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Suggestive Linkage on Chromosome 2, 8, and 17 for Lifetime Major Depression

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It is well established that major depressive disorder (MDD) is partly heritable. We present a genome-wide linkage study aiming to find regions on the genome that influence the vulnerability for MDD. Our sample consists of 110 Australian and 23 Dutch pedigrees with two or more siblings affected with MDD (total N = 278). Linkage analysis was carried out in MERLIN. Three regions showed suggestive linkage signals. The highest LOD-score of 2.1 was found on chromosome 17 at 52.6 cM along with LOD scores of 1.9 and 1.7 on chromosome 8 at 2.7 cM and chromosome 2 at 90.6 cM, respectively. The result on chromosome 8 seems most promising as two previous studies also found linkage in this region, once suggestive and once significant. The linkage peak on chromosome 17 harbors the serotonin transporter gene. In the Australian and Dutch sample, the serotonin transporter length polymorphism did not show evidence for association, thus other genes in this region or other polymorphisms in the serotonin transporter gene might be associated with MDD. Further replication is needed to establish the relevance of our linkage finding on chromosome 2.

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genes influencing the vulnerability for MDD. As part of these efforts, seven genome-wide linkage analyses have been performed, aiming to locate genes for MDD on the genome [Nurnberger et al., 2001; Zubenko et al., 2003; Holmans et al., 2004, 2007; Camp et al., 2005; McGuffin et al., 2005]. Other studies have focused on quantitative traits associated with a diagnosis of MDD, such as neuroticism [Cloninger et al., 1998; Fullerton et al., 2003; Nash et al., 2004; Neale et al., 2005; Kuo et al., 2007; Wray et al., 2008]. Table I summarizes the most promising results of these studies, excluding the study of Holmans et al. [2004] as this is based on the same sample as used by Holmans et al. [2007]. Several regions have shown a linkage signal with an LOD-score >3 in at least one study. The following five regions have reached an LOD-score >3 in one study and an LOD-score >1.5 in a second study: chromosome 1 between 126 and 137 cM [Fullerton et al., 2003; Neale et al., 2005], chromosome 8 between 8 and 38 cM [Cloninger et al., 1998; Fullerton et al., 2003], chromosome 11 between 2 and 35 cM [Zubenko et al., 2003; Camp et al., 2005], chromosome 11 between 85 and 99 cM [Fullerton et al., 2003; Zubenko et al., 2003] and chromosome 12 between 105 and 124 cM [Fullerton et al., 2003; McGuffin et al., 2005].

Five of these studies focused on phenotypes considered to be most influenced by genetic factors, that is recurrent MDD [McGuffin et al., 2005] or recurrent, early-onset MDD [Zubenko et al., 2003; Holmans et al., 2004, 2007; Camp et al., 2005]. This seemed a useful strategy to increase the chance to find genes involved in MDD. However, comparing their results with the linkage analysis of MDD [Nurnberger et al., 2001] or with the additional linkage analysis of major mood disorders [Zubenko et al., 2003] shows that this approach does not seem to result in higher LOD scores than using a diagnosis of lifetime MDD. Although Zubenko et al. [2003] have found the highest LOD score for recurrent, early-onset and recurrent MDD, additional linkage peaks were identified for major mood disorders.

We present a new linkage study of MDD performed on affected sibling pairs selected from the Australian and Dutch Twin Registers. Although these samples are geographically disparate, we have shown previously that the genetic distance between these Dutch and Australian subjects (Fst = 0.30%) is comparable to that between most Northern European populations [Sullivan et al., 2006]. Also, the genetic architecture of MDD is similar in the two countries with a heritability estimate for MDD of 36% and without evidence for sex differences in genetic architecture, allowing us to pool data from men and women in the linkage scan [Middeldorp et al., 2005].

### MATERIALS AND METHODS

#### Subjects

Data were collected in the Australian and Netherlands Twin Registers as part of a project aiming to find the genes underlying the susceptibility to anxiety and MDD. Detailed descriptions of the data collection have been published previously [Boomsma et al.,

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**TABLE I. Most Promising Linkage Results in the Order of the Chromosomes for Neuroticism, Harm Avoidance, MDD or the Subtypes Recurrent MDD (R-MDD) and Recurrent Early-Onset MDD (RE-MDD)**

<table>
<thead>
<tr>
<th>Location (chromosome, cM)</th>
<th>LOD</th>
<th>N subjects*/families</th>
<th>Phenotype</th>
</tr>
</thead>
<tbody>
<tr>
<td>1, 42, and 90 cM [Camp et al., 2005; Nash et al., 2004]</td>
<td>1.7</td>
<td>426/90</td>
<td>MDD-RE</td>
</tr>
<tr>
<td>1, 126–137 cM [Fullerton et al., 2003; Neale et al., 2005]</td>
<td>1.6</td>
<td>711/283</td>
<td>Neuroticism</td>
</tr>
<tr>
<td>2, 237–248 cM [Nurnberger et al., 2001; Zubenko et al., 2003]</td>
<td>4.0 b</td>
<td>1,122/561</td>
<td>Neuroticism</td>
</tr>
<tr>
<td>4, 176 cM Fullerton et al. [2003]</td>
<td>2.5 b</td>
<td>293/129</td>
<td>Neuroticism</td>
</tr>
<tr>
<td>7, 8–38 cM [Cloninger et al., 1998; Fullerton et al., 2003]</td>
<td>2.2</td>
<td>224 possible pairs</td>
<td>MDD comorbid with alcoholism</td>
</tr>
<tr>
<td>10, 5–9 cM [Camp et al., 2005; Wray et al., 2008]</td>
<td>2.5 b</td>
<td>?/81</td>
<td>MDD</td>
</tr>
<tr>
<td>10, 76 cM Zubenko et al. [2003]</td>
<td>3.8 b</td>
<td>1,122/561</td>
<td>Neuroticism</td>
</tr>
<tr>
<td>11, 2–35 cM [Camp et al., 2005; Zubenko et al., 2003]</td>
<td>3.9 b</td>
<td>1,122/561</td>
<td>Neuroticism</td>
</tr>
<tr>
<td>11, 85–99 cM [Fullerton et al., 2003; Zubenko et al., 2003]</td>
<td>3.2</td>
<td>987/105</td>
<td>Harm avoidance</td>
</tr>
<tr>
<td>12, 105–124 cM [Fullerton et al., 2003; McGuffin et al., 2005]</td>
<td>2.9 b</td>
<td>1,122/561</td>
<td>Neuroticism</td>
</tr>
<tr>
<td>13, 64 cM Fullerton et al. [2003]</td>
<td>1.6</td>
<td>426/90</td>
<td>RE-MDD</td>
</tr>
<tr>
<td>18, 73 cM Camp et al. [2005]</td>
<td>2.0</td>
<td>2,030/564</td>
<td>Neuroticism</td>
</tr>
<tr>
<td>18, 109–117 cM [Cloninger et al., 1998; Wray et al., 2008]</td>
<td>3.0</td>
<td>?/81</td>
<td>MDD</td>
</tr>
</tbody>
</table>

*For the MDD studies the number of affected individuals is given.

This is the – log P, not the LOD score.
Genotyping

The genotypic data available for the Australian study resulted from submission of DNA samples to one or more of six genotyping centers, namely Gemini P/L (G), Sequana Therapeutics Inc. (S), Leiden University Medical Center (L), the Center for Mammalian Genetics at the Marshfield Clinic Research Foundation (M), the Australian genotyping research facility (AGRF) (A), and the Finnish Genome Center, University of Helsinki (H). A description of the G, L, M, S genotyping and the subsequent merging and cleaning of the marker data sets is described in detail elsewhere [Cornes et al., 2005; Nyholt et al., 2005]. Since then, additional M, A, and H genotypes have been merged using the same protocol. Family members were submitted to the same genotyping facility. Data cleaning based on Mendelian errors, unlikely genotypes and consistency of pedigree and marker relationships was undertaken as described by Cornes et al. [2005]. In brief, pedigree structures for each scan were examined using graphic representation of relationships (GRR) [Abecasis et al., 2001] and RELPAIR [Epstein et al., 2000; Duren et al., 2003], to identify inconsistencies between the genotypic data and pedigree relationships. Once any discrepancies were resolved, data for the scans were merged and then checked again for pedigree errors by GRR and for Mendelian inconsistencies by SIB-PAIR [Duffy, 2002]. Duplicate markers in the combined genome scan data were included separately on the genetic map, separated by a very small distance (0.001 cM). The consistency of genotype information between these duplicate markers was checked via cross-tabulations of allele calls between different scans. Markers with genotypic data inconsistent between different genome scans were removed from further analysis. The Mendelian error rate, counted as the number of genotypes removed divided by total number of genotypes using all markers genotyped in more than one lab was 0.57%. Unlike genotypes were identified and wiped using MERLIN [Abecasis et al., 2002]. Error rates were similar for the different labs.

For the Dutch samples, DNA was extracted from either whole blood or buccal swabs following standard protocols [Miller et al., 1988; Meulenberg et al., 1995]. Samples were genotyped by M and L. The genotype data from these screens were combined. Allele calling and binning were equalized between markers that were present in multiple scans, using ~30 control samples. In case there were inconsistencies, the data were set to unknown for tested markers (binning and allele calling inconsistencies) and persons (genotyping errors). Sex and prior measured zygosity were checked with the marker data. Pedigree relations in the entire samples were checked with the GRR program [Abecasis et al., 2001]. Errors of Mendelian inheritance were detected with Pedstats [Wigginton and Abecasis, 2005]. Markers and samples were removed if their total error rate was more than 1%; in all other cases the specific erroneous genotypes were set as unknown. The mendelian error rate was 0.58%. Unlikely recombinants were detected using Merlin and erroneous genotypes were removed with Pedwise [Abecasis et al., 2002]. Error rates were similar for the different labs.

Map positions of all genotyped markers were estimated in Kosambi cM by locally weighted linear regression http://www.qimr.edu.au/davidd from the NCBI Build 35.1 physical map positions and published Decode and Marshfield genetic map positions [Duffy, 2006]. Joint analysis of common markers genotyped in the Australian and Dutch samples estimated ΦB between the populations to be only 0.30% implying that these samples can be combined for joint genetic analysis [Sullivan et al., 2006]. Individuals were required to have genotypes on more than 280 markers resulting in an average inter-marker distance of 8.2 cm (AU) and 9.6 cM (NL). The average number of markers genotyped in siblings was 719 in the Australian sample (range 334–1,242) and 400 in the Dutch sample (range 291–761).

Instruments

A telephone interview, during which the computerized version of the composite international diagnostic interview (CIDI) [World Health Organization, 1992] was administered to obtain lifetime DSM-IV diagnoses of mood and anxiety disorders [American Psychiatric Association, 1994]. The CIDI is a fully standardized diagnostic interview. Good reliability and validity have been reported for the CIDI [Andrews and Peters, 1998]. All interviewers were trained by the Dutch and Australian World Health Organization training centers. In order to minimize observer bias, interviewers were unaware of interviewees’ status on the initial selection variables throughout the study. The dependent variable used for
linkage analysis was a lifetime diagnosis of MDD (DSM-IV codes: 296.21, 296.22, 296.23, 296.31, 296.32, and 296.33). Subjects who might have experienced a (hypo)manic episode as measured with the CIDI screening question (Australia) or the total “bipolar disorder” section (The Netherlands) were excluded.

Statistical Analyses

Linkage analysis was carried out using MERLIN [Abecasis et al., 2002]. Estimation of identity by descent (IBD) was based on the allele frequencies in the total genotyped sample and additional genotype data of family members of the affected siblings. Using the Whittemore and Halpern NPL all statistic [Whittemore and Halpern, 1994], MERLIN calculates at which markers affected siblings share more alleles IBD than expected by chance. LOD scores are calculated using the Kong and Cox linear model [Kong and Cox, 1997].

The 1-LOD-drop support interval was used as an estimate for the 90% confidence interval of any QTL locations [Abecasis et al., 2001] and was used to examine linkage replication compared to previously reported locations (using Sullivan Lab Evidence Project http://slep.unc.edu) and candidate genes (using Ensembl: www.ensembl.org and OMIM: www.ncbi.nlm.nih.gov/entrez/query.fcgi?db=omim).

Empirically derived P values were calculated from 1,000 simulated data sets using the gene-dropping method implemented in MERLIN to calculate the threshold LOD-scores for suggestive and significant linkage.

RESULTS

Mean age at the time of the interview was 43.1 years (SD 11.2) and 29.0 years (SD 11.2) for the 1,943 Australian subjects and the 904 Dutch subjects, respectively. Figure 1 shows that 110 Australian and 23 Dutch pedigrees included more than one affected sibling (total N = 278 affected siblings), and thus were eligible for the linkage analysis. From the families with two or more affected offspring, genotypes were available on 35 fathers and 44 mothers and an additional 130 siblings in the Australian sample and on 14 fathers and 14 mothers and an additional 43 siblings in the Dutch sample. Figure 2 shows the linkage results for the analysis of a lifetime diagnosis of MDD and Table II shows the chromosomal regions with an LOD score exceeding 1.0. Simulations yielded an empirically derived suggestive and significant LOD threshold of 1.66 and 2.95. Three regions showed suggestive linkage signals. The highest LOD-score of 2.1 was found on chromosome 17 at 52.6 cm, marker ATA58E08 (90% confidence interval: 46.6–62.5 cM bounded by markers D17S921 and GGA19G04). In addition, LOD scores of 1.9 and 1.7 were found on chromosome 8 at 2.7 cm, marker D8S504 (90% confidence interval: 0–12.4 cM, bounded by 8pter and marker D8S277) and chromosome 2 at 90.6 cM, marker GATA66D01 (90% confidence interval: 78.1–96.9 cM, bounded by markers ATA27D04 and GATA181G08).

DISCUSSION

A linkage analysis on a combined Australian and Dutch sample aiming to identify regions that might harbor genes influencing the vulnerability for MDD found suggestive linkage for two new regions on chromosome 2 and chromosome 17. In addition, we found suggestive linkage on chromosome 8 in a region that showed evidence for linkage in two previous studies, once significant and once suggestive [Cloninger et al., 1998; Fullerton et al., 2003].

The region on chromosome 17 harbors the serotonin transporter gene. In Dutch and Australian samples, partly overlapping with the current samples, no evidence was found for an association with the serotonin transporter gene length s/l polymorphism (5-HTTLPR) and MDD (Dutch and Australian sample) or neuroticism and symptoms of anxiety or depression (in the Dutch sample only) [Gillespie et al., 2005; Middeldorp et al., 2007]. These findings imply that another gene in this region is associated with MDD or another polymorphism in the serotonin transporter gene, for example, a recently identified third variant [Hu et al., 2006]. None of the other usual candidate genes for major depression serotonin receptor 1A (5HT1A), tryptophan hydroxylase (TPH) 1 and 2, catechol-o-methyltransferase (COMT), or brain derived neurotrophic factor (BDNF) were found to lie under one of the linkage peaks [Levinson, 2006].

None of the regions clearly overlapped with syntenic regions found in linkage analyses on emotionality in mice, but the region on

![Fig. 1. Number of Pedigrees (P) with and without affected (A) subjects for the Australian (left) and Dutch (right) samples. Printed in bold are the number of pedigrees with the corresponding number of affected males (M) and females (F) which are included in the linkage analysis.](image-url)
chromosome 2 was very close, with 5 cM between the borders of the confidence intervals [Smoller et al., 2001].

An important limitation of our study, as with many other linkage studies of MDD (Table II), is the small sample size. A power analysis performed in ASP [Krawczak, 2002] showed that we had 10% power to find a P-value < 0.001 (LOD score ~ 2) assuming a lifetime prevalence of MDD of 15%. Therefore, it is likely that other regions that are involved in the development of MDD were not identified. Individual risk variants underlying complex genetic disease are likely to explain only a fraction of the genetic variance [Welcome Trust Case Control Consortium, 2007] but multiple risk variants within a gene or chromosomal region may be more easily detected by linkage than association, and consistently identified regions from linkage analyses are not used to inform prioritization of results from genome-wide association studies. In this study, we identified three regions that exceeded the suggestive threshold for linkage, defined as the threshold expected to be exceeded only once per genome scan.

The current study was carried out in a sub-sample of a larger set of families used for a linkage analysis of the mean neuroticism scores over 4 (Australia) or 5 (The Netherlands) time points [Wray et al., 2008]. Neuroticism is a personality trait measuring emotional instability and those scoring highest on the neuroticism scale express feelings of both anxiety and depression. The motivation for the current study was greater phenotypic homogeneity despite the loss of power in sample size. The three regions of suggestive linkage identified in the neuroticism analysis (10p, 14q, and 18q) showed no evidence for linkage in the current analysis of MDD, although the linkage to 14q is most likely driven by anxiety phenotypes in the Dutch sample [Middeldorp et al., 2008]. Of the three regions identified in the current MDD linkage analysis, chromosome 2 at 90 cM achieved LOD of 1 in the analysis of neuroticism in the Dutch sample, but the regions on chromosomes 8 and 17 showed no evidence for linkage with mean neuroticism score. However, the measure of neuroticism collected at the same interview as the CIDI interview in the Australian sample (AU99) did show evidence for linkage (LOD > 1) across the wide region of 0–85 cM on chromosome 17 [results for individual measures of neuroticism were not presented in Wray et al., 2008], but not at other time points.

To conclude, the region on chromosome 8 around 5 cM has shown most evidence for harboring genes that are involved in the development of MDD. Further research is warranted aiming to identify the specific genes. Future studies are necessary to clarify the significance of our finding on chromosome 2 and 17.

ACKNOWLEDGMENTS

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![FIG. 2. MDD linkage results in a combined Australian and Dutch sample.](image)

**TABLE II. Chromosomal Regions With an LOD Scores >1.0**

<table>
<thead>
<tr>
<th>Chromosome</th>
<th>cM</th>
<th>LOD</th>
</tr>
</thead>
<tbody>
<tr>
<td>17</td>
<td>52.585</td>
<td>2.11</td>
</tr>
<tr>
<td>8</td>
<td>2.652</td>
<td>1.87</td>
</tr>
<tr>
<td>2</td>
<td>90.646</td>
<td>1.7</td>
</tr>
<tr>
<td>15</td>
<td>95.943</td>
<td>1.27</td>
</tr>
<tr>
<td>13</td>
<td>97.493</td>
<td>1.08</td>
</tr>
<tr>
<td>1</td>
<td>221.663</td>
<td>1.04</td>
</tr>
</tbody>
</table>
analyses were carried out on the Genetic Cluster Computer (http://www.geneticcluster.org) which is financially supported by The Netherlands Scientific Organization (NWO 480-05-003). CM was supported by the Hersenstichting Nederland (13F05(2.47) and NWO-ZonMw (91676125). Analytic support was provided by NIMH R01 MH059160 (to PFS). The Australian study was funded by grants to NGM Gavin Andrews from the Australian National Health and Medical Research Council (971232, 339450) and by Gemini Genomics plc (now defunct). We acknowledge the role of Dr. Andrew Heath and NIH grants (AA07535 and AA07728) in earlier projects in which selection variables were collected. For the genome scans we acknowledge and thank: the Mammalian Genotyping Service, Marshfield WI (Director: Dr. James Weber) for genotyping conducted under grants awarded to Dr. Daniel T. O’Connor, Dr. David Duffy, Dr. Dale Nyholt, and Dr. Patrick Sullivan; Dr. Eline Slagboom (Leiden scan); Dr. Peter Reed (Gemini scan); Dr. Jeff Hall (Sequana); Dr. Aarno Palotie (Finnish Genome Center, Helsinki); and Dr. Sue Forrest (Australian Genome Research Facility). We thank our interviewers and clerical and administrative support staff supervised by DJS. We thank David Smyth, Scott Gordon, Harry Beeby, and Olivia Zheng for data management and computer support and our laboratory staff, especially Megan Campbell, Anjali Henders, and Leanne McNeil, for sample processing and preparation. Most of all, we thank the twins and their relatives for their willing participation in the study.

REFERENCES

Duffy DL. 2002. Sib-Pair Version 0.99.9, Brisbane: Queensland Institute of Medical Research.
genome linkage scan of recurrent depressive disorder from the depression network study. Hum Mol Genet 14:3337–3345.


Welcome Trust Case Control Consortium. 2007. Genome-wide association study of 14,000 cases of seven common diseases and 3,000 shared controls. Nature 447:661–678.


