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How should we construct psychiatric family history scores? A comparison of alternative approaches from the Dunedin Family Health History Study

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Abstract

Background—There is increased interest in assessing the family history of psychiatric disorders for both genetic research and public health screening. It is unclear how best to combine family history reports into an overall score. We compare the predictive validity of different family history scores.

Method—Probands from the Dunedin Study ($n=981$, 51% male) had their family history assessed for nine different conditions. We computed four family history scores for each disorder: (1) a simple dichotomous categorization of whether or not probands had any disordered first-degree relatives ; (2) the observed number of disordered first-degree relatives ; (3) the proportion of first-degree relatives who are disordered; and (4) Reed's score, which expressed the observed number of disordered first-degree relatives in terms of the number expected given the age and sex of each relative. We compared the strength of association between each family history score and probands' disorder outcome.

Results—Each score produced significant family history associations for all disorders. The scores that took account of the number of disordered relatives within families (i.e. the observed, proportion, and Reed's scores) produced significantly stronger associations than the dichotomous score for conduct disorder, alcohol dependence and smoking. Taking account of family size (i.e. using the proportion or Reed's score) produced stronger family history associations depending on the prevalence of the disorder among family members.

Conclusions—Dichotomous family history scores can be improved upon by considering the number of disordered relatives in a family and the population prevalence of the disorder.

Keywords

Family history; predictive validity; prevalence; psychiatry

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Declaration of Interest

None.

Introduction

Family history is a major risk factor for most psychiatric disorders (Kendler *et al.* 1997; Miles *et al.* 1998; Sullivan *et al.* 2000; Bandelow *et al.* 2002, 2004; Byrne *et al.* 2002; Klein *et al.* 2003; Qin *et al.* 2005; Newman & Bland, 2006; Coelho *et al.* 2007). There has been a recent revival of interest in the family history method for both genetic research and public health screening. For example, gene-association studies and genome-wide studies often now collect family history information so as to improve statistical power to detect associations (Thompson *et al.* 2004; Prescott *et al.* 2005; Chen *et al.* 2007). Family history is also being promoted for disease prevention (Yoon *et al.* 2002).

Typically, family history is assessed by collecting reports of disorder for members of a target person's (or proband's) family. Once these data have been collected, the question becomes, what to do with them? That is, how should family history reports for a disorder be combined into an overall family history score for that disorder? Will a simple family history positive/negative dichotomy based on the presence or absence of disorder among family members predict proband outcome well enough, or is a more elaborate scoring method warranted?

Surprisingly few studies have compared the performance of different family history scores. Most have focused on whether simple dichotomous scores can perform as well as more complex scores which take account of density of disorder (i.e. the number of family members who have the disorder). Results suggest that density scores perform better than dichotomous scores for alcohol dependence (Turner *et al.* 1993; Stoltenberg *et al.* 1998), but not for depression (Sullivan *et al.* 1996). The relative performance of density *versus* dichotomous scores for other psychiatric disorders has not been investigated. Moreover, whereas research in other health domains has investigated whether factors such as family size and family demographic structure (i.e. the age and sex of each family member) affect the predictive validity of family history scores (Silberberg *et al.* 1999; Murad *et al.* 2007), these factors have yet to be investigated in psychiatry.

In this study we sought to determine the effect of taking account of density of disorder, family size and family demographic structure on the predictive validity of family psychiatric history scores (operationally defined as the ability of a family history score for a disorder to predict the same disorder in probands). We were able to assess the predictive validity of family history scores for seven psychiatric conditions (major depressive episode, anxiety disorder, suicide attempt, schizophreniform disorder, conduct disorder, alcohol dependence and drug dependence) as well as for two non-psychiatric outcomes: smoking and asthma. By assessing both psychiatric disorders and non-psychiatric outcomes, we were able to determine whether our findings about the predictive validity of family history scores were specific to psychiatric disorders or could be generalized to other areas of health.

The findings we report are from a new family history study, the Dunedin Family Health History Study (DFHHS). Thus, this report also serves to describe the methods of the DFHHS.

Method

Participants

All probands and informants gave informed consent before participating. Study protocols were approved by the Otago Ethics Committee.

Probands—Probands are members of the Dunedin Study, a 1-year birth cohort constituted at 3 years of age when investigators enrolled 91% of consecutive eligible births between April 1972 and March 1973 in Dunedin, New Zealand (Moffitt *et al.* 2001). Follow-up assessments were conducted at ages 5, 7, 9, 11, 13, 15, 18, 21, 26 and 32 years. Here we report data from 981 probands (97% of the living sample of 1015, 51% male) who had available family history data. Cohort families represent the full range of socio-economic status in New Zealand's South Island and are primarily white.

Informants—The DFHHS aimed to obtain informant-reported family history for three generations of the proband's family. To achieve this aim, we recruited up to three informants per family: the proband and both of the proband's parents. Fig. 1 summarizes information about informants and family members from the DFHHS.

Proband reports: A total of 937 probands (92% of the living sample) provided information on the psychiatric health history and smoking of their biological siblings (aged >10 years) and parents. Family history interviews typically took place at the Dunedin Research Unit when probands were aged 32 years (2004–2005). Interviews were conducted by trained research interviewers who were blind to the proband's previous data and to the data provided by the parent. Probands did not report on their grandparents as we were not confident that they would have sufficient knowledge to report accurately, nor did they report on asthma as they were not interviewed about their family members' physical health at the age 32 years assessment.

Parent reports: Maternal and paternal informants provided information on the psychiatric health history, smoking and asthma of the probands' biological siblings (aged >10 years), parents and grandparents. Family history interviews typically took place in the home of the parent during 2003–2006, when probands were aged 30–33 years. Interviews were conducted by trained research interviewers who were blind to the data provided by the proband. We aimed to interview the biological mother and father of all living probands ($n=1015$), but sought alternative informants when a biological mother or father was either deceased or unable to be interviewed. A total of 884 biological mothers participated. Of the 131 living probands for whom a biological mother could not be interviewed, 51 had a non-biological mother or a maternal aunt or uncle who agreed to participate. Thus, we achieved maternal informants for 935 of the 1015 living probands (92%). A total of 756 biological fathers participated. Of the 259 living probands for whom a biological father could not be interviewed, 98 had a non-biological father or a paternal aunt or uncle who agreed to be interviewed. Thus, we achieved paternal informants for 854 of the 1015 living probands (84%).

Family members—All three informants reported for members of 790 families (80.5%), two informants reported for members of 154 families (15.7%), and one informant reported for members of 37 families (3.8%). Combining data from the three informants resulted in data for 7856 biological family members from 981 proband families (average of 8.0 family members; range 3–16). These include 3764 first-degree relatives (981 biological mothers, 974 biological fathers and 1809 full siblings) and 4092 second-degree relatives (1845 biological grandmothers, 1817 biological grandfathers and 430 half siblings).

Measures

Family history reports—Family history reports were obtained from each informant as follows. First, we obtained a family pedigree list for the proband by asking about the sex, age (or age at death) and relatedness of the proband's biological family members, following the methods of Thompson *et al.* (1980). Second, the psychiatric health history of each family

member was assessed using the Family History Screen (FHS; Weissman *et al.* 2000). We chose this instrument because it has been shown to be valid and reliable, as well as economical for data collection in large samples, and feasible when some family members are not available for direct interview (Weissman *et al.* 2000). To minimize under-reporting, the FHS uses pairs of questions to ascertain each symptom. A broadly sensitive ‘introductory-screen’ question is first asked to stimulate memory and give the informant time to reflect [e.g. ‘Has (names on the pedigree list) ever had a sudden spell or attack in which they felt frightened or panicked?’]. If any family members are named in response to the introductory question, this is followed by a second, narrower ‘symptom-definition’ question [e.g. ‘Has (names on the pedigree list) had several attacks, of extreme fear or panic, even though there was nothing to be afraid of?’]. For data analysis purposes, only the second question in a pair is used.

To broaden the FHS’s coverage, we added items drawn from the Diagnostic Interview Schedule (DIS; Robins *et al.* 1981, 1995), the Short Michigan Alcoholism Screening Test (Selzer *et al.* 1975) and the Drug Abuse Screening Test (Skinner, 1982). We also added a checklist of psychiatric conditions commonly understood by the public (e.g. ‘alcoholism’, ‘depression’), the asthma item from the National Heart, Lung and Blood Institute Family Heart Study (Higgins *et al.* 1996) and an item asking whether family members were ever a smoker. In total, there were symptom-definition items pertaining to major depressive episode (four items), anxiety (13 items on generalized anxiety, panic, agoraphobia, phobia and obsessive-compulsive disorder), schizophreniform disorder (eight items), conduct disorder (eight items), alcohol dependence (three items), drug dependence (three items), suicide attempt (two items), smoking (one item) and asthma (one item). Following the recommendations of Vandeleur *et al.* (2008), a family member was considered to have a positive history of a disorder if one or more of the disorder’s items were endorsed by at least 50% of informants (i.e. two of three informants, one of two informants, or one of one informant).

Family history scores—One dichotomous score and three density scores were computed. Each score took account of biological relatedness by giving first-degree relatives double the weight of second-degree family members (as described below). The scores were as follows.

(1) Dichotomous: This defines probands as ‘family history positive’ if one or more of a proband’s first-degree family members (or, equivalently, two or more of a proband’s second-degree family members) had a positive history of disorder.

(2) Observed number of disordered family members, weighted by relatedness: This is a simple count of the number of family members with a positive history of disorder, with a count of ‘1’ given for each first-degree family member and a count of ‘0.5’ given for each second-degree family member. This method takes account of density of disorder (i.e. number of family members with disorder).

(3) Proportion of disordered family members, weighted by relatedness: This is the count in (2) above, divided by the number of family members (again, with a count of ‘1’ given for each first-degree family member and a count of ‘0.5’ given for each second-degree family member). This method takes account of density of disorder and family size.

(4) Reed’s score: This is $(O-E)/\sqrt{E}$, where O is the observed number of family members with a positive history of disorder (i.e. the count in (2) above), and E is the expected number of family members with a positive history of disorder, given the age and sex of each family member and the age- and sex-specific prevalence of the disorder. Simulations suggest

Reed's score performs best of all methods relating observed to expected rates (Murad *et al.* 2007). We made two modifications to the method of Reed *et al.* (1986). First, to take account of relatedness (in line with scores above), first-degree relatives were given double the weight of second-degree relatives when calculating observed and expected totals. Second, because we did not collect the data necessary to determine age-at-first-incidence of disorder for each family member in our sample, we calculated expected rates based on prevalence of disorder and not incidence of disorder. Thus, for each disorder, we determined the expected probability of a family member ever having had the disorder by calculating the prevalence of that disorder by sex and 5-year age band (20, 21–25, ..., 86–90, >90 years), using data derived from probands and family members in this sample. Reed's score takes account of density of disorder, family size and family demographic structure (i.e. the age and sex of each family member).

Proband outcomes—There were proband outcomes for each disorder for which family history data were available (i.e. for seven psychiatric disorders, as well as for smoking and asthma).

Psychiatric disorders: The psychiatric assessment of Dunedin probands has been described in detail elsewhere (Kim-Cohen *et al.* 2003). Briefly, proband psychiatric disorder was assessed using the Diagnostic Interview Schedule for Children (Costello *et al.* 1982) at younger ages (11–15 years) and the DIS (Robins *et al.* 1981, 1995) at older ages (18–32 years), with a past-year reporting period at each age. At ages 11, 13 and 15 years diagnoses were made using the then-current *Diagnostic and Statistical Manual of Mental Disorders* (DSM), version 3 (DSM-III; APA, 1980), at ages 18 and 21 years according to the then-current DSM-III-R (APA, 1987) criteria, and at ages 26 and 32 years according to DSM-IV (APA, 1994) criteria. Diagnoses were derived as follows. We assessed conduct disorder in childhood: those who were diagnosed with conduct disorder at any of ages 11, 13, 15 or 18 years were considered to have a childhood diagnosis of conduct disorder. In adulthood we assessed major depressive episode, anxiety (generalized anxiety disorder, phobia, agoraphobia, panic disorder, obsessive-compulsive disorder and post-traumatic stress disorder), schizophreniform disorder, alcohol dependence and drug dependence. For each disorder, those who were diagnosed with that disorder at any of ages 21, 26 and 32 years were considered to have an adult diagnosis of disorder. We assessed suicide attempt in adolescence and adulthood: those who reported a lifetime suicide attempt at any of ages 15, 18, 21, 26 or 32 years were considered to have attempted suicide in their lifetime.

Smoking: Those who reported that they had smoked daily for at least 1 month of the previous year at any of the assessments at ages 15, 18, 21, 26 and 32 years were considered to have 'ever been a smoker'.

Asthma: Those who presented with current asthma at any of the assessments at ages 9, 11, 13, 15, 18, 21, 26 and 32 years were considered to have 'ever had asthma'. Further details on the assessment of asthma are available in Sears *et al.* (2002).

As described above, we based diagnoses for each outcome on a cumulative count of cases, each of which was ascertained in a past-year assessment. Using this prospective approach, cases are not under-counted due to failure to recall criterion symptoms from years past, as occurs in retrospective surveys (Simon & VonKorff, 1995). Moreover, we have shown that there is very little case under-counting as a result of the gaps between past-year assessments. For example, only eight cohort members who had received mental-health services in the years between assessments had not been diagnosed by the study's repeated psychiatric interviews (Moffitt *et al.* 2007). These eight cases had received services for either depression or anxiety; no cases of psychosis, suicide or substance abuse were missed.

Statistical methods

For each disorder, associations between family history and diagnosis of disorder was assessed by conducting four logistic regression models, one with each family history score as a predictor (Table 1). The binary outcome in each model was the proband's disorder. Each model controlled for sex. So that predictive validity could be compared across family history scores, each score was z -standardized (mean=0 and standard deviation=1).

To compare the predictive validity of the dichotomous score *versus* the three density scores, we conducted the following logistic regression model for each disorder outcome:

$$Y_i = \beta_0 + \beta_F X_{ij} + \beta_D X_j + \beta_{FD} (X_{ij} \times X_j), \quad (1)$$

where Y_i represents the disorder status of proband i (0=non-disordered, 1=disordered), X_{ij} represents the family history score for proband i and family history score j (z -standardized as described above) and X_j represents the family history score method j (0=dichotomous, 1=density). The coefficient of interest here is β_{FD} , which tests whether the association between family history and disorder varies according to whether the method used to estimate family history is dichotomous or takes account of density of disorder (Table 1; statistical comparisons). As there are four family history scores (one dichotomous and three density), this model contains four observations per proband.

To compare the predictive validity of the three density scores, we restricted analyses to those three scores and conducted a second logistic regression model for each disorder outcome. We chose the simplest, observed score as the reference category when comparing family history scores. Thus, we modelled:

$$Y_i = \beta_0 + \beta_F X_{ij} + \beta_P X_P + \beta_{FP} (X_{ij} \times X_P) + \beta_R X_R + \beta_{FR} (X_{ij} \times X_R), \quad (2)$$

where the Y_i and X_{ij} terms are the same as in model (1), X_P represents the effect of the proportion score relative to the observed score (1=proportion score, 0=other scores), and X_R represents the effect of Reed's score relative to the observed score (1=Reed's score, 0=other score). The coefficients of interest here are β_{FP} , which tests whether the association between family history and disorder is different for the proportion *versus* the observed score (Table 1; statistical comparisons), and β_{FR} , which tests whether the association between family history and disorder is different for Reed's score *versus* the observed score (Table 1; statistical comparisons). As there are three family history scores, this model contains three observations per proband.

Because models (1) and (2) involve multiple observations per subject, standard error estimates were adjusted based on the sandwich or Huber–White variance estimator (Williams, 2000) to account for the lack of independence in the data. Both models (1) and (2) controlled for sex.

All analyses were conducted using STATA 9.1 (StataCorp, 2005).

Results

Table 1 shows the association between family history and disorder outcome for the four family history scores, and statistical comparisons of the associations produced by these four scores. The table shows that there were significant associations between family history and proband disorder outcome for all psychiatric disorders, as well as for smoking and asthma. These results held across the four family history scores, with odds of disorder associated

with a 1 standard deviation increase in family history score ranging from 1.2 (asthma) to 1.8 (conduct disorder).

The statistical comparisons of Table 1 reveal two main findings. First, there was some evidence that density scores had greater predictive validity than the dichotomous score: the density scores performed slightly better for all disorders, and performed significantly better for conduct disorder, alcohol dependence and smoking.

Second, among density scores, there was evidence that the proportion score and Reed's score were more predictive than the observed score (i.e. the observed number of family members with a positive history of disorder) for one outcome only: smoking. Moreover, compared with the observed score, Reed's score was actually less predictive of disorder for suicide attempt. We were surprised that the proportion score and Reed's score performed no better than the observed score for psychiatric disorders and asthma, because it suggests that the risk conveyed by having a number of family members with disorder is the same regardless of family size and family demographic structure. As smoking was far more prevalent among family members (prevalence=59.6%) than the other disorders (prevalence ranged from 3.3% for suicide attempt to 25.8% for major depressive episode), we hypothesized that taking account of family size and family demographic structure may be important for prevalent disorders only. To test this, we plotted the odds ratios assessing the difference between the proportion score and the observed score against the prevalence for each disorder (Fig. 2a) and the odds ratios assessing the difference between Reed's score and the observed score against the prevalence for each disorder (Fig. 2b). In support of our hypothesis, there was a strong positive correlation between the extent to which the proportion score was an improvement on the observed score and prevalence of disorder in family members ($r=0.93$, Fig. 2a). Similarly, there was a strong positive correlation between the extent to which Reed's score was an improvement on the observed score and prevalence of disorder in family members ($r=0.91$, Fig. 2b). Correlations were still strong if smoking, which is likely to exert a strong effect on the magnitude of the correlations, was excluded from analyses ($r=0.65$ and $r=0.58$, respectively).

Discussion

This study described the methods of the DFHHS and estimated associations between family history and disorder using four different family history scores. We found significant associations for all disorders across all family history scores, with odds of disorder associated with a 1 standard deviation increase in family history score ranging from 1.2 (asthma) to 1.8 (conduct disorder).

Are family history associations stronger for density versus dichotomous scores?

For all disorders, the measures we tested that took account of density of disorder (i.e. the number of family members with disorder) performed slightly better than a measure that dichotomized probands based on the presence or absence of disorder in at least one first-degree relative. Taking account of density of disorder is likely to be most beneficial for alcohol dependence, conduct disorder and smoking, for which the density measures produced family history associations that were significantly stronger than the family history associations produced by the dichotomous method.

Does taking account of family size and family demographic structure produce stronger family history associations?

The extent to which taking account of family size produced stronger family history associations was determined in part by the prevalence of disorder in family members. Thus,

for most disorders, we found very little difference between the density scores we computed based on a count of disordered family members ('observed') and the density scores we computed that took account of family size ('proportion' and 'Reed's score'). However, for smoking, a very prevalent phenotype among family members, both the proportion score and Reed's score produced stronger family history associations. Conversely, for suicide attempt, a rare phenotype among family members, Reed's score actually produced weaker family history associations. A possible explanation for this pattern of results is that an increase in family size is more likely to be associated with an increase in affected family members for prevalent than for non-prevalent disorders. Thus the need to take account of family size is likely to increase with increasing prevalence of disorder.

Taking account of family demographic structure did not appear to convey any benefits over and above those conveyed by taking account of family size.

Do our findings using psychiatric disorders generalize to non-psychiatric disorders?

Our finding that density scores tend to produce stronger family history associations than dichotomous scores generalized to the two non-psychiatric outcomes we tested: smoking and asthma. Also, our finding that the best method to assess family history depends on the prevalence of disorder appeared to apply to smoking and asthma as well as to psychiatric disorders. More work is needed to determine whether this phenomenon is universal across health domains.

Strengths

The present study has a number of strengths. First, we obtained family history reports from multiple informants. This has been shown to help avoid under-reporting, a common problem of the family history method (Kosten *et al.* 1992; Weissman *et al.* 2000; Hardt & Franke, 2007). Moreover, by collecting reports from multiple informants we avoided the biases associated with estimating family history solely from proband reports. That is, because informants with a psychiatric history of disorder tend to over-report that same disorder in family members (Kendler *et al.* 1991; Chapman *et al.* 1994; Roy *et al.* 1996; Heun *et al.* 1997; Coelho *et al.* 2006), relying solely on probands as informants would result in artefactual differences between disordered and non-disordered probands in terms of their family history.

Second, we were able to take advantage of our longitudinal design to prospectively assess disorder in the probands at a number of ages, and derived diagnoses by combining data across ages. In this way we were able to assess the association between family history and disorder using a measure of disorder that is not prone to biases associated with retrospective recall (Simon & Vonkorff, 1995).

Third, our findings were derived from a representative, population-based sample, so are likely to be applicable to population-based uses of family history (e.g. whole genome scans, public health screening programmes).

Fourth, we compared results across a number of psychiatric disorders as well as two non-psychiatric outcomes, so were able to test the generality of our findings. Moreover, studying a number of disorders at the one time enabled us to uncover patterns across disorders, e.g. the correlation between prevalence of disorder and the extent to which taking account of family size produced stronger family history associations. This correlation would not have been apparent if we had limited our analyses to just one or two disorders.

Limitations

Our findings should be interpreted in the context of the following limitations. First, our sample of probands consists of a primarily white birth cohort born in New Zealand in 1972–1973. Although our findings regarding the familiarity of psychiatric disorder are consistent with the literature (Kendler *et al.* 1997; Miles *et al.* 1998; Sullivan *et al.* 2000; Bandelow *et al.* 2002, 2004; Byrne *et al.* 2002; Klein *et al.* 2003; Qin *et al.* 2005; Newman & Bland, 2006; Coelho *et al.* 2007), our new findings regarding the relative merits of density and dichotomous scores need to be tested in other settings and with other ethnic groups.

Second, although we did our best to ensure the validity of family history reports (e.g. by using a validated instrument and by obtaining reports from multiple informants), the validity of the family history method is known to be inferior to directly interviewing every relative (Orvaschel *et al.* 1982; Andreason *et al.* 1986; Rice *et al.* 1995; Roy *et al.* 1996; Davies *et al.* 1997; Vandeleur *et al.* 2008). However, we chose to use the family history method because it is an inexpensive way to collect family history data on all relatives in a family. Also, it is not subject to the ascertainment bias associated with direct interviews, whereby those relatives able to be interviewed may have less psychopathology than those relatives unable to be interviewed due to death, estrangement or unwillingness to take part (Davies *et al.* 1997).

Third, when calculating Reed's score to take account of family demographic structure, we calculated expected rates based on prevalence of disorder rather than incidence of disorder. We took this approach because, to our knowledge, there are no published sex- and age-specific incidence rates for psychiatric disorders for the population we investigated. It is possible that this approach compromised our ability to assess the merits of Reed's score. However, studies of cardiovascular disease that have used incidence rates to calculate Reed's score have reached the same conclusion as us: that it adds little or no benefit over and above a proportion score (Silberberg *et al.* 1999; Murad *et al.* 2007).

Implications for family history researchers

With these limitations in mind, implications of our findings can be noted. First, for conduct disorder, alcohol dependence and smoking, researchers would benefit from assessing family history using density scores. Second, for other disorders, in particular depression, anxiety, suicidality and schizophrenia, family history can be adequately assessed using dichotomous scores. Third, the choice of density score depends upon the prevalence of disorder in family members. For disorders with very low prevalence (e.g. suicide), a simple count of family members with disorder should be preferred. For disorders with low to moderate prevalence, either a count score or a score calculated as the proportion of family members with disorder will suffice. For disorders with high prevalence (e.g. smoking), a proportion score should be preferred.

The use of family history data is vital for many areas of psychiatry. However, family history will only be as good as the scores that are used to measure it, so getting this right is important. We hope this comparative analysis will help researchers decide the best family history score to use.

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References

- Andreasen NC, Rice J, Endicott J, Reich T, Coryell W. The family history approach to diagnosis. *Archives of General Psychiatry*. 1986; 43:421–428. [PubMed: 3964020]
- APA. *Diagnostic and Statistical Manual of Mental Disorders*. 3. American Psychiatric Association; Washington, DC: 1980.
- APA. *Diagnostic and Statistical Manual of Mental Disorders*. 3. American Psychiatric Association; Washington, DC: 1987. revised
- APA. *Diagnostic and Statistical Manual of Mental Disorders*. 4. American Psychiatric Association; Washington, DC: 1994.
- Bandelow B, Späth C, Tichauer GA, Broocks A, Hajak G, Rütger E. Early traumatic life events, parental attitudes, family history, and birth risk factors in patients with panic disorder. *Comprehensive Psychiatry*. 2002; 43:269–278. [PubMed: 12107864]
- Bandelow B, Torrente AC, Wedekind D, Broocks A, Hajak G, Rütger E. Early traumatic life events, parental rearing styles, family history of mental disorders, and birth risk factors in patients with social anxiety disorder. *European Archives of Psychiatry and Clinical Neuroscience*. 2004; 254:397–405. [PubMed: 15538600]
- Byrne M, Agerbo E, Mortensen PB. Family history of psychiatric disorders and age of first contact in schizophrenia: an epidemiology study. *British Journal of Psychiatry*. 2002; 181 (Suppl 43):s19–s25.
- Chapman TF, Mannuzza S, Klein DF, Fyer AJ. Effects of informant mental disorder on psychiatric family history data. *American Journal of Psychiatry*. 1994; 151:574–579. [PubMed: 8147456]
- Chen X, Wang X, Hossain S, O'Neill FA, Walsh D, van den Oord E, Fanous A, Kendler KS. Interleukin 3 and schizophrenia: the impact of sex and family history. *Molecular Psychiatry*. 2007; 12:273–282. [PubMed: 17179997]
- Coelho HF, Cooper PJ, Murray L. Impact of psychiatric disturbance on identifying psychiatric disorder in relatives: study of mothers and daughters. *British Journal of Psychiatry*. 2006; 188:288–289. [PubMed: 16507974]
- Coelho HF, Cooper PJ, Murray L. A family study of co-morbidity between generalized social phobia and generalized anxiety disorder in a non-clinic sample. *Journal of Affective Disorders*. 2007; 100:103–113. [PubMed: 17113155]
- Costello, A.; Edelbrock, C.; Kalas, R.; Kessler, M.; Klaric, S. *Diagnostic Interview Schedule for Children (DISC)*. National Institute of Mental Health; Rockville, MD: 1982.
- Davies NJ, Sham PC, Gilvarry C, Jones PB, Murray RM. Comparison of the family history with the family study method: report from the Camberwell Collaborative Psychosis Study. *American Journal of Medical Genetics, Part B: Neuropsychiatric Genetics*. 1997; 74:12–17.
- Hardt J, Franke P. Validity, reliability and objectivity of the family history method in psychiatry: a meta analysis. *European Psychiatry*. 2007; 22:49–58. [PubMed: 17188848]
- Heun R, Maier W, Müller H. Subject and informant variables affecting family history diagnoses of depression and dementia. *Psychiatry Research*. 1997; 71:175–180. [PubMed: 9271790]
- Higgins M, Province M, Heiss G, Eckfeldt J, Ellison RC, Folsom AR, Rao DC, Sprafka JM, Williams R. NHLBI Family Heart Study: objectives and design. *American Journal of Epidemiology*. 1996; 143:1219–1228. [PubMed: 8651220]
- Kendler KS, Davis KL, Kessler RC. The familial aggregation of common psychiatric and substance use disorders in the National Co-morbidity Survey: a family history study. *British Journal of Psychiatry*. 1997; 170:S41–S48.
- Kendler KS, Silberg JL, Neale MC, Kessler RC, Heath AC, Eaves LJ. The family history method: whose psychiatric history is measured? *American Journal of Psychiatry*. 1991; 148:1501–1504. [PubMed: 1928463]
- Kim-Cohen J, Caspi A, Moffitt TE, Harrington H, Milne BJ, Poulton R. Prior juvenile diagnoses in adults with mental disorder: developmental follow-back of a prospective-longitudinal cohort. *Archives of General Psychiatry*. 2003; 60:709–717. [PubMed: 12860775]
- Klein DN, Lewinsohn PM, Rohde P, Seeley JR, Shankman SA. Family study of co-morbidity between major depressive disorder and anxiety disorders. *Psychological Medicine*. 2003; 33:703–714. [PubMed: 12785472]

- Kosten TA, Anton SF, Rounsaville BJ. Ascertaining psychiatric diagnoses with the family history method in a substance abuse population. *Journal of Psychiatric Research*. 1992; 26:135–147. [PubMed: 1613680]
- Miles DR, Stallings MC, Young SE, Hewitt JK, Crowley TJ, Fulker DW. A family history and direct interview study of the familial aggregation of substance abuse: the adolescent substance abuse study. *Drug and Alcohol Dependence*. 1998; 49:105–114. [PubMed: 9543647]
- Moffitt, TE.; Caspi, A.; Rutter, M.; Silva, PA. *Sex Differences in Antisocial Behaviour: Conduct Disorder, Delinquency, and Violence in the Dunedin Longitudinal Study*. Cambridge University Press; Cambridge, UK: 2001.
- Moffitt TE, Harrington H, Caspi A, Kim-Cohen J, Goldberg D, Gregory AM, Poulton R. Depression and generalized anxiety disorder: cumulative and sequential co-morbidity in a birth cohort followed prospectively to age 32 years. *Archives of General Psychiatry*. 2007; 64:651–660. [PubMed: 17548747]
- Murad H, Kalter-Leibovici O, Chetrit A, Freedman LS. A statistical comparison of different family history scores. *Statistics in Medicine*. 2007; 26:2785–2798. [PubMed: 17133629]
- Newman SC, Bland RC. A population based family study of DSM-III generalized anxiety disorder. *Psychological Medicine*. 2006; 36:1275–1281. [PubMed: 16700965]
- Orvaschel H, Thompson WD, Belanger A, Prusoff BA, Kidd KK. Comparison of the family history method to direct interview: factors affecting the diagnosis of depression. *Journal of Affective Disorders*. 1982; 4:49–59. [PubMed: 6461687]
- Prescott CA, Sullivan PF, Myers JM, Patterson DG, Devitt M, Halberstadt LJ, Walsh D, Kendler KS. The Irish affected sib pair study of alcohol dependence: study methodology and validation of diagnosis by interview and family history. *Alcoholism: Clinical and Experimental Research*. 2005; 29:417–429.
- Qin P, Xu H, Laursen TM, Vestergaard M, Mortensen PB. Risk for schizophrenia and schizophrenia-like psychosis among patients with epilepsy: population based cohort study. *British Medical Journal*. 2005; 331:23. [PubMed: 15964859]
- Reed T, Wagener DK, Donahue RP, Kuller LH. Young adult cholesterol as a predictor of familial ischemic heart disease. *Preventive Medicine*. 1986; 15:292–303. [PubMed: 3749009]
- Rice JP, Reich T, Bucholz K, Neuman RJ, Fishman R, Rochberg N, Hesselbrock VM, Nurnberger JJ, Schuckit MA, Begleiter H. Comparison of direct interview and family history diagnoses of alcohol dependence. *Alcoholism: Clinical and Experimental Research*. 1995; 19:1018–1023.
- Robins LN, Helzer HE, Croughan J, Ratcliff KS. National Institute of Mental Health Diagnostic Interview Schedule: its history, characteristics, and validity. *Archives of General Psychiatry*. 1981; 38:381–389. [PubMed: 6260053]
- Robins, LN.; Cottler, L.; Bucholz, K.; Compton, W. *Diagnostic Interview Schedule for DSM-IV*. Washington University Press; St Louis, MO: 1995.
- Roy M-A, Walsh D, Kendler KS. Accuracies and inaccuracies of the family history method: a multivariate approach. *Acta Psychiatrica Scandinavica*. 1996; 93:224–234. [PubMed: 8712019]
- Sears MR, Greene JM, Willan AR, Taylor DR, Flannery EM, Cowan JO, Herbison GP, Poulton R. Long-term relationship between breastfeeding and development of atopy and asthma in children and young adults: a longitudinal study. *Lancet*. 2002; 360:901–907. [PubMed: 12354471]
- Selzer ML, Vinokur A, Van Rooijen L. A self-administered Short Michigan Alcoholism Screening Test (SMAST). *Journal of Studies on Alcohol*. 1975; 36:117–126. [PubMed: 238068]
- Silberberg JS, Fryer J, Wlodarczyk J, Robertson R, Dear K. Comparison of family history measures used to identify high risk of coronary heart disease. *Genetic Epidemiology*. 1999; 16:344–355. [PubMed: 10207716]
- Simon GE, VonKorff M. Recall of psychiatric history in cross-sectional surveys: implications for epidemiologic research. *Epidemiologic Reviews*. 1995; 17:221–227. [PubMed: 8521941]
- Skinner HA. The drug abuse screening test. *Addictive Behaviors*. 1982; 7:363–371. [PubMed: 7183189]
- Statacorp. *Stata Statistical Software: Release 9.1*. Stata Corporation; College Station, TX: 2005.
- Stoltenberg SF, Mudd SA, Blow FC, Hill EM. Evaluating measures of family history of alcoholism: density vs dichotomy. *Addiction*. 1998; 93:1511–1520. [PubMed: 9926555]

- Sullivan PF, Neale MC, Kendler KS. Genetic epidemiology of major depression: review and meta-analysis. *American Journal of Psychiatry*. 2000; 157:1552–1562. [PubMed: 11007705]
- Sullivan PF, Wells JE, Joyce PR, Bushnell JA, Mulder RT, Oakley-Browne MA. Family history of depression in clinic and community samples. *Journal of Affective Disorders*. 1996; 40:159–168. [PubMed: 8897115]
- Thompson D, Witte JS, Slattery M, Goldgar D. Increased power for case–control studies of single nucleotide polymorphisms through incorporation of family history and genetic constraints. *Genetic Epidemiology*. 2004; 27:215–224. [PubMed: 15389928]
- Thompson WD, Kidd JR, Weissman MM. A procedure for the efficient collection and processing of pedigree data suitable for genetic analysis. *Journal of Psychiatric Research*. 1980; 15:291–303. [PubMed: 551167]
- Turner WM, Cutter HSG, Worobec TG, O'Farrell TJ, Bayog RD, Tsuang MT. Family history models of alcoholism: age of onset, consequences and dependence. *Journal of Studies on Alcohol*. 1993; 54:164–171. [PubMed: 8459710]
- Vandeleur CL, Rothen S, Jeanpretre N, Lustenberger Y, Gamma F, Ayer E, Ferrero F, Fleischmann A, Besson J, Sisbane F, Preisig M. Inter-informant agreement and prevalence estimates for substance use disorders: direct interview *versus* family history method. *Drug and Alcohol Dependence*. 2008; 92:9–19. [PubMed: 17643870]
- Weissman MM, Wickramaratne P, Adams P, Wolk S, Verdelli H, Olfson M. Brief screening for family psychiatric history: the family history screen. *Archives of General Psychiatry*. 2000; 57:675–682. [PubMed: 10891038]
- Williams RL. A note on robust variance estimation for cluster-correlated data. *Biometrics*. 2000; 56:645–646. [PubMed: 10877330]
- Yoon PW, Scheuner MT, Peterson-Oehlke KL, Gwinn M, Faucett A, Khoury MJ. Can family history be used as a tool for public health and preventive medicine? *Genetics in Medicine*. 2002; 4:304–310. [PubMed: 12172397]

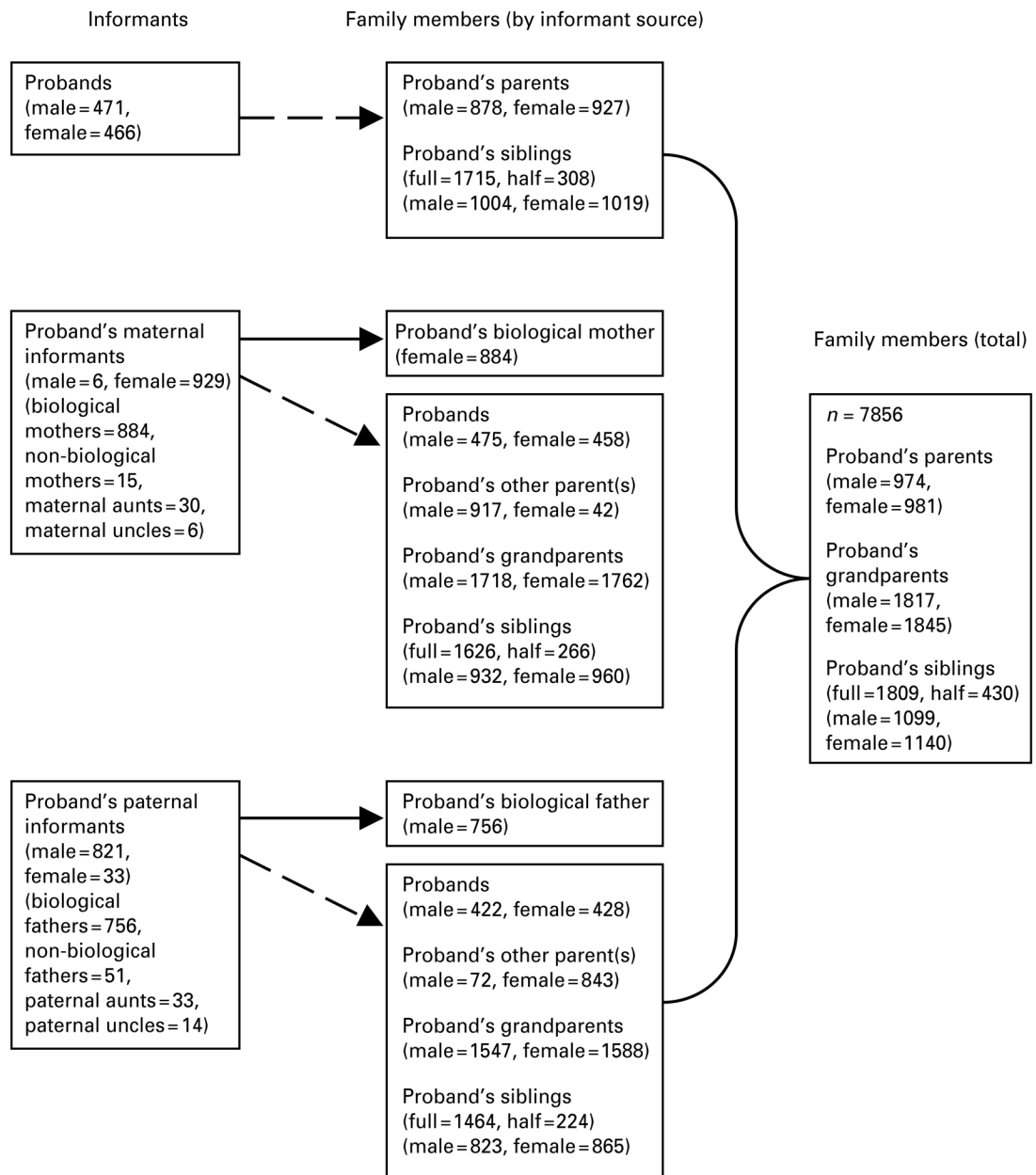


Fig. 1. Informants and family members from the Dunedin Family Health History Study. →, Information provided by self-report ; ->, information provided by informant-report.

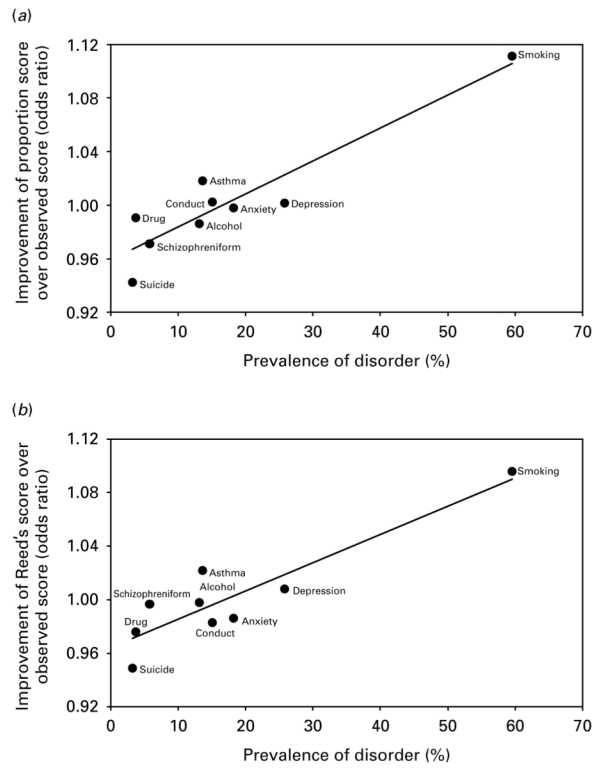


Fig. 2. Scatter-plots of prevalence of psychiatric disorders, smoking and asthma in family members *versus* the odds ratio assessing the difference in estimates of family history associations between the proportion score and the observed score (a), and *versus* the odds ratio assessing the difference in estimates of family history associations between Reed's score (Reed *et al.* 1986) and the observed score (b). The solid lines in each plot are fitted regression lines ; Pearson's correlation coefficients are $r=0.93$ for (a) and $r=0.91$ for (b). Correlations were still strong if smoking, which is likely to exert a strong effect on the magnitude of the correlations, was excluded from analyses [$r=0.65$ for (a) and $r=0.58$ for (b)].

Table 1

Family history associations using four different family history scores and statistical comparison of the family history score methods

Disorder	Associations using different family history scores ^a				Statistical comparisons ^b			
	Density scores		Proportion	Reed's score ^c	Density scores v. dichotomous score	Proportion score v. observed score	Reed's score v. observed score	
	Dichotomous score	Observed						
Major depressive episode	1.40 *** (1.21-1.62)	1.42 *** (1.24-1.63)	1.42 *** (1.24-1.63)	1.43 *** (1.25-1.64)	1.02 (0.91-1.14)	1.00 (0.95-1.06)	1.01 (0.96-1.05)	
Anxiety	1.45 *** (1.27-1.66)	1.47 *** (1.29-1.69)	1.47 *** (1.29-1.68)	1.45 *** (1.27-1.66)	1.01 (0.92-1.11)	1.00 (0.95-1.05)	0.99 (0.95-1.02)	
Suicide attempt	1.31 ** (1.07-1.62)	1.37 ** (1.14-1.66)	1.29 ** (1.07-1.56)	1.30 ** (1.07-1.59)	1.01 (0.91-1.11)	0.94 (0.88-1.01)	0.95 * (0.90-1.00)	
Schizophreniform disorder	1.45 * (1.09-1.94)	1.49 ** (1.17-1.91)	1.45 ** (1.16-1.81)	1.49 ** (1.17-1.89)	1.01 (0.86-1.19)	0.97 (0.86-1.10)	1.00 (0.95-1.05)	
Conduct disorder	1.43 *** (1.22-1.68)	1.79 *** (1.54-2.08)	1.80 *** (1.55-2.09)	1.76 *** (1.51-2.05)	1.24 *** (1.12-1.38)	1.00 (0.95-1.06)	0.98 (0.93-1.04)	
Alcohol dependence	1.21 ** (1.05-1.40)	1.46 *** (1.27-1.69)	1.44 *** (1.25-1.66)	1.46 *** (1.27-1.68)	1.20 *** (1.11-1.30)	0.99 (0.93-1.04)	1.00 (0.96-1.04)	
Drug dependence	1.28 ** (1.09-1.50)	1.38 *** (1.19-1.60)	1.36 *** (1.17-1.58)	1.34 *** (1.16-1.56)	1.06 (0.97-1.16)	0.99 (0.95-1.03)	0.98 (0.92-1.04)	
Ever smoked	1.26 ** (1.07-1.47)	1.51 *** (1.32-1.72)	1.67 *** (1.46-1.92)	1.65 *** (1.44-1.89)	1.28 ** (1.08-1.53)	1.11 * (1.02-1.21)	1.09 * (1.01-1.18)	
Asthma	1.20 * (1.04-1.39)	1.29 *** (1.12-1.48)	1.31 *** (1.14-1.51)	1.32 *** (1.14-1.52)	1.09 (0.99-1.19)	1.02 (0.98-1.06)	1.02 (0.98-1.06)	

^a Sex-adjusted odds (95% confidence intervals) of the proband having the disorder associated with a 1 standard deviation change in the family history score for that disorder (family history scores have been z-standardized).

^b Statistical comparison of the family history score methods. Odds ratios (95% confidence intervals) for tests of whether the associations between family history and disorder differ for different family history score methods.

^c Reed *et al.* (1986).

* $p < 0.05$,

** $p < 0.01$,

*** $p < 0.001$.