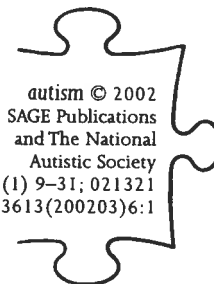


The CAST (Childhood Asperger Syndrome Test)

Preliminary development of a UK screen for mainstream primary-school-age children

autism © 2002
SAGE Publications
and The National
Autistic Society
Vol 6(1) 9-31; 021321
1362-3613(200203)6:1



FIONA J. SCOTT University of Cambridge, UK

SIMON BARON-COHEN University of Cambridge, UK

PATRICK BOLTON University of Cambridge, UK

CAROL BRAYNE University of Cambridge, UK

ABSTRACT The article describes a pilot and follow-up study of the preliminary development of a new tool to screen for Asperger syndrome (AS) and related social and communication conditions (the Childhood Asperger Syndrome Test, CAST) in children aged 4–11 years, in a non-clinical setting. In the pilot study, parents of 13 children with AS and of 37 typically developing children completed the CAST. There were significant differences between the AS and typical sample means. The pilot was used to establish preliminary cut-off scores for the CAST. In the main study, parents of 1150 primary-school-age children were sent the CAST, and 174 took part in the full data analysis. Results suggest that compared with other tools currently available, the CAST may be useful for identifying children at risk for AS and related conditions, in a mainstream non-clinical sample. Further research is ongoing.

KEYWORDS

Asperger
syndrome;
autism
spectrum;
epidemiology;
screening

ADDRESS Correspondence should be addressed to: FIONA J. SCOTT, Autism Research Centre, Department of Psychiatry, University of Cambridge, 18b Trumpington Road, Cambridge CB2 2AH, UK

Background

Classic autism is now routinely identified by the age of 3 years (Howlin and Moore, 1997), and can be identified by as young as 18 months of age (Baird et al., 2000; Baron-Cohen et al., 1996). However, other conditions on the autism spectrum are not as easily identified, even though the prevalence of autism spectrum conditions may be around 60 per 10,000 (Baird

giving varying rates of true and false positives (children who score as AS and who really do have AS, versus children who score as AS but who do not have AS). For parent ratings, the optimal ASSQ cut-off score derived was 19, giving a true positive rate of 62 percent (false positive 10 percent). Teacher ratings had an optimal cut-off of 22, leading to a slightly better true positive rate of 70 percent (false positive 9 percent).

Two other tools with some level of validation are as follows. The first is the Social Communication Questionnaire (SCQ, previously the Autism Spectrum Questionnaire or ASQ: Kazak-Berument et al., 1999), which has been developed to differentiate PDD from non-PDD children in a clinical sample but which, like the ASSQ, has not been validated on a non-clinical population. Indeed, the authors suggest that it would not be a good screening tool for use at a population level (Bailey, 2001). The SCQ does not differentiate AS from other autism spectrum conditions, or differentiate between different 'points' on the autism spectrum.

The second is the Pervasive Developmental Disorders Questionnaire (PDDQ: Baird et al., 2000), which has been developed and tested with a younger sample of children (age 5 years), and asks developmentally appropriate questions for that age. The PDDQ has advantages in that it addresses the broader autism spectrum, but its specificity does not appear to be well developed as yet. Whilst the PDDQ has been piloted with 40 children already diagnosed with AS, and 37 of those children (92.5 percent) failed five or more of the key items, its sampling in a broader population has limitations. Out of a sample of 7766 5-year-old children, 63 scored above cut-off on the PDDQ. Assessments were conducted on 29 of those children, with 11 (37.9 percent) meeting criteria for autism or pervasive developmental disorder (Baird et al., 2000). The PDDQ may suffer from being a very brief screen, with only 18 questions, of which nine are AS-relevant.

The National Screening Committee (1998) recommends that screening for identification of as yet unidentified cases should only be conducted where it can be shown that earlier identification coupled with treatment or intervention has some beneficial outcome on that population. Additionally, the NSC recommends that screening tools should strive for as high a level of sensitivity, specificity and positive predictive value as possible. That is, a tool should identify as many of the true cases as possible in a population, without picking up too many non-cases (those who score positive on the screen but are later shown not to have the specified condition), and it should be the case that the likelihood of having the specified condition if one is positive on the screen is high. It is certainly felt that there is a need for development of a UK screener for AS and the broader autism spectrum, particularly as this is the area where there is the greatest current shortage of knowledge coupled with increasing demand (Howlin, 2000), and early

and related social and communication difficulties in mainstream primary-school-age children (4–11 years) in the UK.

Pilot study

Participants

The participants were 13 children already diagnosed with Asperger syndrome (AS) or autism (age 3–9 years, mean 6:11, SD 1:11), and 37 normally developing control children (age 6–9 years, mean 6:7, SD 0:7).

Screen

The screening instrument being developed is the Childhood Asperger Syndrome Test (CAST). It is based on a variety of behavioural descriptions of the ICD-10 (World Health Organization, 1993) and DSM-IV core features of the autism spectrum (social impairments, communication impairments and repetitive or stereotyped behaviours). Some items in the CAST were based on items appearing in two other screening tools: the Pervasive Developmental Disorders Questionnaire (PDDQ: Baird et al., 2000) and the Asperger Syndrome Screening Questionnaire (ASSQ: Ehlers et al., 1999).² The PDDQ and the ASSQ were not considered appropriate tools for screening of AS in primary-school-age children for the reasons outlined in the background – namely that the ASSQ has only been validated on a clinical sample, and the PDDQ is itself in very early stages of development and has not been designed to focus on Asperger syndrome.

The AS-relevant questions in the CAST were designed to cover as wide a range of behaviours as possible, so as to facilitate detecting the high-functioning end of the autism spectrum. The CAST has 37 items in total, of which 31 are key items contributing to a child's total score. The remaining six items are control questions on general development and these are not scored. The six control items are items 3, 4, 12, 22, 26 and 33. The maximum a child can score is 31. The CAST is shown in Appendix 1.

Procedure

The CAST was completed by the parents of 13 children with an existing diagnosis of AS, and by the parents of 37 normally developing children aged 6–9 years attending a mainstream primary school outside the region. Parents were informed that we were developing a new screening tool to identify possible cases of AS and related social communication difficulty in primary-school-age children, and that their input would help us establish provisional cut-off scores and understanding of 'typical' scoring on the CAST. Additionally, for the pilot stage parents were invited to complete the CAST only if there were no special needs requirements

Table 1 Number of children scoring at or above each point on the CAST (pilot study)

Total CAST score	Number (and %) of children with AS	Number (and %) of control children
0	13 (100)	37 (100)
1	13 (100)	34 (92)
2	13 (100)	31 (86)
3	13 (100)	29 (73)
4	13 (100)	22 (64)
5	13 (100)	15 (35)
6	13 (100)	10 (27)
7	13 (100)	6 (16)
8	13 (100)	6 (16)
9	13 (100)	6 (16)
10	13 (100)	5 (14)
11	13 (100)	4 (11)
12	13 (100)	4 (11)
13	13 (100)	3 (8)
14	13 (100)	0
15	13 (100)	0
16	10 (77)	0
17	10 (77)	0
18	9 (69)	0
19	7 (46)	0
20	6 (38)	0
21	6 (38)	0
22	6 (38)	0
23	5 (31)	0
24	5 (31)	0
25	5 (31)	0
26	5 (31)	0
27	3 (15)	0
28	3 (15)	0
29	3 (15)	0
30	2 (8)	0
31	1 (8)	0

Discussion of pilot study

As can be seen, all of the AS sample scored equal to or greater than 15, whilst none of the controls did so. These results suggested that choosing a provisional cut-off of 15 for our preliminary study would not generate any false positives, or lead to many (if any) cases needing an assessment for a possible social and communication condition, in a typical mainstream

primary-age population. Neither would it risk generating many (if any) false negatives (i.e. missing too many possible positive cases of AS). Although three of the CAST questions showed no significant differences between the two groups in the pilot study, and one just missed significance, we decided initially to retain all the questions for the main study. This decision was made in part because of the small sample sizes involved in the pilot study. The results show that the majority of the CAST questions differentiate very clearly between the AS/HFA group and the normally developing controls, and it may be that one could safely drop the non-significant questions. This issue would be addressed following the main study.

Main study

Participants

The participants were 199 mainstream primary-school-age children (age 4–11 years, mean 8:1, SD 1:9).

Procedure

The CAST was sent to the parents of 1150 children age 4–11 attending mainstream primary schools in Cambridgeshire. The schools involved were informed of the purpose of the study – that we were developing a potential new tool to screen for possible cases of AS and related social communication difficulties in primary-school-age children – and the CAST was distributed via schools to parents with an accompanying explanatory letter (Appendix 2). Parents were informed that the questionnaire they had received was part of a study exploring social and communication development in primary-school-age children, looking at the differences seen and the difficulties some children have. It was explained that a small percentage of children have severe difficulties in social communication, and that these children might have a condition such as Asperger syndrome. Parents were asked to indicate if they would be willing to be approached by the research team at a later date for face-to-face assessments, and it was made clear that this was not necessarily an indication of a difficulty on their child's part. Ethical agreement for the study was established on the basis that we would indicate to a family if there was a problem and the family were concerned about their child's development, and that the family would be counselled about further action to take as necessary. Children clearly requiring further clinical assessment or intervention were thus linked into appropriate services. The research team has strong links with child clinical services in the area, and these services were readily available when required.

Additionally, the schools involved were visited by the research team,

or at or above cut-off on both instruments, were assessed using the ADOS-G (Lord et al., 1999) or the ADI-R (Lord et al., 1994). Additionally, we assessed those children scoring near cut-off (up to 3 points below) on either screen.

Those children within this sample who had not already received a definitive clinical diagnosis of autism spectrum disorder were assessed by the first author using either the ADI-R or the ADOS-G. In practice, the majority of cases were assessed using the ADOS-G, as the ADI-R takes around 3 hours to complete. All assessments were videotaped with consent. These assessments were not to provide clinical diagnoses, as the ADI-R or ADOS-G used alone cannot provide diagnoses. Rather, the assessments were to establish whether children met research criteria on established and standardized tools for autism spectrum condition. However, as has been outlined earlier, where a child met criteria for an autism spectrum condition on the ADOS-G or ADI-R the family was given feedback about the possibility of requiring further clinical assessment if they wished, and were put in touch with clinical services as appropriate.

Whilst the first author is fully trained in the use of the ADI-R and ADOS-G, reliability was checked on a random sample of cases via ADOS-G consensus meetings with other researchers and clinicians qualified with the tool, but not involved in the present study.

Results

Results at screening stage The response rate from the 1150 families approached through the local primary schools was low. Of 1150 families, we had 199 replies (17.3 percent) for our screen. However, this return rate is not unexpected in postal survey research studies. This was probably due to the CAST being distributed close to the summer break. Of these 199, 25 indicated that they did not wish to take part further in the study. The remaining 174 were sent the SCQ, and replies were received from 139 (79.9 percent).³

Table 3 shows the number and percentage of children from the sample of 199 who scored at or above each point on the CAST. Table 4 shows the number and percentage of children scoring at or above each point on the SCQ. Table 5 shows the percentage of children scoring at or above cut-off on both the SCQ and the CAST, or on one but not the other, or scoring near cut-off on either, or scoring below cut-off on either.

As can be seen from Table 3, 6.5 percent of children in this random sample scored at or above the cut-off of 15. Since the percentage of children with a possible autism spectrum condition would not be expected to exceed around 0.6 percent (e.g. Baird et al., 2000), this suggests either that

Table 4 Number of children (*N* = 139) scoring at or above each point on the SCQ

Total SCQ score	Number (and %) of children
0	139 (100%)
1	123 (88.5%)
2	110 (79.1%)
3	91 (65.5%)
4	78 (56.1%)
5	62 (44.6%)
6	53 (38.1%)
7	49 (35.3%)
8	44 (31.7%)
9	37 (26.6%)
10	26 (18.7%)
11	21 (15.1%)
12	17 (12.2%)
13	15 (10.8%)
14	12 (8.6%)
15	9 (6.5%)
16	7 (5.0%)
17	6 (4.3%)
18	6 (4.3%)
19	5 (3.6%)
20	4 (2.9%)
21	3 (2.2%)
22	2 (1.4%)
23	2 (1.4%)
24	1 (0.7%)
25	0
26	0
27	0
28	0
29	0
30	0
31	0
32	0
33	0
34	0
35	0
36	0
37	0
38	0
39	0
40	0

Prevalence implications

If we consider these results in relation to the sample size originally contacted of 1150 (being conservative owing to the likelihood of a bias in the 199 responders to the screen), this equates to a prevalence of 70 in 10,000. Whilst this number sounds high, it is in line with recent findings suggesting prevalence rates of around 60 in 10,000 (Baird et al., 2000; Scott et al., in press).

Sensitivity and specificity

This preliminary study did not allow for assessment of all children whose parents responded to the CAST; thus it is not possible to establish precise sensitivity and specificity data. To do so would require knowing the number of children who have AS or related social communication difficulty who did not score above cut-off on the CAST (i.e. the false negatives). However, we can report initial positive predictive value and specificity data on the basis of the results to date, but with the addendum that this assumes that the children reported with AS or related conditions versus those without are correctly classified. Additionally, because the screen includes a section asking for details of existing diagnoses, we know that there were no children with existing AS or autism spectrum that were missed by either the CAST or the SCQ. Thus of the *known* cases of AS or ASD, *none* were misclassified by the screening tools as non-cases.

With a cut-off on our screen of 15, it can be seen that 82 percent of children scoring at or above this point either met criteria for an autism spectrum condition or had a deficit relating to social communication (e.g. language delay or social anxiety disorder). We did not conduct further diagnostic assessments of these children at this point, so we do not know what form the language problems took, for example. The positive predictive value (PPV) of the CAST for AS and related social communication conditions was 0.82, with a specificity of 0.99. This compares with a PPV for the SCQ of 0.75, and a specificity of 0.99, for these conditions.

Looking at only AS and autism spectrum criteria, the CAST correctly identified 87.5 percent of cases. However, 36.4 percent of those scoring above cut-off did not meet criteria for AS or autism spectrum (even though 50 percent of those did meet other social communication difficulty criteria). The PPV for AS and autism spectrum for the CAST was 0.64, with a specificity of 0.98. In comparison, the SCQ correctly identified 62.5 percent of AS or autism spectrum cases, with 37.5 percent of those scoring above cut-off failing to meet criteria (of which 67 percent had other social communication difficulties). The PPV for AS and autism spectrum for the SCQ was 0.63, with a specificity of 0.98.

If the cut-off for the CAST were to be raised to 17 for identification of

possible AS or autism spectrum cases, the specificity increases to 0.99, with a PPV of 0.86. Sensitivity is likely to be worsened, as this cut-off catches only 75 percent of true cases (as opposed to 87.5 percent with a cut-off of 15), but there are fewer false positives, with only 14.3 percent of those scoring above 17 failing to meet criteria.

Discussion

The aim of this study was the preliminary development of a UK screening tool for Asperger syndrome (AS) and related social communication difficulties in primary-school-age children. Other screens that have been developed in the past either have only been assessed with known clinical populations, and thus may not be relevant to screening in a non-clinical sample, or have had limited success at identifying children at the higher-functioning end of the spectrum. In fact, there have been few tools developed specifically to identify Asperger syndrome (Howlin, 2000).

Results of this study suggest that the CAST (Childhood Asperger Syndrome Test) may be effective at screening for AS and related social communication conditions in primary-school-age (4–11 years) children in the general population. Compared with the SCQ, the CAST was better able to detect in this sample those children at risk for AS and related disorders who had not already received clinical diagnoses. Seven of the eight children (87.5 percent) who met criteria for autism spectrum conditions were identified by the CAST, whilst the SCQ identified five (62.5 percent). This suggests that the SCQ may perhaps not be as suitable either for use with a non-clinical population, or for identifying less clear-cut cases of AS or related conditions (i.e. those children who are being missed at this age by services).⁴

However, it was clear that with a cut-off of 15 the CAST picks up 6.5 percent of the overall sample (as did the SCQ). This cut-off may thus be deemed to be too low if one wishes to concentrate only on AS and autism spectrum conditions, although identification of children at risk for a broader range of social and communication difficulties may be appropriate at this point. Finding an appropriate balance between specificity and sensitivity is of utmost importance, and an issue which this preliminary research cannot fully address. Establishing accurate sensitivity and specificity data will require longer-term research. With the average age of diagnosis for AS and the higher-functioning end of the autism spectrum currently being about 11 years of age (Howlin and Moore, 1997), one would need to re-examine the sample over a minimum of 7 years, in order to establish whether those children who were age 4 when first assessed had been diagnosed with AS or a related condition by around 11 years of age, and how many of those were picked up or were missed by the CAST. The

floor effects explain the non-significance of question 30. Future research is needed to establish the effect of removing or rewording such questions on the overall sensitivity and specificity of the CAST, and the ongoing study is looking at this initially using latent trait analysis of each CAST question in relation to identification of AS and related conditions.

In summary, these preliminary results indicate that the CAST may be an effective tool for the early screening of primary-school-age (4–11 years) children at risk for AS and related conditions, in a non-clinical sample. With ongoing development it could be established as a UK screener for the broader autism spectrum to be used within that population of children who are currently mislabelled as 'naughty' or 'disruptive', and whose educational and personal development is being compromised owing to lack of or delay in established diagnosis.

Appendix I: the Childhood Asperger Syndrome Test (CAST)

Child's name:..... Age:..... Sex: male/female
 Birth order:..... Twin or single birth:.....
 Parent/guardian:.....
 Parent(s) occupation:.....
 Age parent(s) left full-time education:.....
 Address:.....

 Tel. no.:..... School:.....

Please read the following questions carefully, and circle the appropriate answer. All responses are confidential.

- | | | | |
|----|--|-----|----|
| 1 | Does s/he join in playing games with other children easily? | Yes | No |
| 2 | Does s/he come up to you spontaneously for a chat? | Yes | No |
| 3 | Was s/he speaking by 2 years old? | Yes | No |
| 4 | Does s/he enjoy sports? | Yes | No |
| 5 | Is it important to him/her to fit in with the peer group? | Yes | No |
| 6 | Does s/he appear to notice unusual details that others miss? | Yes | No |
| 7 | Does s/he tend to take things literally? | Yes | No |
| 8 | When s/he was 3 years old, did s/he spend a lot of time pretending (e.g. play-acting being a superhero, or holding teddy's tea parties)? | Yes | No |
| 9 | Does s/he like to do things over and over again, in the same way all the time? | Yes | No |
| 10 | Does s/he find it easy to interact with other children? | Yes | No |
| 11 | Can s/he keep a two-way conversation going? | Yes | No |
| 12 | Can s/he read appropriately for his/her age? | Yes | No |
| 13 | Does s/he mostly have the same interests as his/her peers? | Yes | No |

Appendix 2: parental cover letter

QUESTIONNAIRE INFORMATION SHEET

Dear Parent

We would like to invite you to take part in a research study being conducted by the University of Cambridge exploring how social and communication skills develop in primary-school-age children.

Children develop such skills in very different ways. Some children are very outgoing and sociable, others more quiet and reserved. A few children may be very shy. A very small number of children may have difficulties in their social development. This can be for a variety of reasons. For example social anxiety problems may underlie the difficulty in mixing. Very occasionally the difficulties may be due to Asperger syndrome or an autism spectrum condition (conditions where children have significant problems understanding social and emotional situations).

We are interested in exploring the full range of development of social and communication skills in children from the whole population. This research will then help us to better understand when children do have difficulties.

We are inviting parents of children aged 4 to 11 from Cambridge, Huntingdon and Fenland areas to help us. This involves filling in the questionnaire provided, which takes about 10 minutes. A Freepost envelope is provided so you can post this directly to us. In order for us to get a truly representative picture of the range of social communication styles, it is important we receive replies from everyone willing to participate.

After we have received all the questionnaires, we would like to invite around 10 percent of people to take part in more detailed face-to-face assessments to see how accurately our questionnaire works in characterizing social communicative style. Participation with the questionnaire survey does not commit you to helping with these more detailed assessments. We will write to families again asking if they would be willing to help with the second part of the survey, and if you wish you could decline to participate further at that or any other stage. If you have any concerns about your child or if we identify a possible developmental problem, we will be happy to discuss these with you and if you are in agreement arrange for a clinical specialist to see you and advise further.

If you are happy to take part in this study, we would be grateful if you would complete and return the questionnaire(s) in the Freepost envelope provided. All information you supply will be confidential to the research team. We would be interested to receive your questionnaire even if you do not wish to take part in later stages of the project.

You are of course free to withdraw from the study at any stage without providing an explanation, should you wish to do so. Neither participation nor non-participation in this research will affect any treatments or services your child may be receiving or be entitled to receive.

Should you wish to discuss this research further, or have any questions, the Project Coordinator, Dr Fiona Scott, can be contacted on 01223 746113 (fax: 01223 746122; e-mail: fjs25@cam.ac.uk).

Acknowledgements

This study was funded by the Inge Wakehurst Trust. In addition, FJS was supported by the Isaac Newton Trust, and the NHS R&D. SBC was supported by the MRC (UK), the Three Guineas Trust, and the Shirley Foundation.

Notes

- 1 Whilst children with AS are 'higher-functioning' in terms of cognitive and language development, there remain severe difficulties in social interaction,

- HOWLIN, P. & MOORE, A. (1997) 'Diagnosis in Autism: A Survey of over 1200 Patients in the UK', *Autism* 1: 135–62.
- KAZAK-BERUMENT, S., RUTTER, M., LORD, C., PICKLES, A. & BAILEY, A. (1999) 'Autism Screening Questionnaire: Diagnostic Validity', *British Journal of Psychiatry* 175: 444–51.
- KLIN, A. & VOLKMAR, F. (1997) 'Asperger's Syndrome', in D. COHEN & F. VOLKMAR (eds) *Handbook of Autism and Pervasive Developmental Disorders*, 2nd edn. New York: Wiley.
- KUGLER, B. (1998) 'The Differentiation between Autism and Asperger Syndrome', *Autism* 2: 11–32.
- LAW, J., BOYLE, J., HARRIS, F., HARKNESS, A. & NYE, C. (2000) 'The Feasibility of Universal Screening for Primary Speech and Language Delay: Findings from a Systematic Review of the Literature', *Developmental Medicine and Child Neurology* 42: 190–200.
- LORD, C., RUTTER, M. & LECOUEUR, A. (1994) 'Autism Diagnostic Interview—Revised: A Revised Version of a Diagnostic Interview for Carers of Individuals with Possible Pervasive Developmental Disorders', *Journal of Autism and Developmental Disorders* 24: 659–85.
- LORD, C., RUTTER, M., DILAVORE, P. & RISI, S. (1999) *Autism Diagnostic Observation Schedule—Generic (ADOS—G)*. Los Angeles, CA: Western Psychological Services.
- NATIONAL SCREENING COMMITTEE (1998) *First Report of the National Screening Committee*. Department of Health, UK. www.nsc.nhs.uk/pdfs/nsc_firstreport.pdf or www.open.gov.uk/doh/nsc/nsch.htm.
- ROBINSON, R. (1998) 'Effective Screening in Child Health', *British Medical Journal* 316: 1–2.
- SCOTT, F.J., BARON-COHEN, S., BOLTON, P. & BRAYNE, C. (in press) 'Brief Report: Prevalence of Autism Spectrum Conditions in Children Aged 5–11 years in Cambridgeshire, UK'. *Autism: International Journal of Research and Practice*.
- TONGE, B.J., BRERETON, A.V., GRAY, K.M. & EINFELD, S.L. (1999) 'Behavioural and Emotional Disturbance in High-Functioning Autism and Asperger Syndrome', *Autism* 3: 117–30.
- VOLKMAR, F.R. (1998) 'Categorical Approaches to the Diagnosis of Autism: An Overview of DSM-IV and ICD-10', *Autism* 2: 45–59.
- WING, L. & GOULD, J. (1979) 'Severe Impairments of Social Interactions and Associated Abnormalities in Children: Epidemiology and Classification', *Journal of Autism and Developmental Disorders* 9: 11–29.
- WORLD HEALTH ORGANIZATION (1993) *The ICD-10 Classification of Mental and Behavioural Disorders: Diagnostic Criteria for Research*. Geneva: WHO.