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Meta-analysis of 32 genome-wide linkage studies of schizophrenia

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Abstract

A genome scan meta-analysis (GSMA) was carried out on 32 independent genome-wide linkage scan analyses that included 3255 pedigrees with 7413 genotyped cases affected with schizophrenia (SCZ) or related disorders. The primary GSMA divided the autosomes into 120 bins, rank-ordered the bins within each study according to the most positive linkage result in each bin, summed these ranks (weighted for study size) for each bin across studies and determined the empirical probability of a given summed rank (P_{SR}) by simulation. Suggestive evidence for linkage was observed in two single bins, on chromosomes 5q (142-168 Mb) and 2q (103-134 Mb). Genome-wide evidence for

linkage was detected on chromosome 2q (119-152 Mb) when bin boundaries were shifted to the middle of the previous bins. The primary analysis met empirical criteria for 'aggregate' genome-wide significance, indicating that some or all of 10 bins are likely to contain loci linked to SCZ, including regions of chromosomes 1, 2q, 3q, 4q, 5q, 8p and 10q. In a secondary analysis of 22 studies of European-ancestry samples, suggestive evidence for linkage was observed on chromosome 8p (16-33 Mb). Although the newer genome-wide association methodology has greater power to detect weak associations to single common DNA sequence variants, linkage analysis can detect diverse genetic effects that segregate in families, including multiple rare variants within one locus or several weakly associated loci in the same region. Therefore, the regions supported by this meta-analysis deserve close attention in future studies.

Keywords

genome; human; humans; schizophrenia/genetics; genetic predisposition to disease; linkage (genetics); meta-analysis

Introduction

We report here on a new genome scan meta-analysis (GSMA)¹⁻³ of genome-wide linkage scans (GWLS) of schizophrenia (SCZ). We previously published a GSMA of 20 scans that included 1208 pedigrees with 2945 genotyped affected individuals.⁴ Here we analyze 32 scans that included 3255 pedigrees with 7413 affected individuals.

Genome-wide association (GWA) study methods have proven more successful than GWLS for common diseases: they detect smaller effects of common single nucleotide polymorphisms (SNPs)⁵ as well as some copy number variants (CNVs),⁶ and have identified many common disease associations.⁷ But an adequately powered GWLS can detect a signal from many kinds of susceptibility variants (common SNPs, multiple rare SNPs, tandem repeats and heritable structural variations) in one or more⁸ loci in a region. Meta-analysis can achieve much larger sample sizes than single studies. Consensus linkage regions may deserve further study using methods such as high-throughput resequencing.

Family and twin data⁹ suggest that most of the genetic risk to SCZ is conferred by multiple interacting loci, each causing a small increase in risk. As SCZ has similar symptoms and roughly similar prevalence throughout the world,¹⁰ a reasonable hypothesis is that at least some loci have effects in many populations. SCZ linkage findings do not replicate consistently, and although there may be some true positives in small samples from unique populations, in general, power is probably limited by inadequate sample size and differences in ascertainment, marker sets, ancestry and statistical methods. There must also be some measurement error due to limitations in diagnostic methods. Current diagnostic criteria produce the largest known estimates of heritability (twins) and of increased risk to first-degree relatives,^{11,12} and there is good diagnostic reliability within and across research groups.¹³ But diagnostic judgments are based on data from interview and medical records that vary in completeness and quality. Fortunately diagnostic differences are usually among psychotic disorders that coaggregate in families with similar risks.¹⁴ It is not known whether a further subdivision of cases would increase power.

We present a GSMA of all 32 scan analyses, and a secondary analysis of 22 European-ancestry analyses. The new samples (some of them large) are from European, non-European and isolate populations, but are too diverse to consider a subset other than Europeans. We comment briefly below on an analysis of heterogeneity between Asian and European samples. Statistical significance has been evaluated with permutation tests and simulations. In the primary analysis,

2 bins met empirical genome-wide criteria for suggestive linkage; and a set of 10 bins (in eight chromosomal regions) met an empirical aggregate criterion for genome-wide significant linkage, indicating that loci linked to SCZ are present in some or all of these bins. Five of these bins were consistent with our previous report.⁴ When bin boundaries were shifted to the middle of the previous bins, one genome-wide significant result was observed on chromosome 2q, consistent with our previous report.⁴

Materials and methods

Study selection

We identified GWLS of SCZ (in addition to those studied previously⁴) through PubMed literature searches, conference abstracts and personal contacts with investigators, selecting the largest analysis for each study. Study characteristics are summarized in Table 1. Investigators of each of these studies agreed to contribute data to this analysis. All scans ascertained probands with SCZ by one of three closely related sets of diagnostic criteria. Most scans included relatives with SCZ and those with one or all subtypes of schizoaffective disorder (SA). A few scans included only SCZ cases, or included broader diagnoses in alternative models. Scans were included if informative families ascertained through a SCZ case had at least 30 genotyped SCZ and SA cases. We included only primary genome-wide analyses with few gaps in marker coverage, and did not consider subsequent ‘fine mapping’ with additional markers or pedigrees in selected regions. We excluded a small Utah study⁴⁸ whose RFLP markers were difficult to place on the current map; a study of a large Costa Rican pedigree⁴⁹ with too few affected cases; a small study from an Italian isolate population⁵⁰ that combined SCZ and bipolar cases; and a study of isolated villages in Daghestan⁵¹ that used a very broad phenotype definition. We lacked detailed results for the Icelandic study of Stefansson *et al.*⁵² We included two earlier studies of Icelandic families for which we had complete results,^{15,20} and discuss below a secondary analysis substituting results from Stefansson *et al.* inferred from published graphs.

A companion paper³⁸ reports on a combined analysis of studies 24-31 (Table 1). Older data for all or parts of six of those data sets were included as separate scans in our previous GSMA, so all eight are treated separately here using the newer data.³⁸ To eliminate sample overlap, investigators provided us with data for NUH (study 28) that excluded families that were also in the US/International study (study 9); and data for sample 15 (US/Sweden) without families in sample 26 (Cardiff). The investigators determined from genotypes that there were no overlapping families in samples 10 and 22.

There were 14 studies of European-ancestry pedigrees (including partial isolates such as Finland and Ashkenazim); 8 of both European and non-European families; and 10 non-European samples, including 3 Asian (Japanese, Han Chinese and Indonesian), 2 Latino, 1 Arab, 2 Pacific Island isolates (Palau and Kosrae) and 2 entirely (study 20) or predominantly (study 6) African-American samples. Study 6 (Texas) included a few European-ancestry families for which we lacked separate data. The ALL analysis included all 32 samples; the EUR (European-ancestry) analysis included 22 samples or subsamples (Table 1). Data for chromosome X markers were available for 23 studies, so this chromosome was considered in a secondary analysis.

Linkage scores were obtained from investigators or internet postings. Data for each study were rank ordered based on the designated ‘primary’ analysis, and if this included more than one test or model, we took the most positive linkage score across tests for each chromosomal bin. From our previous GSMA,⁴ data were carried over unchanged for studies 1-10 in Table 1; the Utah study⁴⁸ was omitted as discussed above; and nine studies were updated with new genotyping results, some with larger samples. Studies 13 and 24-31 used SNP markers from

the Illumina version 4 panel; all other studies used microsatellite markers at approximately 8-10 cM density.

Marker mapping

Marker locations were determined from the Rutgers combined linkage map (Build 36),⁵³ or interpolated based on physical position, other maps or flanking markers. Chromosomes were divided into bins of approximately equal genetic widths—120 30-cM bins (our primary analysis), or (for secondary analyses) 179 20-cM bins and 88 40-cM bins. Six additional 30-cM bins were defined on the X chromosome (female map length). If a study had no marker in a bin (for example, the narrower 20 cM bins), we used the mean linkage score for the two flanking markers, or the single closest marker for an empty telomeric bin. Bin boundaries in cM and Mb are listed in online Supplementary Table 1.

Statistical analysis

GSMA is a nonparametric method to combine data generated with different maps and statistical tests.¹⁻³ A GSMA divides the genome into approximately equal-width bins (N_{bins}), labeled $ch.K$, where ch = chromosome and K = bin number (that is, 1.4 is the fourth bin of chromosome 1). For each study, bins are ranked by their highest LOD, NPL or Z score or minimum P -value. Ranks for each sample was weighted by the square root of the number of genotyped affected cases, and then, for comparison, analyzed without weights (results of unweighted analyses are provided in the online Supplementary Materials). Note that '120' was the best rank, and higher summed ranks indicate stronger evidence for linkage (whereas in our previous report,⁴ '1' was the best rank in a study and we reported averaged rather than summed ranks).

We used GSMA software² to evaluate nominal (binwise) significance by randomly permuting the ranks for bins within each study and then resumming the ranks across studies to determine the empirical summed rank probability (P_{SR})—the nominal probability that a bin would achieve or exceed the observed summed rank under the null hypothesis of no linkage in the bin. We also determined the empirical ordered rank P -value (P_{OR})—the probability that the k th highest summed rank achieves or exceeds an observed summed rank under the null). Thresholds for genome-wide significant and suggestive evidence for linkage⁵⁴ were determined empirically (see below). For the primary analysis, these thresholds were $P_{\text{SR}} < 0.00037$ and $P_{\text{SR}} < 0.0077$ respectively.

Strong linkage signals can cover a wide genetic region in complex disorders, and often produce high summed ranks in adjacent bins,¹ but GSMA can miss a weak linkage signal near the boundary of two bins. Therefore, regions showing suggestive evidence for linkage were reanalyzed using 20 and 40 cM bin widths, and using 30 cM bins starting at the midpoint of the bins used in the primary analysis (Supplementary Figure 1).

We tested for heterogeneity in study ranks between three outbred Asian studies (Taiwan, Japan, Indonesia) and the 22 EUR studies, using the nonparametric Wilcoxon rank sum test. We did not test for heterogeneity arising from an arbitrary set of studies;⁵⁵ these tests detect bins where study ranks have a wider spread than expected by chance, but they have low power and do not identify the study subgroups contributing to the heterogeneity.⁵⁶ We also carried out sensitivity analyses, dropping one study at a time and analyzing the remaining studies with GSMA (weighted and unweighted analyses, 30 cM bins).

Simulation study of power and significance thresholds

We performed a simulation study to determine: (1) power; (2) permutation-based estimates of type I error and (3) aggregate criteria for genome-wide significance. Details are provided in Supplementary Online Materials and in a previous report.¹ Briefly, genome-wide data were

simulated for affected sibling pairs under the assumption of no linkage or assuming a range of genetic models (1-10 linked bins with locus-specific sibling relative risks (λ_S) of 1.15 or 1.3, at the edge or near the middle of chromosomes), and 1000 replicates of the 32-sample data set (or of alternative subsets of the data) were created by replacing each sample with a number of ASPs with roughly comparable linkage information.

For estimates of type 1 error, we determined empirical bin-wise thresholds for genome-wide suggestive linkage (the nominal P -value observed on average once per GSMA replicate) or genome-wide significant linkage (the nominal P -value observed on average in 5% of GSMA replicates).

We also determined criteria for aggregate genome-wide significant linkage, that is, that the total number of bins achieving a specified nominal P -value threshold was greater than that observed in 5% of genome-wide replicates with no linkage present. This criterion was determined for ALL and EUR studies by establishing a P -value threshold, P_0 , for which exactly 5% of unlinked replicates had N or more bins with $P_{SR} < P_0$. P_0 was initially set at 0.05, and the number of unlinked replicates with $\geq N$ bins achieving $P_{SR} < P_0$ was tabulated (for $N = 1, 2, 3, \dots$). P_0 was then reduced incrementally until exactly 5% of simulations had at least N bins with $P_{SR} < P_0$, for some value N . For example, for the primary weighted analysis of 32 studies, the 5% genome-wide threshold by this procedure was 10 bins with $P < 0.046$; for EUR studies, it was 10 bins with $P < 0.048$.

For power calculations, we determined how often, on average, the linked bins exceeded the P_{SR} threshold for ALL and for EUR studies, and, to investigate heterogeneity, for ALL studies assuming that linkage arose only in EUR studies.

Results

Evidence for linkage (ALL and EUR studies)

Figure 1 illustrates P -values for all bins in the ALL and EUR analyses, and the figure legend states the empirical P -value thresholds for each study. For ALL studies (3255 pedigrees, 7413 genotyped cases), Table 2 lists the 10 bins (in eight nonadjacent chromosomal regions) that achieved $P_{SR} < 0.05$ in the primary analysis (weighted, 30 cM bins). These 10 bins met the empirical aggregate criterion for genome-wide significant linkage (10 bins with nominal $P_{SR} < 0.046$). Suggestive evidence for linkage was observed for single bins on chromosome 5q (bin 5.6, $P_{SR} = 0.0046$) and chromosome 2q (bin 2.5, $P_{SR} = 0.0075$). Figure 1 illustrates the results for all bins (ALL and EUR analyses), and Figure 2 illustrates the relative ranks of bins in each study. (Results for all analyses are available in Online Supplementary Tables; ranks for each study are available online at www.kcl.ac.uk/mmg/ngdata.html.)

Table 3 summarizes results for 22 EUR samples or subsamples (1813 pedigrees, 4094 genotyped cases). Suggestive evidence for linkage was observed on chromosome 8p ($P_{SR} = 0.00057$, close to the genome-wide threshold for 22 EUR studies of 0.00044), with a further five bins in four different regions achieving nominal significance.

On chromosome X, there were no nominally significant bins for ALL studies; there was one such bin (X.5, 130-162 cM, 119-141 Mb, $P_{SR} = 0.0247$) for EUR studies.

When estimated ranks (interpolated from published graphs) for the Icelandic study of Stefansson *et al.*⁵² were substituted for the results of Gurling *et al.*²⁰ and Moises *et al.*,¹⁵ results were similar to those shown in Table 1 except that bin 3.4 failed to achieve nominal significance. For EUR studies, results again were similar except that bins 8.2 and 2.8 both achieved suggestive significance.

Effect of bin width

Online Supplementary Figures 2-4 illustrate results of analyses using four alternative bin widths (20, 30 and 40 cM, and 30 cM bins shifted by 50% for chromosomes with suggestive evidence for linkage (chromosomes 2 and 5 for ALL, chromosome 8 for EUR). On chromosome 2, genome-wide significance was obtained for the shifted 30 cM bin ($P_{SR} = 0.00035$, 132.2-161.6 cM, 118.7-152 Mb), and suggestive linkage evidence was seen in the adjacent bin (102.8-132.2 cM, 84.9-118.7 Mb). For the region of chromosome 5q that produced suggestive evidence for linkage in the primary analysis, the signal was similar regardless of bin width. On chromosome 8, using 20 cM bins extended the signal by approximately 5 cM in both directions; if linkage is present, this pattern could indicate a broader signal (multiple loci) or the effects of a single strong signal on adjacent bins.

Heterogeneity and sensitivity analyses

In tests of differences between 22 EUR vs 3 Asian studies, none of the 120 bins exceeded nominal $P < 0.01$, and only 4 bins achieved nominal $P < 0.05$, thus, no significant difference was observed. The most significant evidence for heterogeneity between the Asian and European studies was in bin 1.5, but this did not reach the suggestive threshold (online Supplementary Figure 5). Results of sensitivity analyses, omitting each study in turn, are shown in online Supplementary Figures 6 and 7. The largest changes: bin 8.2 (8p) achieves the suggestive linkage threshold without the Taiwan data set; bin 2.8 (2q) becomes substantially more significant (but not achieving genome-wide significance) without the Japan data set; bin 5.6 (5q) becomes substantially more significant (but not genome-wide significant) without the Suarez *et al.* data set (study 21); and bin 6.3 (6pq) would join the list of nominally significant bins without either the Taiwan or Japan data set.

Type I error

Type I error was calculated for the unlinked GSMA simulations (120 000 30-cM bins) by determining the number of observed bins in which the GSMA program computed P_{SR} values (based on permutation alone) that were less than the theoretical threshold for nominal significance (0.05); for genome-wide suggestive linkage, or a value expected once per GSMA ($1/120 = 0.00833$) or genome-wide significant linkage ($0.05/120 = 0.00042$). The GSMA program's permutation procedure produced type I errors that were slightly liberal for ALL (0.053, 0.0096 and 0.00051) and less liberal for EUR studies (0.051, 0.0091 and 0.00042). The source of the discrepancy is not clear, but we have used the permutation-based suggestive and significant values here for single bins (0.0077 and 0.00037 for ALL, and 0.0078 and 0.00044 for EUR studies) and an empirical threshold for aggregate genome-wide significance as described above.

Power

Table 4 summarizes the results of power calculations from simulation studies. Power is lower for edge bins for multipoint analyses, so we computed a weighted average of power assuming 20% edge and 80% mid-chromosome bins (see Table 4 legend). Power was excellent to detect significant linkage at multiple loci with observed population-wide $\lambda_S = 1.3$ (30% increased risk in sibs) across ALL studies or in EUR studies alone. For $\lambda_S = 1.15$, there was excellent power to detect significant linkage in multiple bins (for ALL studies), or to detect suggestive evidence for linkage in multiple bins (if limited to EUR studies); there was reduced power in the presence of substantial heterogeneity. Genetic effects limited to a defined subset (EUR) were more readily detected in a separate analysis.

Discussion

Meta-analysis of 32 GWLS, encompassing 7413 individuals affected with SCZ or related disorders in 3255 pedigrees, produced evidence for significant linkage in the genome based on empirical aggregate criteria: 10 30-cM bins that achieved bin-wise $P_{SR} < 0.046$. Genome-wide evidence for linkage was detected on chromosome 2q (118.7-152 Mb) in a secondary analysis that shifted bin location by 50%, consistent with our previous finding of significant linkage in the same region in our previous GSMA of 20 studies.⁴ Suggestive evidence for linkage was detected in ALL studies in two single bins, on chromosomes 5q and 2q. Separate analysis of European-ancestry samples produced linkage evidence falling just short of the genome-wide significance in bin 8.2 (15.7-32.7 Mb).

It is widely assumed that many loci contribute to SCZ susceptibility. Table 4 shows that for the GSMA method, power to detect at least one true linkage goes up with the number of linked loci, but the power to detect any specific locus decreases, because in a rank-ordered test there is a finite number of high ranks in each study. But if there were many loci with locus-specific λ_S values of 1.3 or even 1.15 (averaged across all study populations and accounting for the effects of various sources of measurement error), we would expect to detect genome-wide significant linkage in several individual bins, and suggestive evidence for linkage in many more than two bins. It would be reasonable to conclude that there are probably no loci with such strong effects worldwide or across all European populations, although they might exist within particular populations for which samples of this size are not available or within subsets of families, which we cannot yet identify in advance.

The power of linkage analysis for a locus is predicted by its contribution to the relative risk (RR) to siblings of probands (λ_S),⁵⁷ whereas the power of association analysis is predicted by allelic or genotypic RRs, that is, the increased risk to individual carriers.⁵ GWA studies have produced robust association findings for many complex genetic disorders⁵⁸⁻⁶⁰ typically with RRs of 1.1-1.5. A variant found on 10% of chromosomes conferring a 'large' RR of 1.5 per allele in a multiplicative model (2.25 for homozygous carriers) would produce a population-wide sibling RR of only 1.02 (undetectable by linkage). But GWA chips are unlikely to tag all pathogenic variants. For example, in NOD2 there are three rare variants, each with a frequency of less than 5%, that confer moderate risks of Crohn's disease. Their individual effects generate only weak signals when assayed by the Affymetrix 500K chip because they are poorly tagged; but combined, they confer a sibling RR of 1.16, which is detectable in a very large linkage sample.^{58,61} This illustrates a strength of linkage analysis: allelic heterogeneity is difficult to detect in association studies, as the genetic effect is split across variants, but linkage can detect the pooled effect of all variants (not only SNPs) within one or more susceptibility genes or elements in a region.

Readers may be interested to know the location of SCZ candidate genes or regions in relation to bins that produced evidence for linkage here. On chromosome 1q, NOS1AP (160 Mb; previously known as CAPON),^{62,63} RGS4 (161.3 Mb)^{64,65} and UHMK1 (160.7 Mb)⁶⁶ are near the telomeric edge of bin 1.6 (114.6-162.1 Mb), and the rare (usually *de novo*) SCZ-associated CNV is located in a different part of the same bin (145-146 Mb).^{67,68} On chromosome 8p, PPP3CC (22.35 Mb)⁶⁹ and NRG1 (31.6-32.7 Mb)^{52,70} are both within bin 8.2 (15.7-32.7 Mb). On chromosome 2q, ZNF804A, which contains a SNP that produced genome-wide significant evidence for association in a combined SCZ-bipolar disorder sample,⁷¹ is in bin 2.7 at 185 Mb. This bin was not nominally significant in the primary analysis, but 185 Mb is within nominally significant bins in two secondary analyses (one that shifted bin boundaries by 50% and one that considered 40 cM bin widths). DTNBP1, DAOA, TRAR4/STX7, CHRNA7, COMT/ARVCF, DISC1, AKT1, HTR2A and DRD2 are not in bins listed in Tables 2 or 3. Many of the research groups whose samples are included here are actively

investigating association of SCZ to sequence variants in genes identified in linkage regions or by the more recent genome-wide association methodology.

In conclusion, this updated GSMA of 32 SCZ studies showed significant evidence for linkage on chromosome 2q in a secondary analysis, suggestive evidence for linkage on chromosomes 5q and 2q in the primary analysis, and suggestive evidence for linkage on 8p in European samples. Genome-wide significant aggregate evidence for linkage (that is, more modestly significant results than expected by chance) was observed based on results for 10 bins in 8 nonadjacent chromosomal regions. As results from genome-wide association studies emerge, it may be important to keep in mind that there may be SCZ susceptibility loci in some of these consensus linkage regions that cannot be detected by common tag SNPs. It is likely that diverse methods will be required to identify specific DNA sequence variants and to confirm and define their role in the etiology of SCZ.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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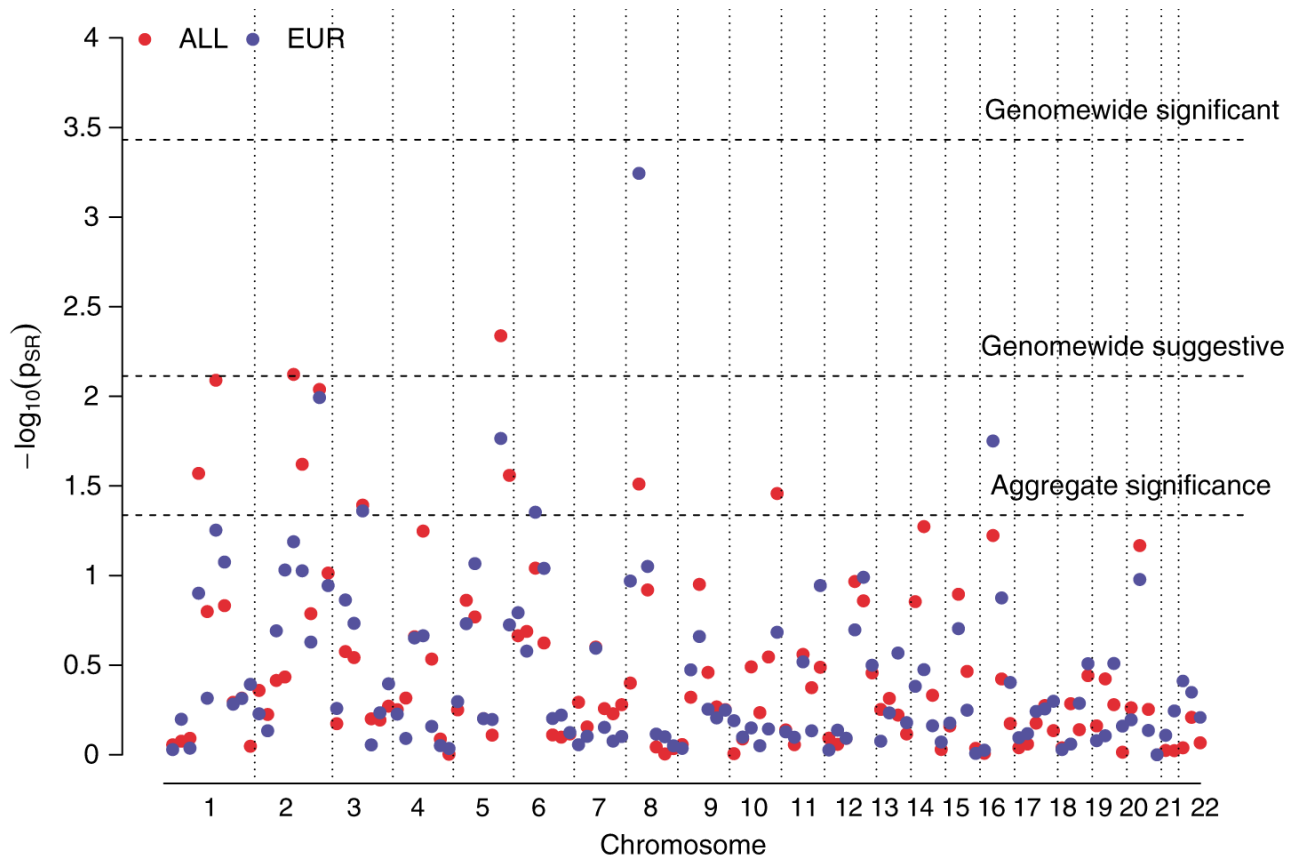


Figure 1.

Genomewide results for primary analysis (weighted, 30 cM bins) for ALL studies (red) and EUR studies (blue). The two upper dashed lines represent empirical thresholds for genomewide significant (theoretical $P_{SR} = 0.00037$) and suggestive (theoretical $P_{SR} = 0.0077$) evidence for linkage for a single bin for ALL studies; and the lower dashed line represents the threshold for aggregate significance for ALL studies (theoretical $P_{SR} = 0.046$ in 10 or more bins). The thresholds for EUR studies were slightly less stringent (P_{SR} values of 0.00044, 0.0078 and 0.0475 (in 10 or more bins) respectively), but the Figure shows all EUR bins within the region for their correct threshold of significance.

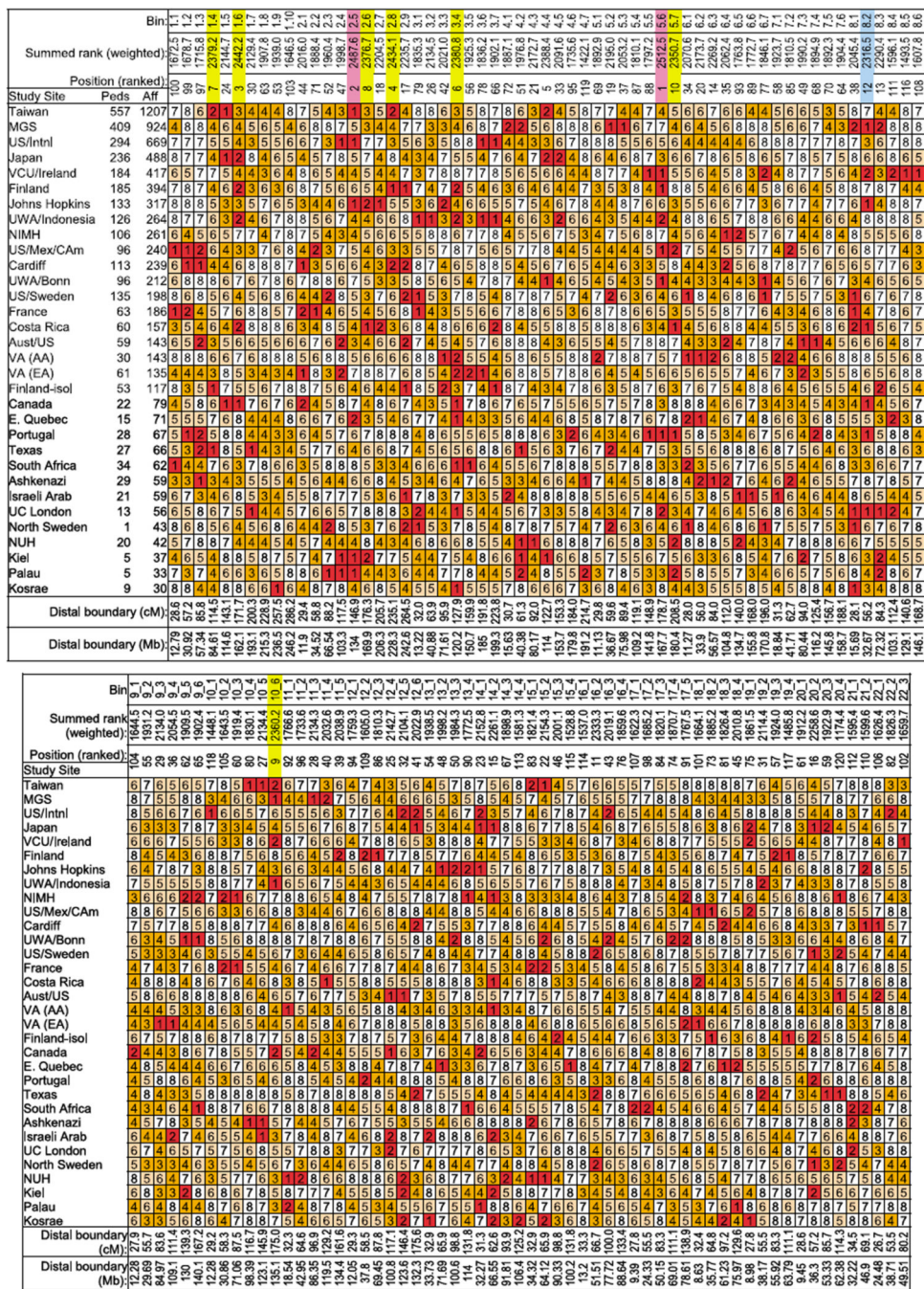


Figure 2. Ranks by study and summed ranks. Shown for each study is the within-study rank of each bin. '1' against a red background indicates bins with ranks of 116-120 (the bin containing the most significant linkage score in a study is ranked 120); 2 indicates ranks 111-115; 3—ranks 101-110; 4—ranks 81-100; 5—ranks 61-80; 6—ranks 41-60; 7—ranks 21-40; and 8—ranks 1-20 (cells with 7 and 8 have a white background). Shown at the top of each part of the figure are the bin number, the weighted summed rank for each bin across studies (i.e., weighted for each sample by the square root of the N of genotyped affected individuals), and the rank-ordered position of that bin in the ALL analysis (i.e., 1 is the best position, see details in Table 2). Purple background indicates bins with empirical suggestive evidence for linkage in the ALL.

analysis; 10 bins shaded in purple, yellow or blue met the threshold for aggregate genome-wide evidence for linkage (10 bins with nominal $P < 0.046$); and bin 8.2 (shaded blue) also met empirical suggestive evidence for linkage in the EUR analysis. Shown at the bottom is the distal boundary of each bin in Rutgers cM and in Mb (build 36). Note that tied ranks can result in uneven numbers of bins in a grouping, particularly for lower ranks when there are many zero or negative scores. See Table 1 for the references associated with each study.

Table 1
Genome-wide linkage scan characteristics (ALL and European-ancestry analyses)

	Study site or project ^a	Diagnostic criteria ^b	ALL analysis		EUR analysis		Markers	Program	Statistics	Ethnicity ^c
			Ped	Aff	Ped	Aff				
1	Kiel ¹⁵ (p)	SCZ, SA (RDC or D3R)	5	37	5	37	413	WRPC	P-values	Icelandic
2	Palau ¹⁶ (p)	SCZ, SAS (RDC)	5	33			496	LINKAGE	LOD	Palau
3	Canada ¹⁷ (p)	SCZ, SA (D3R)	22	79	22	79	381	FASTLINK	HLOD	Celtic Canada
4	Finland— ¹⁸ isolate (p)	SCZ; +SA (D4)	53	117	53	117	345	MLINK	LOD	Finnish
5	Finland— ¹⁸ national (p)	SCZ; +SA (D4)	185	394	185	394	345	MLINK	LOD	Finnish
6	Texas ¹⁹ (p)	SCZ (D4)	27	66			406	Genehunter Plus	NPL	AA, few EA
7	UC London ²⁰ (p)	SCZ,SA,UFP (RDC)	13	56	13	56	365	MFLINK	P-values	Icelandic
8	North Sweden ²¹ (p)	SCZ;+SAD;+PNOS;+SAB (D4)	1	43	1	43	371	LINKAGE	LOD	Swedish
9	US/International ²² (p)	SCZ, SA (D3R)	294	669	294	669	396	Mapmaker/Sibs	MLS	EA (inc Chile)
10	Costa Rica ²³ (p)	SCZ, SA (D4)	60	157			404	Genehunter Plus	KCLOD	Costa Rican
11	Kosrae ²⁴ (n)	SCZ, SA (D4)	9	30			398	FASTLINK	LOD	Kosrae
12	Israeli Arab ²⁵ (n)	SCZ, SAD (D4)	21	59			350	Allegro	NPL	Arab
13	Japan ²⁶ (n)	SCZ, SA (D4)	236	488			5861	Merlin	KCLOD	Japanese
14	Ashkenazi ²⁷ (n)	SCZ, SA (D4)	29	59	29	59	382	Genehunter	LOD	Ashkenazi
15	US/Sweden ²⁸ (m)	SCZ, SA (D4)	135	198	135	198	392	Mapmaker/Sibs	MLS	Swedish, US
16	Portugal ²⁹ (n)	SCZ, SAD (D4)	28	67	28	67	366	Genehunter	NPL	Portuguese
17	South Africa ³⁰ (n)	SCZ, SAD (D4)	34	62	34	62	388	Merlin	KCLOD	Afrikaner
18	Eastern Quebec ³¹ (n)	SCZ (D3R)	15	71	15	71	350	FASTLINK	LOD	Quebecois
19	VA (EA) ³² (n)	SCZ, SAD (D3R)	61	135	61	135	414	Merlin	KCLOD	EA
20	VA (AA) ³² (n)	SCZ, SAD (D3R)	60	143			414	Merlin	KCLOD	AA
21	MGS ³³ (n)	SCZ, SA (D4)	409	924	263	571	400	Genehunter Plus	Zlr	EA, AA
22	US/Mexico/Central America ³⁴ (n)	SCZ, SA (D4)	96	240			404	Merlin	KCLOD	Mexican, Cent Am
23	Taiwan ³⁵ (n)	SCZ (D4)	557	1207			286	Merlin	Zlr	Han Chinese
24	Aust/US ³⁶⁻³⁸ (m)	SCZ, SA (D3R)	59	143	49	114	5868	Allegro	Zlr	EA, AA, Other

	Study site or project ^a	Diagnostic criteria ^b	ALL analysis		EUR analysis		Markers	Program	Statistics	Ethnicity ^c
			Ped	Aff	Ped	Aff				
25	UWA/Bonn ^{38,39} (m)	SCZ, SA (D3R)	96	212	96	212	5868	Allegro	Zlr	German, Israeli
26	Cardiff ^{28,38} (m)	SCZ, SA (D3R)	113	239	113	239	5868	Allegro	Zlr	UK
27	France/LaR�union ^{38,40,41} (n)	SCZ, SA (D3R)	63	186	29	75	5868	Allegro	Zlr	EA, AA, Other
28	NUH ^{38,42} (n)	SCZ, SA (D3R)	20	42	19	40	5868	Allegro	Zlr	EA, AA
29	Johns Hopkins ^{38,43} (m)	SCZ, SA (D3R)	133	317	124	295	5868	Allegro	Zlr	EA, AA, Other
30	NIMH ^{38,44,45} (m)	SCZ, SA (D3R)	106	261	61	142	5868	Allegro	Zlr	EA, AA, Other
31	VCU/Ireland ^{38,46} (m)	SCZ, SAPO (D3R)	184	417	184	417	5868	Allegro	Zlr	Irish
32	UWA/Indonesia ^{38,47} (n)	SCZ (D3R)	124	267			400	Genehunter	MLS	Indonesian
	Total		3283	7476	1835	4141				

Other abbreviations: Ped = informative pedigrees; Aff = genotyped affected cases. VA = US Veterans’ Administration; MGS = Molecular Genetics of Schizophrenia collaboration; Aus/US = Australia/US collaboration; NUH = NorthShore University HealthSystem Research Institute; NIMH = National Institute of Mental Health Schizophrenia Genetics Initiative; VCU = Virginia Commonwealth University; UWA = University of Western Australia. LOD = log of the odds ratio under one or more parametric models; HLOD = heterogeneity LOD scores (generally, maximized over at least two models); NPL = nonparametric linkage Z-score; MLS = maximum LOD score (affected sib-pair analysis, constrained to possible triangle); Zlr = Z-likelihood ratio statistic; KCLOD = Kong-Cox equivalent LOD score computed from Zlr.

^aSorted by year of publication of the data used in this analysis. The letter in parentheses indicates whether the dataset was included in the previous analysis (p), has been modified with new genotyping and/or additional pedigrees (m), or is new (n).

^bD3R = DSM-III-R criteria; D4 = DSM-IV criteria; RDC = Research Diagnostic Criteria; SA = schizoaffective disorder (SAS = RDC mainly schizophrenic type; SAD = depressed type; SAB = bipolar type; SAPO = poor outcome; PNOS = psychosis not otherwise specified; UFP = unspecified functional psychosis (RDC). Diagnoses separated by commas constitute one model; sets separated by semicolons are alternative models over which scores were maximized, with ‘+’ indicating that diagnoses were added to the ones in the next narrower model.

^cEA = mixed European ancestry (usually from the US, but also mixed US, European and Australian samples); AA = African American.

Table 2
Bins achieving nominally significant evidence for linkage (ALL studies)

<i>Pos</i>	<i>Bin</i>	<i>cM</i>	<i>Mb</i>	<i>P_{SR}</i>	<i>P_{OR}</i>
1	5.6*	148.9-178.7	141.8-167.7	0.00459	0.43156
2	2.5*	117.5-146.9	103.3-134.0	0.00755	0.22798
3	1.6*	143.1-171.7	114.6-162.1	0.00814	0.06639
4	2.8	205.7-235.1	206.3-228.3	0.00916	0.01970
5	2.6*	146.9-176.3	134.0-169.9	0.02395	0.14269
6	1.4	85.8-114.5	57.3-84.6	0.02692	0.08449
7	5.7	178.7-208.5	167.7-180.4	0.02760	0.03430
8	8.2*	28.1-56.2	15.7-32.7	0.03086	0.02070
9	10.6	145.9-175.0	123.1-135.1	0.03487	0.01350
10	3.4	95.9-127.9	71.6-120.2	0.04047	0.01140

Shown are the bins with nominally significant evidence for linkage ($P_{SR} < 0.05$), for ALL studies (30cM bins, weighted analysis), ordered by their position (1 = best) according to weighted analysis P_{SR} (summed rank P -value). P_{OR} : ordered rank P -value. Bold P_{SR} values achieved an empirical threshold for suggestive evidence for linkage ($P_{SR} < 0.0077$). No single bin achieved an empirical genome-wide significant threshold (0.00037). Empirically (see text), 10 or more bins with nominal $P_{SR} < 0.046$ was observed in 5% of simulated replicates of this dataset in the absence of linkage. Therefore, it is likely that at least some of the bins listed here contain loci linked to SCZ. Asterisks indicate which regions were consistent with our previous GSMA⁴ of 20 schizophrenia scans (i.e., bins that overlap with those in the previous analysis that met aggregate significance criteria—that analysis used the Marshfield rather than the Rutgers map, so bin boundaries were not identical). See Figure 2 for further details.

Table 3
Bins achieving nominally significant evidence for linkage (European studies)

<i>Pos</i>	<i>Bin</i>	<i>cM</i>	<i>Mb</i>	<i>P_{SR}</i>	<i>P_{OR}</i>
1	8.2*	28.1-56.2	15.7-32.7	0.00057	0.06659
2	2.8	205.7-235.1	206.3-228.3	0.01016	0.35077
3	5.6*	148.9-178.7	141.8-167.7	0.01718	0.33717
4	16.2*	33.3-66.7	13.2-51.5	0.01775	0.15808
5	3.4	95.9-127.9	71.6-120.2	0.04359	0.62724
6	6.3*	56.0-84.0	33.9-56.6	0.04433	0.44626

Shown are bins with nominally significant evidence for linkage ($P_{SR} < 0.05$), in 22 samples or subsamples of European-ancestry families (see Table 1); the bolded p-value met empirical criteria for suggestive evidence for linkage ($P_{SR} < 0.0078$). The results here did not meet aggregate criteria for significant genome-wide evidence for linkage in this subset of the data (10 bins with nominal $P_{SR} < 0.0475$). See legend for Table 2 for additional details.

Table 4

Power to detect linkage in simulation studies (weighted GSMA)

Level of significance												
Number of linked bins:	Nominal				Suggestive				Genome-wide			
	1	5	10	1	5	10	1	5	10	1	5	10
$\lambda_S = 1.3$												
ALL studies	1.000	1.000	1.000	1.000	1.000	0.998	1.000	0.996	0.998	1.000	0.996	0.975
European samples	1.000	0.998	0.993	1.000	0.992	0.953	1.000	0.912	0.953	0.950	0.912	0.745
ALL, Heterogeneity ^a	0.988	0.952	0.918	0.898	0.834	0.705	0.898	0.474	0.705	0.600	0.474	0.325
$\lambda_S = 1.15$												
ALL studies	1.000	1.000	0.989	0.996	0.980	0.932	0.996	0.850	0.932	0.904	0.850	0.705
European samples	0.970	0.948	0.908	0.874	0.822	0.697	0.874	0.468	0.697	0.598	0.468	0.323
ALL, Heterogeneity ^a	0.864	0.774	0.740	0.592	0.546	0.403	0.592	0.180	0.403	0.186	0.180	0.116

Shown is the power of weighted GSMA to detect linkage in the simulation study of samples of affected sibling pairs with information approximately comparable to the 32 studies in the schizophrenia GSMA (120 30-cM bins). λ_S = relative risk to siblings of case probands vs population risk. For 1 linked bin per simulated genome, power is a weighted average assuming that 20% of bins lie at a chromosome end ('edge,' see text), and 80% of are mid-chromosomal ('mid'). Simulations with 5 linked loci included 1 edge and 4 mid bins; those with 10 linked included 2 edge and 8 mid bins.

^a Heterogeneity analyses modelled cases where all samples were included, but European samples were linked as shown while non-European studies were unlinked.