).

METHODS

Eligible Children

Children ($N=677$) referred and assessed at XXXXX from 2008-2010 were identified from an existing clinical database. Study approval was granted by the SCHN Human Research Ethics Committee (LNR-2011-02-07) and was conducted in accordance with the provisions of the Declaration of Helsinki (2013).

Assessments

Developmental paediatricians and clinical psychologists completed child assessments and caregiver interviews. A clinical nurse specialist or social worker was present as required. When ASD concern was noted by referrers or identified by XXXXX clinicians, behavioural characteristics were elicited through parental interview and direct child observation. Reports from external informants were also collated and the Childhood Autism Rating Scale (CARS)¹⁴ completed. Clinical diagnoses were made based on all available information. Using DSM-IV criteria,¹⁵ the classifications were (1) autistic disorder, (2) pervasive developmental disorder-not otherwise specified (PDD-NOS), (3) Asperger's syndrome, or (4) no ASD. For the purposes of this study, children with diagnoses (1)-(3) were collectively categorised as ASD.

The Stanford-Binet Intelligence Scale – Fifth Edition (SBIS-5)¹⁶ and the Griffiths Mental Development Scales – Extended Revised (GMDS-ER)¹⁷ or 1996 Revised version (if aged under 2 years; GMDS-1996)¹⁸ evaluated each child's cognitive or developmental level. Children were classified as within the *Average* range (SBIS-5 or GMDS/GMDS-ER classifications within the Borderline range or higher), or with *Mild*, *Moderate*, or *Severe* delay (as per standard SBIS-5 or GMDS/GMDS-ER classifications). Children with substantial variability in subscale scores, preventing allocation of global category level, were classified as *Variable*. For the SBIS-5, a difference of greater than 10 points between verbal and non-verbal cognitive abilities, and/or non-uniform profile across subscale classifications (as described above), was defined as *Variable*. For the GMDS-ER or GMDS-1966, individuals with a disparity

between domains greater than one category (e.g., average fine motor development but moderate or severe delay in communication and language) were classified as *Variable*.

Data Extraction

Data were extracted from the XXXXX database. Data accuracy was verified by reviewing clinical records. Children were divided into two groups based on the referral reason: (1) ASD concern, including children with ASD features noted by referrers and others with a previous ASD diagnosis by a paediatrician or psychologist, and (2) without ASD concern.

The data included age at date of referral and assessment; referral sources; referral reason (with/without ASD concern); diagnostic outcome; assessment tools used and test scores; and known associated conditions, including language delay, congenital abnormality, sensory (hearing or vision) impairment, and behavioural disorders (attention deficit hyperactivity disorder, ADHD; oppositional defiant disorder, ODD).

Of the 677 children referred during the study period, 300 were referred and assessed in 2008, 186 in 2009, and 191 in 2010. The reduction in the number of assessments following 2008 was due to team relocation, involving re-housing and reduced availability of medical and psychological staff.

Statistical analysis

De-identified data were analysed using SAS version 9.3. Continuous variables were described by mean and range, or median and inter-quartile range. Categorical variables were described by frequencies and percentages, and compared using chi-square tests for independent groups. McNemar's test was used to examine the level of agreement between the reason for referral and ASD diagnosis at assessment, and a Kappa coefficient was calculated. There was no adjustment made for multiple statistical comparisons.

RESULTS

Child Characteristics

Of the 677 children reviewed by XXXXX from 2008-2010, 35% were referred with ASD concern. This was similar across all three years, with 37%, 31% and 34% referred with ASD concerns in 2008, 2009, and 2010, respectively.

There was evidence of an association between gender and ASD diagnosis ($P=0.0006$), with males ($n=151$) more commonly diagnosed with ASD than females ($n=34$); male to female ratio approximately 4:1. Among the non-ASD group ($n=492$), the gender ratio approximated 2:1 (M:F=336:156).

Age at Referral and Diagnosis

The mean age at referral was 39 months (range 14-76 months). The majority of children were referred between 24-59 months, with a similar distribution of age of referral for children referred with and without ASD concerns (Figure 1).

(Insert Figure 1)

For children referred with ASD concern, the length of time between referral and ASD diagnosis ranged from 1-37 months ($Md=5.6$, $IQR=3-7$ months). ASD was diagnosed in 185 (27%) children aged between 19 to 126 months; median age of diagnosis was 44 months ($IQR=35-50$ months). Age of assessment for children not diagnosed with ASD was similar ($Md=47$ months, $IQR=35-54$).

Agreement between Referral Reason and Diagnosis

The referral reason and diagnosis agreed in 592 (87%) of cases (Table 1). The kappa statistic (proportion of agreement above what would be expected by chance) was 0.71. Of the 85 for whom referral reason disagreed with diagnosis, 67 (79%) were referred with ASD concerns but not diagnosed with ASD, whereas the remaining 18 (21%) were referred with other concerns but diagnosed with ASD ($P<0.0001$, McNemar's test).

(Insert Table 1)

There was no evidence that the agreement between referrer's concern and the positive ASD diagnosis differed between age groups ($P=.11$; Table 2).

(Insert Table 2)

When consistency between referral reason and diagnostic outcome was considered for the full sample, the level of agreement was at least 80% for all referral sources ($P=0.42$). Table 3 displays the agreement between reason for referral and diagnostic outcome for each referral source, with the sample differentiated according to referral reason. Of the 234 children referred with concern about ASD, the percentage of cases with confirmed diagnosis was similar across all referral sources.

(Insert Table 3)

There was no evidence that the level of agreement between referral reason and diagnosis differed according to the developmental level of the child ($P=0.20$). As displayed in Table 4, 64% ($n=435$) of children referred to XXXXX had variation of ability for different aspects of their IQ or developmental profiles. Of these, 39% ($n=171$) were referred with ASD concern, with 30% ($n=129$) meeting diagnostic criteria for ASD.

(Insert Table 4)

The number and proportion of children with a language delay, known congenital condition, sensory (hearing or vision) impairment, or ADHD/ODD are shown in Table 5, according to referral reason and diagnosis.

(Insert Table 5)

The level of agreement between referral reason and diagnosis differed according to the presence of associated conditions ($P<0.0001$). Of 324 children with no associated conditions, agreement between referral reason and diagnostic outcome was 89% ($n=288$). Agreement between referral

reason and diagnostic outcome was 75% ($n=126$) in children with language delay ($n=168$). Of the 42 children with language delay where the referral disagreed with diagnosis, 40 (95%) were referred with ASD concern but did not meet diagnostic criteria, while only two children (5%) referred without ASD concern were diagnosed with ASD following assessment.

DISCUSSION

In this study, a substantial minority (approximately 30%) of children referred for developmental assessment with ASD concern did not meet ASD diagnostic criteria, with similar patterns of agreement between referral reason and diagnosis regardless of child age and referral source. Only a small proportion (4%) of children referred without any ASD concerns were diagnosed with ASD. This indicates that, when developmental concerns were identified and further assessment was sought, few children meeting ASD criteria had been missed by referrers. Children referred with concern about ASD who did not meet diagnostic criteria frequently had delayed language development. Variability in cognitive or developmental profiles according DQ or IQ assessments was common, with no evidence of difference between children diagnosed with ASD or other disabilities.

Recent estimates of the Australian ASD prevalence approximate 2%.¹⁹ This has increased over time in association with changing diagnostic criteria and inclusion of less severe differences for each criterion. The resultant wide variability in the clinical presentation of ASD can add to the complexity of diagnostic decision-making. Diagnostic complexity and growing evidence supporting benefits of early diagnosis and intervention likely steers clinicians towards referring children for more comprehensive assessment when uncertain. In Australia, definitive ASD diagnosis prior to age six is a pre-requisite for accessing national funding. This has created a potential conflict for clinicians, where a diagnosis may be made to support a child's access to early intervention services, despite there being diagnostic uncertainty.²⁰ Greater community awareness also facilitates earlier detection and referral for ASD diagnostic assessments. In doing so, however, this increases pressure on diagnostic and intervention services.

Increasingly, there is a focus on ASD surveillance and screening from early childhood ages.²¹⁻²³ This promotes early referral for diagnostic assessment when ASD is suspected, and timely access to intervention services when appropriate. However, there is debate about whether there is sufficient evidence for this approach, with potential negative impacts acknowledged for families in which children were falsely identified.^{24, 25} Our results showed a substantial minority of children referred for

ASD diagnostic assessment did not meet diagnostic criteria. This is consistent with recent research in the USA, where 39% of referrals for ASD diagnostic assessments did not have ASD.⁶ This places greater burden on already pressed diagnostic services, potentially delaying access to assessments and intervention for children who do have ASD. ASD diagnostic assessment when ASD is not the underlying cause of the child's difficulties can also increase parental anxiety. It is therefore important to carefully consider factors that may contribute to inappropriate referrals for ASD diagnostic assessments.

We have found that language delay, and not the level of global or cognitive development, may be an important clinical marker contributing to referral for ASD assessment in cases where ASD diagnostic criteria are not subsequently met. This suggests that referrers are overly concerned about ASD in children with language delays. While delayed language development has commonly been identified as one of the earliest markers raising concerns about ASD,²⁶⁻²⁸ such difficulties are not specific to ASD,²⁹ and 60-80% of children with delayed expressive language as toddlers 'catch up' by preschool age.³⁰⁻³² Thus, in cases where language delay is a primary referral concern, seeking expert speech and language assistance early, and adopting a 'watch-and-wait' approach when language differences are mild, may be justified.

This study raises broader considerations regarding service configuration of developmental assessment services. It is not unusual for assessment and intervention services to be separate, and only loosely connected to primary care services. Increasingly, autism specific services are also developing. Evidence that approximately 30% of ASD referrals do not meet ASD diagnostic criteria suggests these approaches may be problematic. There is potential to focus on ASD specific behaviours, or lack thereof, rather than interpreting functioning within a broader developmental differences framework. There is also the risk of assessment findings not being optimally linked to delivery of care and intervention.³³ Emerging evidence supports the importance of service structure emphasising links between multidisciplinary assessments and secondary care, with parents valuing information regarding their child's strengths and weakness gained from multidisciplinary assessments, but finding recommendations from sole paediatricians more useful.³⁴

This study offered the unique opportunity to compare the consistency between referral reason and diagnostic outcome across a number of referral sources. To the author's knowledge, this is the first study to explore Australian clinical data in this way. Evidence that the level of agreement was

comparable between paediatricians and other referral sources, including non-professionals, was somewhat unexpected. This is not taken to suggest that paediatricians were underperforming in this regard; as paediatricians are able to diagnose ASD independently, without necessarily needing input from external assessment agencies, this may reflect a tendency for paediatricians to refer in cases where complexity and diagnostic uncertainty is greatest.

Limitations

Findings of this study relate to children aged 6 years and under, and cannot be extrapolated to older age groups. The data were gathered from a single diagnostic service only; the level of agreement between referral reason and diagnosis may differ to that found in other clinics. Data was limited to information available at the time of referral and initial assessment only. Given recent reports of lack of stability in ASD diagnosis over time,³⁵ repeat assessment during later childhood years would have been informative.

Conclusions

Current evidence that a substantial minority of children suspected of having ASD did not receive a diagnosis indicates that attempts to differentiate between diagnoses is challenging for pre-schoolers prior to comprehensive assessment. Referral for ASD diagnostic assessments in cases where this is not the underlying cause for the presenting difficulties places increasing pressure on services, results in longer wait times, and creates barriers to accessing intervention services in instances where a diagnosis is required for entry. Similarly, referral to autism-specific early intervention services prior to comprehensive assessment would have over-burdened those services, and potentially prevented access to them for children with ASD. Moving forward, a developmental differences approach where children are referred for comprehensive assessment because of functional or behavioural concerns could prevent unnecessary anxiety for families and reduce waiting times for assessments, if in turn that approach reduced the need for ASD specific assessments for some children.

Acknowledgments

We are grateful to psychologists XX XX and XX XX for their assistance in data entry. Special thanks to Dr XX XX for her support during the data verification process.

REFERENCES

1. Glasson EJ, MacDermott S, Dixon G, Cook H, Chauvel P, Maley-Berg A, et al. Management of assessments and diagnoses for children with autism spectrum disorders: The Western Australian model. *Med J Aust*. 2008;**188**(5):288-91.
2. Kawamura Y, Takahashi O, Ishii T. Reevaluating the incidence of pervasive developmental disorders: Impact of elevated rates of detection through implementation of an integrated system of screening in Toyota, Japan. *Psychiatry Clin Neurosci*. 2008;**62**(2):152-9.
3. Mayor S. Medical Research Council review sets research agenda for autism. *BMJ*. 2002;**324**(7328):10.
4. Landa RJ, Kalb LG. Long-term outcomes of toddlers with autism spectrum disorders exposed to short-term intervention. *Pediatrics*. 2012;**130** Suppl 2:S186-90.
5. Ogiwara H, Takahashi O. The developmental characteristics of autistic children receiving very early intervention. *Jpn J Child Adol Psy*. 2005;**46**(Supplement):40-60.
6. Monteiro SA, Spinks-Franklin A, Treadwell-Deering D, Berry L, Sellers-Vinson S, Smith E, et al. Prevalence of autism spectrum disorder in children referred for diagnostic autism evaluation. *Clin Pediatr (Phila)*. 2015;**54**(14):1322-7.
7. World Health Organisation. *The ICD-10 Classification of Mental and Behavioural Disorders: Diagnostic Criteria for Research*. World Health Organization, editor. Geneva: World Health Organization; 1993.
8. American Psychiatric Association. *Diagnostic and Statistical Manual of Mental Disorders: DSM-5*. 5th ed. ed. American Psychiatric A, American Psychiatric Association DSMTF, American Psychiatric P, editors. Arlington, VA: Arlington, VA : American Psychiatric Association; 2013.
9. Autism Spectrum Disorders Special Interest Group. *Best practice guidelines for the assessment and diagnosis of autistic spectrum disorders for children and adolescents (birth to 18 years)*2012.
10. California Department of Developmental Services. *Autism Spectrum Disorder: Best Practice Guidelines for Screening, Diagnosis and Assessment*. Sacramento, CA: California Department of Developmental Services; 2002.
11. Lord C, Jones Rebecca M. Annual research review: Re-thinking the classification of autism spectrum disorders. *J Child Psychol Psychiatry*. 2012;**53**(5):490-509.
12. Western Australian Autism Diagnosticians' Forum Inc. *Autism Spectrum Disorders: Assessment Process in Western Australia*. Western Australia: Western Australian Autism Diagnosticians' Forum Inc.; 2012.

13. National Collaborating Centre for Women's and Children's Health. *Autism: Recognition, Referral and Diagnosis of Children and Young People on the Autism Spectrum*. London: RCOG Press, National Institute for Health and Clinical Excellence: Guidance; 2011.
14. Schopler E, Reichler RJ, DeVellis RF, Daly K. Toward objective classification of childhood autism: Childhood Autism Rating Scale (CARS). *J Autism Dev Disord*. 1980;**10**(1):91-103.
15. American Psychiatric Association. *Diagnostic and Statistical Manual of Mental Disorders: DSM-IV-TR*. 4th ed., text revision 2000. ed. American Psychiatric A, American Psychiatric Association. Task Force on D-I, editors. Washington, DC: Washington, DC : American Psychiatric Association; 2000.
16. Roid GH. *Stanford-Binet Intelligence Scales for Early Childhood, Fifth Edition*. Sink CA, Eppler C, Vacca JJ, editors. Rolling Meadows, IL: Riverside Publishing; 2005 //.
17. Luiz DM, Foxcroft CD, Povey J-L. The Griffiths Scales of Mental Development: A factorial validity study. *S Afr J Psychol*. 2006;**36**(1):192-214.
18. Huntley M. *The Griffiths Mental Development Scales: From Birth to 2 Years*: Association for Research in Infant and Child Development (ARICD); 1996.
19. Randall M, Sciberras E, Brignell A, Ihsen E, Efron D, Dissanayake C, et al. Autism spectrum disorder: Presentation and prevalence in a nationally representative Australian sample. *Aust N Z J Psychiatry*. 2015.
20. Skellern C, Schluter P, McDowell M. From complexity to category: Responding to diagnostic uncertainties of autistic spectrum disorders. *J Paediatr Child Health*. 2005;**41**(8):407-12.
21. Barbaro J, Halder S. Early identification of autism spectrum disorder: Current challenges and future global directions. *Curr Dev Disord Rep*. 2016;**3**(1):67-74.
22. Barbaro J, Ridgway L, Dissanayake C. Developmental surveillance of infants and toddlers by maternal and child health nurses in an Australian community-based setting: Promoting the early identification of autism spectrum disorders. *J Pediatr Nurs*. 2011;**26**(4):334-47.
23. Veenstra-VanderWeele J, McGuire K. Rigid, inflexible approach results in no recommendation for autism screening. *JAMA psychiatry*. 2016;**73**(4):327-8.
24. Siu AL, Bibbins-Domingo K, Grossman DC, Baumann LC, Davidson KW, Ebell M, et al. Screening for autism spectrum disorder in young children: US Preventive Services Task Force recommendation statement. *Jama*. 2016;**315**(7):691-6.
25. Al-Qabandi M, Gorter JW, Rosenbaum P. Early autism detection: Are we ready for routine screening? *Pediatrics*. 2011;**128**(1):e211-e7.
26. De Giacomo A, Fombonne E. Parental recognition of developmental abnormalities in autism. *Eur Child Adolesc Psychiatry*. 1998;**7**(3):131-6.

27. Herlihy L, Knoch K, Vibert B, Fein D. Parents' first concerns about toddlers with autism spectrum disorder: Effect of sibling status. *Autism*. 2015;**19**(1):20-8.
28. Noterdaeme M, Hutzelmeyer-Nickels A. Early symptoms and recognition of pervasive developmental disorders in Germany. *Autism*. 2010;**14**(6):575-88.
29. Tager-Flusberg H, Paul R, Lord C. Language and Communication in Autism. In: Volkmar FR, Paul R, Klin A, Cohen DJ, editors. *Handbook of Autism and Pervasive Developmental Disorders*: John Wiley & Sons; 2005. p. 335-64.
30. Ellis EM, Thal DJ. Early language delay and risk for language impairment. *Perspect Lang Learn Educ*. 2008;**15**(3):93-100.
31. Dale PS, Price TS, Bishop DV, Plomin R. Outcomes of early language delay: I. Predicting persistent and transient language difficulties at 3 and 4 years. *Journal of speech, language, and hearing research : JSLHR*. 2003;**46**(3):544-60.
32. Petruccelli N, Bavin EL, Bretherton L. Children with specific language impairment and resolved late talkers: Working memory profiles at 5 years. *Journal of speech, language, and hearing research : JSLHR*. 2012;**55**(6):1690-703.
33. Williams K, Prior M, Bartak L, Woolfenden S, Roberts J, Rodger S. Autism in context 2: Assessment, intervention and services in Australia. *J Paediatr Child Health*. 2014.
34. Mittal R, Sciberras E, Sewell J, Efron D. Assessment of children with learning and behavioural problems: Comparison of a multidisciplinary and sole paediatrician model. *J Paediatr Child Health*. 2014;**50**(2):135-40.
35. Woolfenden S, Sarkozy V, Ridley G, Williams K. A systematic review of the diagnostic stability of autism spectrum disorder. *Res Autism Spect Dis*. 2012;**6** (1):345-54.

Tables

Table 1

Agreement between reason for referral and final diagnosis

	ASD Diagnosis	Other Diagnosis	Total
Referral Reason	<i>n</i>	<i>n</i>	<i>N</i>
ASD referral	167	67	234
Other referral reason	18	425	443
Total	185	492	677

Table 2

Number and percentage with agreement between referral reason and diagnosis by age

Age	Referral: ASD Concern				Referral: No ASD Concern			
	<i>n</i> =234				<i>n</i> =443			
	ASD Diagnosis		Other Diagnosis		ASD Diagnosis		Other Diagnosis	
	<i>n</i> =167		<i>n</i> =67		<i>n</i> =18		<i>n</i> =425	
	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%
<24 months	3	75	1	25	1	3	35	97
24-35 months	36	72	14	28	8	10	76	90
36-47 months	63	78	18	22	4	3	117	97
48-59 months	51	68	24	32	3	2	162	98
>= 60 months	14	58	10	42	2	5	35	95

Table 3

Referral sources and XXXXX diagnosis for children with and without ASD concern

Referral Sources	Referral: ASD Concern				Referral: No ASD Concern			
	n=234				n=443			
	ASD Diagnosis		Other Diagnosis		ASD Diagnosis		Other Diagnosis	
	n=167		n=67		n=18		n=425	
	n	%	n	%	n	%	n	%
Paediatrician	77	70	33	30	6	5	121	95
ADHC	7	100	0	0	2	6	29	94
Psychologist	3	100	0	0	0	0	0	0
Carers	25	68	12	32	4	4	86	96
Community RN	6	100	0	0	2	9	21	91
DOCS	3	75	1	25	0	0	16	100
EI therapy team	29	71	12	29	2	2	85	98
GPs	2	50	2	50	1	6	15	94
Preschool staff	15	68	7	32	1	2	52	98

ADHC: State Department of Ageing, Disability & Home Care providing therapy and case work services. **DOCS:** State Department of Community Services. **EI therapy team:** Early Intervention services providing physiotherapy, occupation and speech therapies.

Table 4

Level of agreement between referral concern, XXXXX diagnosis and child intellectual/developmental level

	Referral: ASD Concern				Referral: No ASD Concern			
	<i>n</i> =234				<i>n</i> = 443			
	ASD		Other		ASD		Other	
	Diagnosis		Diagnosis		Diagnosis		Diagnosis	
	<i>n</i> =167		<i>n</i> =67		<i>n</i> =18		<i>n</i> =425	
Intellectual/ Developmental Level	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%
Average	5	36	9	64	1	2	42	98
Mild	15	62	9	38	1	2	51	98
Moderate	5	45	6	55	1	4	25	94
Severe	6	100	0	0	3	5	53	95
Variable profile	129	75	42	25	12	5	252	95
Not Assessed	7	87	1	13	0	0	2	100

Author Manuscript

Table 5

Associated condition by reason for referral and diagnosis

	Referral: ASD Concern				Referral: No ASD Concern			
	<i>n</i> =234				<i>n</i> =443			
	ASD Diagnosis		Other Diagnosis		ASD Diagnosis		Other Diagnosis	
	<i>n</i> =167		<i>n</i> =67		<i>n</i> =18		<i>n</i> =425	
Associated Condition	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%	<i>n</i>	%
None	132	79	23	34	13	72	156	37
Language Delay	25	15	40	60	2	11	101	24
Congenital	9	5	2	3	2	11	165	39
Special Senses	1	1	0	0	1	6	3	1
ADHD or ODD	0	0	2	3	0	0	0	0

Language Delay: specific language and communication delay according to Griffith's Mental Developmental Scales, as defined by score <70 on Scale C only (Language and Communication scale). **Special senses:** Hearing/sight impaired under one associated condition. **Congenital:** all known syndromes, congenital abnormality, epilepsy and prenatal insults. **ADHD/ODD:** attention deficit hyperactive disorder/oppositional defiant disorder.

Figure Legends

Figure 1. Age distribution at time of referral and diagnosis.

Author Manuscript

Figure 1_Blinded.tiff

Original Article

Agreement between concern about autism spectrum disorder at the time of referral and diagnosis, and factors associated with agreement

Bee Hong Lo,¹ Felicity Kloppe,² Elizabeth H Barnes,³ and Katrina Williams^{2,4,5}

¹Parramatta Early Childhood Assessment Team, Children's Hospital Westmead, Sydney, Australia

²Murdoch Childrens Research Institute, Melbourne, Australia

³NHMRC Clinical Trials Centre, University of Sydney, Sydney, Australia

⁴Developmental Medicine, The Royal Children's Hospital, Melbourne, Australia

⁵Department of Paediatrics, University of Melbourne, Melbourne, Australia

Correspondence

Dr Bee Hong Lo, Parramatta Early Childhood Assessment Team (PECAT), Children's Hospital Westmead, Locked Bag 4001, Westmead, NSW 2145. E: bee.lo@health.nsw.gov.au. T: (02) 9891 7200. F: (02) 9891 7222

Conflict of Interest

None declared

Minerva Access is the Institutional Repository of The University of Melbourne

Author/s:

Lo, BH;Klopper, F;Barnes, EH;Williams, K

Title:

Agreement between concern about autism spectrum disorder at the time of referral and diagnosis, and factors associated with agreement

Date:

2017-08-01

Citation:

Lo, B. H., Klopper, F., Barnes, E. H. & Williams, K. (2017). Agreement between concern about autism spectrum disorder at the time of referral and diagnosis, and factors associated with agreement. JOURNAL OF PAEDIATRICS AND CHILD HEALTH, 53 (8), pp.742-748.
<https://doi.org/10.1111/jpc.13511>.

Persistent Link:

<http://hdl.handle.net/11343/292746>