Heritability of Autistic Traits in the General Population

Rosa A. Hoekstra, MSc; Meike Bartels, PhD; Catharina J. H. Verweij, MSc; Dorret I. Boomsma, PhD

Objective: To explore genetic and environmental influences on individual differences in autistic traits in early adulthood and to test if there is assortative mating (non-random partner choice) for autistic traits in the general population.

Design: Twin family study using structural equation modeling.

Setting: Population-based twin family sample from the Netherlands.

Participants: Twins aged 18 years (n=370) and their siblings (n=94); parents of twins (128 couples).

Main Outcome Measure: Self-reported Autism-Spectrum Quotient (AQ) scores, a quantitative measure of autistic traits.

Results: Autistic traits were continuously distributed in the population. Twins and siblings did not significantly differ in AQ scores; men obtained significantly higher AQ scores than women (in twin-sibling sample, \( P = .02 \); twin-parent sample, \( P = .02 \)). Individual differences in endorsement on autistic traits show substantial heritability (57%). No significant shared environmental influences were detected. The genes affecting autistic traits appear to be the same across the sexes. The correlation in AQ score between spouses was low and not significant (Pearson \( r = .05; P = .59 \)).

Conclusions: Previous general population twin studies reported high heritability for autistic traits in childhood and early adolescence. This study extends these findings to late adolescence and yields no evidence for sex-specific genetic influences on autistic traits in later stages of development. As autistic traits show substantial variation in the general population, future genetic studies may be facilitated by measuring autistic traits on a continuous scale like the AQ. No evidence for assortative mating for autistic traits was found, suggesting that, in the general population, there is no passive or active partner selection for autistic traits.

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Author Affiliations:
Department of Biological Psychology, VU University Amsterdam, Amsterdam, the Netherlands.
ences were found. In more than 3000 7-year-old male, female, and opposite-sex twin pairs, both social and non-social autistic behaviors were found to be highly heritable. A study in the same sample 1 year later reported high heritability of autistic traits and no shared environmental influences. The latter study also used a categorical approach (extreme vs typical endorsement on autistic traits) and found results similar to the dimensional approach, yielding no indication that etiology is different at the extreme end of the spectrum. These studies were all conducted in children and young adolescents. No studies of the heritability of autistic traits at later ages have been reported yet, and none have included siblings of twins. Furthermore, the studies mentioned were all based on parent or teacher ratings. Previous studies of behavioral problems have shown that external and self-rated reports may yield other results, as different raters can provide different perspectives on behavior. 

Our study aims to examine genetic and environmental influences on self-reported autistic traits in a sample of 18-year-old twins and their siblings using the Autism-Spectrum Quotient (AQ), a well-validated instrument used to quantify autistic traits (R.A.H., unpublished data, 2006).

Additionally, assortative mating (nonrandom partner choice) for autistic traits will be examined. Assortative mating could influence the frequency of the genotypes related to autistic traits and bias correlations in first-degree relatives and, consequently, heritability estimates. One previous study of assortative mating for autistic traits reported a spouse correlation of $r = 0.38$. However, in this study, spouses rated each other and not themselves on the endorsement of autistic traits. Shared beliefs or perceptions about the couple's relationship may have inflated the results. Various studies have explored partner resemblance for personality traits and reported modest to moderate similarity for introversion and modest similarity in preference for consistency and routine. We examined assortative mating for autistic traits in a general population sample using self-reported AQ scores.

**METHODS**

**PARTICIPANTS**

The twin families participating in the heritability study were recruited via the Netherlands Twin Register kept by the Department of Biological Psychology at the VU University in Amsterdam. The current study sample comprised 194 families and is part of an ongoing longitudinal project examining development of cognition and behavioral problems. Participation rate for this data collection was 54%. Participating families did not significantly differ from nonparticipating families in socioeconomic status (Mann-Whitney test, $U = 10382.00; P = .23$; effect size, $r = 0.07$), but parental education level was slightly higher in participating families (education of mother, $U = 9538.00, P = .05, r = 0.12$; father, $U = 7773.00, P = .01, r = 0.16$). No information about ASD diagnoses was available. Mean age of the twins was 18.18 years (SD, 0.22; range, 17.61-18.99); mean sibling age was 18.77 years (SD, 4.71; range, 10.52-35.39). Most twin families ($n = 184$) completed the AQ in the university laboratory as part of an extensive test protocol. The other families ($n = 10$) filled out the questionnaire at home. The sample consisted of 36 monozygotic male twin pairs, 35 dizygotic male twin pairs, 45 pairs of monozygotic female twins, 39 pairs of dizygotic female twins, and 39 dizygotic twin pairs of opposite sex. Zygosity of the same-sex twin pairs ($n = 135$) was determined by DNA analyses ($n = 101$), blood group polymorphisms ($n = 45$), or discriminant analyses of longitudinally collected questionnaire items ($n = 9$). This method has been proven to be of sufficient reliability. This study was approved by the Central Committee on Research Involving Human Subjects and the institutional review board of the VU University Amsterdam. Written informed consent was obtained from all participating subjects.

To study assortative mating for autistic traits, parents of twins (unrelated to the twin families mentioned) were asked to fill out the AQ during an informational day for parents of multiples. They either completed the AQ during the day or returned it to our research group by mail. The response rate was 62%; no information was available about nonresponders. The mean age of the participants was 35.68 years (SD, 6.33). Only data of male-female couples were included; complete partner data on the AQ were available for 128 pairs. All couples were either living together or married.

**DUTCH AUTISM-SPECTRUM QUOTIENT**

The AQ consists of 50 items assessing personal preferences and habits. Subjects rate to what extent they agree or disagree with the statements on a 4-point Likert scale, with the following answer categories: 1 representing definitely agree; 2, slightly agree; 3, slightly disagree; and 4, definitely disagree. Total AQ scores were calculated as the sum of the Likert scale scores. For items in which an agree response was characteristic for autism, the scoring was reversed (definitely agree scored 4 points; slightly agree, 3 points; slightly disagree, 2 points; and definitely disagree, 1 point). The minimum AQ score (50) indicates no autistic traits; the maximum score (200) suggests full endorsement on all autistic items.

The original English version of the AQ was translated into Dutch using a backward translation procedure. After comparing the outcome of the retranslated version with the original text, a final version was established. The Dutch translation of the AQ has good internal consistency (Cronbach's $\alpha = 0.79$) and test-retest reliability (Pearson's $r = 0.78$ in a group of 75 subjects with a 1- to 6-month time interval) (R.A.H., unpublished data, 2006).

If more than 5 items were left blank, the AQ was considered incomplete and the data were discarded in subsequent analyses ($n = 7$ in the twin-sibling sample). Complete AQs were available for 370 twins and 94 siblings. If 5 or fewer answers were missing, the AQ score was corrected for the number of missing items by making the following calculation: total AQ score + (mean item score × number of missing items). Twenty-one individuals were missing 1 answer, and 3 individuals were missing 2 answers.

**DATA ANALYSES**

Descriptive statistics were calculated using SPSS 13.0 for Windows (SPSS Inc, Chicago, Ill). Twin-sibling differences in AQ score and effects of birth order, zygosity, age, and sex were examined using a saturated model in the structural equation modeling program Mx. Twin and twin-sibling correlations for AQ scores were estimated for each zygosity group. Sex differences in mean AQ score in the twin-parent sample were
examining the deterioration in model fit after constraining
the magnitude of additive genetic influences, shared envi-
ronmental influences, and nonshared environmental influ-
ences to be equal across the sexes. The significance of the
contribution of additive genetic influences and shared envi-
ronmental influences was tested by assessing the deteriora-
tion in model fit after each component was dropped from
the full model.

**RESULTS**

**Table 1** presents the descriptive statistics for AQ scores
in the twins and their siblings, and in spouses drawn from
the sample of parents of twins. Autism-Spectrum Quo-
tient scores were continuously distributed (Figure 2).
No differences in mean AQ scores between twins and sib-
lings were found (102.1 vs 102.9, respectively; \( \chi^2 = 1.18; \)
\( P = .28 \)). Moreover, no effects of birth order (\( \chi^2 = 1.66; \)
\( P = .44 \)), zygosity (\( \chi^2 = 1.44; P = .23 \)), or age (\( \chi^2 = 0.59; \)
\( P = .44 \)) could be detected. A significant sex effect on
the mean was found; mean AQ scores were signifi-
cantly higher in men than in women (104.0 vs 100.8,
respectively; \( \chi^2 = 12.97; P < .001 \); effect size, \( d = 0.30 \)).
Similarly, in the twin-parent sample, men obtained
significantly higher AQ scores than women (mean 106.0
vs 102.8, respectively; \( F_{1,254} = 5.32; P = .02 \); \( d = 0.28 \)). No evidence for assortative mating for autis-
tic traits was found. The partner correlation for AQ
scores was \( r = 0.05 \) (\( P = .59 \)).

Twin and twin-sibling correlations are presented in
**Table 2**. Inspection of the monozygotic, dizygotic,
and twin-sibling correlations gives a first impression of what
factors influence individual differences in AQ scores. Al-
though the confidence intervals overlap, the estimates for
monozygotic correlations are higher than dizygotic and
twin-sibling correlations, indicating that genetic factors
may play a role. As twin correlations in opposite-sex twins
are not attenuated, compared with the correlations in
same-sex dizygotic twins, there is no indication for sex-
specific genes influencing variance in AQ scores. The
monozygotic correlations are not twice as high as the di-
yzgotic correlations and twin-sibling correlations, sug-
gest that shared environmental factors could also be
of importance.

Model-fitting statistics for the full model, including
both additive genetic, shared environmental, and non-
shared environmental influences (referred to as the
ACE model), and various submodels are presented in
**Table 3**. Constraining the parameters that represent
the effect of additive genetic, shared environmental, and
nonshared environmental influences as equal across the
sexes did not significantly worsen the fit (\( \chi^2 = 2.88; \)
\( P = .41 \)), confirming that the relative effects of these
components were the same in men and women. Dropping
the shared environmental component from the model
did not result in a worse model fit (\( \chi^2 = 0.78; \)
\( P = .37 \)). The genetic effects, however, were of signifi-
cant importance (\( \chi^2 = 4.33; P = .04 \)). In the best-fitting
parsimonious model, individual differences in autistic
traits were explained by additive genetic influences (ac-
counting for 57% of the variance) and nonshared envi-
ronmental influences (accounting for 43% of the variance).

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**Table 1. Sample Size and AQ Score in Twins, Their Siblings, and in Spouses**

<table>
<thead>
<tr>
<th>Subject</th>
<th>Sample Size</th>
<th>AQ Score, Mean ± SD</th>
</tr>
</thead>
<tbody>
<tr>
<td>Twin sample</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Twin</td>
<td>370</td>
<td>102.1 ± 10.5</td>
</tr>
<tr>
<td>Sibling</td>
<td>94</td>
<td>102.9 ± 11.0</td>
</tr>
<tr>
<td>M</td>
<td>217</td>
<td>104.0 ± 10.5</td>
</tr>
<tr>
<td>F</td>
<td>247</td>
<td>100.8 ± 10.5</td>
</tr>
<tr>
<td>All</td>
<td>464</td>
<td>102.3 ± 10.6</td>
</tr>
<tr>
<td>Spouse sample*</td>
<td></td>
<td></td>
</tr>
<tr>
<td>M</td>
<td>128</td>
<td>106.0 ± 10.9</td>
</tr>
<tr>
<td>F</td>
<td>128</td>
<td>102.8 ± 11.5</td>
</tr>
<tr>
<td>All spouses</td>
<td>256</td>
<td>104.4 ± 11.3</td>
</tr>
</tbody>
</table>

*Parents of twins not in the twin sample.

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**Figure 1.** Univariate path diagram representing the contribution of additive genetic (A), shared environmental (C), and nonshared environmental (E) influences to the trait under investigation (Autism-Spectrum Quotient scores). The factor loadings of these influences are represented by a, c, and e. The correlation of the additive genetic factors is 1.0 in monozygotic twins and, on average, 0.5 in dizygotic twins and between twins and siblings. The correlation of the shared environmental effects is 1.0 between twins and between twins and siblings. E represents effects unique to a family member and are thus uncorrelated.
The present study shows that autistic traits, as measured by the AQ in the general population, are continuously distributed; show a significant sex difference in mean scores, with men scoring higher than women; and are unrelated to age or to being born a twin or singleton. Moreover, individual differences in autistic traits show substantial heritability, and are influenced by the same additive genetic factors in men and women. No evidence for assortative mating for autistic traits was found.

The finding of a significant sex difference in mean AQ scores in both the twin-sibling and the twin-parent sample is in concordance with findings using other measures of autistic traits, such as the Social Responsiveness Scale and the Childhood Asperger Syndrome Test, suggesting cross-cultural similarities. All studies report higher endorsement on autistic traits in men than in women, which is in line with the observation that ASDs are more common in men than in women.

We did not find a difference in mean AQ scores between twins and singletons. Some studies have suggested the process of twinning as a risk factor for the development of autism. Large population-based studies did not support these findings. Our results indicate that in the general population, endorsement of autistic traits is unrelated to being born a twin or singleton.

In 18-year-old twins and their siblings, variance in autistic traits is largely explained by additive genetic effects (57%). Shared environmental effects were not of significant importance; nonshared environmental effects accounted for 43% of the variance. Twin correlations in same-sex dizygotic twins were of similar magnitude as the correlation in opposite-sex twins, yielding no evidence for sex-specific genes for autistic traits. Comparable dizygotic same-sex and opposite-sex twin correlations were found in twin studies in childhood and early childhood.

**Table 2. Twin and Twin-Sibling Correlations**

<table>
<thead>
<tr>
<th>Subject Pair</th>
<th>Complete Pair, No.</th>
<th>Incomplete Pair, No.</th>
<th>Correlation (95% Confidence Interval)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Monozygotic male twin pair</td>
<td>33</td>
<td>3</td>
<td>.59 (.32 to .74)</td>
</tr>
<tr>
<td>Dizygotic male twin pair</td>
<td>43</td>
<td>2</td>
<td>.51 (.23 to .68)</td>
</tr>
<tr>
<td>Monozygotic female twin pair</td>
<td>45</td>
<td>3</td>
<td>.43 (.07 to .65)</td>
</tr>
<tr>
<td>Dizygotic female twin pair</td>
<td>37</td>
<td>3</td>
<td>.35 (.11 to .55)</td>
</tr>
<tr>
<td>Dizygotic twin pair of opposite sex</td>
<td>37</td>
<td>3</td>
<td>.35 (.11 to .55)</td>
</tr>
<tr>
<td>All monozygotic twin pairs</td>
<td>76</td>
<td>5</td>
<td>.55 (.37 to .67)</td>
</tr>
<tr>
<td>All dizygotic twin pairs</td>
<td>110</td>
<td>11</td>
<td>.57 (.30 to .65)</td>
</tr>
<tr>
<td>Male twin-sibling pairs</td>
<td>102</td>
<td>11</td>
<td>.57 (.30 to .65)</td>
</tr>
<tr>
<td>Female twin-sibling pairs</td>
<td>58</td>
<td>2</td>
<td>.47 (.24 to .62)</td>
</tr>
<tr>
<td>Twin-sibling pair of opposite sex</td>
<td>77</td>
<td>5</td>
<td>.33 (.12 to .50)</td>
</tr>
<tr>
<td>All twin-sibling pairs</td>
<td>180</td>
<td>8</td>
<td>.28 (.12 to .41)</td>
</tr>
<tr>
<td>All first-degree relative pairs</td>
<td>282</td>
<td>19</td>
<td>.21 (.19 to .43)</td>
</tr>
</tbody>
</table>

**Figure 2.** Distribution of Autism-Spectrum Quotient (AQ) scores in twins and their siblings.
adolescence. Our results indicate that even in more advanced stages of development, the set of genes influencing autistic traits is the same across the sexes.

The correlations in first-degree relatives in this study (r = 0.32) are similar to the dizygotic twin correlations reported in other studies assessing autistic traits on a continuum but are considerably larger than dizygotic twin concordance rates and sibling prevalence rates for clinical diagnosis of autism (0%-5%). It is found that dizygotic twin concordance rates and sibling prevalence increase when diagnostic criteria are relaxed and include a broader phenotype of autistic traits. Genetic studies, as the strong disparity between monozygotic and dizygotic twin concordance rates for diagnosed autism have led to the hypothesis that gene-gene interactions play an important role in the risk for autism.

Our results suggest that shared environmental effects are of major importance in explaining the variance of autistic traits, but the power to detect such effects was limited with our sample size. In previous studies including larger sample sizes, 1 study found no shared environmental effects, and 1 reported a small but significant contribution of shared environment in girls but not in boys, and 1 reported moderate shared environmental influences in both sexes. The studies reporting significant shared environmental effects were both based on parental reports. As the parent rates the behavior of both members of the twin pair, rater bias may have inflated the shared environmental effects in these studies.

Our study relied on self-report measurement of autistic traits. As subjects with autism may underestimate the extreme. However, the real test for this should come from genetic studies.

No evidence for assortative mating for autistic traits was found. The correlation of AQ scores between partners was close to zero. This is in contrast to a previous study that reported a spouse correlation of r = 0.38. How- ever, as their assessment of autistic traits was based on spouse report, shared perceptions about the relationship may have inflated the correlation. All participants in our assortative mating study were recruited on an informational day for parents of multiples. Individuals who dislike being confronted with large crowds may be unlikely to attend this event. Our sample may therefore not be completely representative of the general population. The results suggest that in the general population, people do not actively or passively select their partner for autistic characteristics. One theory has proposed assortative mating for extreme autistic traits as a risk factor for having a child with an ASD. Our sample included insufficient numbers of extreme AQ scorers to test this hypothesis.

From these data, we conclude that variance in autistic traits, as measured with the AQ, show substantial heritability. There is no indication that the heritability estimate reported here is confounded by assortative mating. This study shows that the strong heritability is not limited to the clinical autism spectrum, but also accounts for variance in autistic traits in the general population. Singletons do not differ from twins in endorsement of autistic traits. Genetic studies may be facilitated by measuring autistic traits on a continuous scale like the AQ. Such studies can elucidate whether the genes associated with the clinical spectrum are also associated with normal variation in autistic traits.

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Correspondence: Rosa A. Hoekstra, MSc, Department of Biological Psychology, Vrije University, Van der Boechorststraat 1, Amsterdam 1081 BT, the Netherlands (ra.hoekstra@psy.vu.nl).

Author Contributions: Study concept and design: Hoekstra, Bartels, and Boomsma. Acquisition of data: Hoekstra and Verweij. Analysis and interpretation of data: Hoekstra, Bartels, Verweij, and Boomsma. Drafting of the manuscript: Hoekstra, Bartels, Verweij, and Boomsma. Critical revision of the manuscript for important intellectual content: Hoekstra, Bartels, and Boomsma. Statistical analysis: Hoekstra, Bartels, Verweij, and Boomsma. Obtained funding: Boomsma.

Table 3. Model Fit Statistics and Parameter Estimates for the Best-Fitting Model

<table>
<thead>
<tr>
<th>Model</th>
<th>−2LL</th>
<th>df</th>
<th>Compared With Model</th>
<th>( \chi^2 )</th>
<th>( P ) Value</th>
<th>A (95% CI)</th>
<th>E (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>ACE, sex differences</td>
<td>3444.32</td>
<td>454</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
</tr>
<tr>
<td>ACE, no sex differences</td>
<td>3447.19</td>
<td>457</td>
<td>1</td>
<td>2.88</td>
<td>.41</td>
<td>.57 (.43-.68)</td>
<td>.43 (.32-.57)</td>
</tr>
<tr>
<td>AE, no sex differences</td>
<td>3447.98</td>
<td>458</td>
<td>2</td>
<td>0.78</td>
<td>.38</td>
<td>.57 (.43-.68)</td>
<td>.43 (.32-.57)</td>
</tr>
<tr>
<td>CE, no sex differences</td>
<td>3451.54</td>
<td>458</td>
<td>2</td>
<td>4.35</td>
<td>.04</td>
<td>.57 (.43-.68)</td>
<td>.43 (.32-.57)</td>
</tr>
<tr>
<td>E, no sex differences</td>
<td>3496.69</td>
<td>459</td>
<td>3</td>
<td>48.71</td>
<td>&lt;.001</td>
<td>.57 (.43-.68)</td>
<td>.43 (.32-.57)</td>
</tr>
</tbody>
</table>

Abbreviations: A, relative contribution of genetic influences; C, relative contribution of shared environmental influences; CI, confidence interval; E, relative contribution of nonshared environmental influences; NA, not applicable; −2LL, minus twice the log likelihood.
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REFERENCES


Simply imagine that it’s not your child, but someone else’s. Everybody knows how to bring up other people’s children.
—Russian proverb