Familial clustering of suicide risk: a total population study of 11.4 million individuals

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Background. Research suggests that suicidal behaviour is aggregated in families. However, due to methodological limitations, including small sample sizes, the strength and pattern of this aggregation remains uncertain.

Method. We examined the familial clustering of completed suicide in a Swedish total population sample. We linked the Cause of Death and Multi-Generation Registers and compared suicide rates among relatives of all 83 951 suicide decedents from 1952–2003 with those among relatives of population controls.

Results. Patterns of familial aggregation of suicide among relatives to suicide decedents suggested genetic influences on suicide risk; the risk among full siblings (odds ratio 3.1, 95% confidence interval 2.8–3.5, 50% genetic similarity) was higher than that for maternal half-siblings (1.7, 1.1–2.7, 25% genetic similarity), despite similar environmental exposure. Further, monozygotic twins (100% genetic similarity) had a higher risk than dizygotic twins (50% genetic similarity) and cousins (12.5% genetic similarity) had higher suicide risk than controls. Shared (familial) environmental influences were also indicated; siblings to suicide decedents had a higher risk than offspring (both 50% genetically identical but siblings having a more shared environment, 3.1, 2.8–3.5 v. 2.0, 1.9–2.2), and maternal half-siblings had a higher risk than paternal half-siblings (both 50% genetically identical but the former with a more shared environment). Although comparisons of twins and half-siblings had overlapping confidence intervals, they were supported by sensitivity analyses, also including suicide attempts.

Conclusions. Familial clustering of suicide is primarily influenced by genetic and also shared environmental factors. The family history of suicide should be considered when assessing suicide risk in clinical settings or designing and administering preventive interventions.

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Key words: Family studies, intergenerational transmission, nationwide registers, suicidal behaviour, suicide.

Introduction

Suicide and attempted suicide are leading causes of death and morbidity worldwide. According to projections by the World Health Organization, approximately 1.5 million people will die from suicide in 2020 (Bertolote & Fleischmann, 2005). Suicidal behaviour runs in families (Brent et al. 1996; Runeson, 1998; Brent & Mann, 2005) and the familial transmission appears, at least partly, to be independent of psychiatric disorder (Agerbo et al. 2002; Qin et al. 2002, 2003; Runeson & Åsberg, 2003; Kim et al. 2005; McGirr et al. 2009; Brezo et al. 2010).

Theoretically, familial aggregation of suicides could be explained by shared genes, shared environments or both. In fact, results from prior family, adoption, twin, molecular genetic, geographic and immigrant studies do suggest that familial clustering of suicidal acts has genetic as well as environmental causes (Turecki, 2001; Baldessarini & Hennen, 2004; Brent & Mann, 2005; Voracek & Loibl, 2007, 2008; Wasserman et al. 2007). However, due to underpowered studies and limitations of data available for study, the strength and patterns of genetic and environmental impact on familial aggregation of suicidal behaviour remain uncertain (Baldessarini & Hennen, 2004; Baud, 2005; Bondy et al. 2006; Brent & Melhem, 2008; Brezo et al. 2008, 2010; Currier & Mann, 2008).

Three large twin studies yielded estimates of the genetic contribution to risk of a suicide attempt

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individual at birth or immigration and used for health and welfare statistics (Ludvigsson et al. 2009). The Cause of Death Register, held by the National Board of Health and Welfare, includes all deaths among persons registered as residents of Sweden at the time of death. Established in 1952, the register covers >99% of all deaths from 1961 and onwards, including those occurring abroad (National Board of Health and Welfare, 2010). Each record contains the date of death and codes for causes of death, including definite and uncertain suicide, in accordance with ICD-6, -7, -8, -9 and -10. The National Inpatient Register (maintained by the National Board of Health and Welfare) covers all in-patient health care (including the few existing private hospitals). The register has complete national coverage for all psychiatric admissions from 1973 and onwards. Each record contains admission and discharge dates, diagnoses and codes for attempted suicide (method used and if definite or uncertain) according to ICD-8, -9 and -10. Diagnoses and codes for suicide attempts are determined and recorded by the physician at discharge. The Multi-Generation Register (maintained by Statistics Sweden) contains information about child–parent relationships for all children born in Sweden since 1932 and who were still alive in 1961. Swedish-born children and children who immigrated to Sweden with at least one parent and were granted citizenship before age 18 years are linked to their parents. On 31 December 2004, the register contained 7,969,645 index persons together with their biological and adoptive parents; in total, 11,384,649 individuals. From these child–parent dyads, it is possible to construct larger pedigrees, including relatives at increasing distances (genetic and environmental) from each index person.

From the Cause of Death Register, we identified all Swedish residents who died from a definite suicide (ICD-6 [1951–1957]: E970–9; ICD-7 [1958–1968]: E971.9–979.9; ICD-8 [1969–1986], ICD-9 [1987–1996]: E950–9; ICD10 [1997–]: X60–84) or an uncertain suicide (ICD-8, ICD-9: E980–9; ICD10: Y10–34) between 1 January 1952 and 31 December 2003 (n = 83,951). Uncertain suicides (not coded in ICD-6 and ICD-7) were included to avoid underestimation of suicide rates (Neeleman & Wessely, 1997). Further, we identified all cases of a definite or uncertain suicide attempt from the National Inpatient Register (ICD-8, -9 and -10, codes correspond to those for completed suicide) between 1 January 1973 and 31 December 2003.

**Statistical analyses**

We used a matched case–control design to estimate familial suicide risks for different classes of relatives and other close relationships. We studied relatives at
varying distances from probands who had committed suicide. For instance, for each suicide proband, we specified all possible case-sibling pairs consisting of the proband and each of his/her full siblings (who had or had not committed suicide). For each case-sibling pair, we randomly selected five control-sibling pairs matched to case pairs by gender and birth year. The controls were selected through risk set sampling; that is, individuals were eligible as controls if they were alive and living in Sweden at the time of the case person’s suicide, regardless of whether they later committed suicide. This matching procedure was used for the probands’ first-, second- and third-degree relatives, as well as monozygotic twins, spouses/unrelated partners and adopted children. First-degree relatives comprised parents, full siblings (including dizygotic twins) and children. Second-degree relatives were grandparents, uncles/aunts, half-siblings, nephews/nieces and grandchildren, while third-degree relatives consisted of cousins. To increase statistical power, suicidal behaviour (completed suicide or suicide attempt leading to hospital care) was used as a broader suicidal behaviour phenotype for classes of relatives with few pairs concordant for completed suicide.

We calculated odds ratios (OR) using conditional logistic regression in PROC PHREG in SAS version 9.2 (SAS Institute, USA). Since multiple probands were possible in each family constellation, we adjusted for the non-independence of probands by computing corrected (less narrow) confidence intervals (CI) with a robust sandwich estimator (covsandwich option in PHREG). A similar case–control method was previously applied to calculate familial risks of schizophrenia, bipolar disorder (Lichtenstein et al. 2009) and violent offending (Frisell et al. 2011). For twin analyses, asymmetric CI for proportions were calculated with PROC FREQ in SAS version 9.2 (SAS Institute).

### Results

This total population sample included almost 8.0 million individuals of known parentage, which, together with their parents, constituted 11.4 million unique individuals. A total of 83,951 individuals committed suicide during the study period. Table 1 shows the relative risks of suicide in biological and non-biological relatives of probands who died by suicide compared to risks in relatives of matched control individuals. First, risk patterns suggested genetic influences on suicide risk; the risk for full siblings (OR 3.1, 95% CI 2.8–3.5, 50% genetic similarity) was significantly higher than that for maternal half-siblings (OR 1.7, 95% CI 1.1–2.7, 25% genetic similarity).

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### Table 1. Relative risks of suicide in relatives of all probands who committed suicide in Sweden during 1952–2003 (n = 83,951) compared with relatives of matched controls, in a total population cohort of 11,384,649 individuals

<table>
<thead>
<tr>
<th>Relation to proband</th>
<th>Number of dyads</th>
<th>Concordant pairs</th>
<th>0</th>
<th>1</th>
<th>2</th>
<th>3</th>
<th>4</th>
<th>Matched OR (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>First-degree relatives</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sibling</td>
<td>10,451,878</td>
<td>486</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>3.1 (2.8–3.5)</td>
</tr>
<tr>
<td>Child</td>
<td>13,714,253</td>
<td>923</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>2.0 (1.9–2.2)</td>
</tr>
<tr>
<td>Adopted-away child</td>
<td>66,564</td>
<td>12</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>1.4 (0.7–2.6)</td>
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<tr>
<td>Second-degree relatives</td>
<td></td>
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<td></td>
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<td></td>
</tr>
<tr>
<td>Maternal half-sibling</td>
<td>1,121,888</td>
<td>34</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>1.7 (1.1–2.7)</td>
</tr>
<tr>
<td>Paternal half-sibling</td>
<td>1,482,456</td>
<td>34</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>1.2 (0.8–1.9)</td>
</tr>
<tr>
<td>Nephew or niece</td>
<td>11,247,095</td>
<td>154</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>1.6 (1.4–1.9)</td>
</tr>
<tr>
<td>Grandchild</td>
<td>12,698,233</td>
<td>247</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>1.3 (1.2–1.5)</td>
</tr>
<tr>
<td>Third-degree relatives</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
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</tr>
<tr>
<td>Cousin</td>
<td>20,542,261</td>
<td>134</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>1.5 (1.3–1.8)</td>
</tr>
<tr>
<td>Non-biological relatives</td>
<td></td>
<td></td>
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<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Spouse/unrelated partner</td>
<td>7,538,548</td>
<td>882</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>2.3 (2.2–2.5)</td>
</tr>
<tr>
<td>Adopted child</td>
<td>206,692</td>
<td>11</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>1.3 (0.7–2.5)</td>
</tr>
<tr>
<td>Twins</td>
<td></td>
<td></td>
<td>0</td>
<td>10</td>
<td>20</td>
<td>30</td>
<td>40</td>
<td>50</td>
</tr>
<tr>
<td>Twin (monozygotic)</td>
<td>14,018</td>
<td>4</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>15.4 (5.4–43.8)</td>
</tr>
<tr>
<td>Twin (dizygotic, same gender)</td>
<td>19,664</td>
<td>3</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>3.0 (1.0–9.7)</td>
</tr>
<tr>
<td>Twin (dizygotic, opposite gender)</td>
<td>21,814</td>
<td>4</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>5.2 (1.9–14.4)</td>
</tr>
</tbody>
</table>

Relative risks were calculated with conditional logistic regression by comparing suicide rates in each proband-relative dyad type with population control-relative dyads matched 1:5 by relative category, gender, birth year and time at risk. The graph depicts odds ratios (OR) and 95% confidence intervals (CI) for all dyads (●). Each individual in the population could appear multiple times in different categories (e.g. sibling, child, cousin, etc.) depending on family pedigree.

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Familial clustering of suicide risk

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Discussion

This is the first total population study providing robust estimates of familial suicide risk in relatives at varying genetic and environmental distance from each other; including half-siblings, grandchildren, cousins and spouses. Importantly, since this investigation was four times larger than any previous study, we had the power to identify shared environmental effects on the risk of completed suicide. In line with our two hypotheses, we found evidence for familial aggregation of completed suicide influenced primarily by genetic and also by shared environmental factors. For example, we found a twofold increased risk of suicide in children and a threefold risk in siblings of suicide probands, compared with corresponding relatives of controls. These risk estimates are somewhat lower than the average size reported among first-degree relatives (children and siblings) in a smaller Danish study of a rather comparable design (Qin et al. 2003). A few limitations should be noted. The study was based solely on register data, although with excellent coverage and validity >40 years. Second, longitudinal registers are subject to 'left censoring', or missing data before the date the register started. However, since we matched for birth year and time at risk, such losses would be similar for both case and control dyads and not affect the estimates of familial aggregation (i.e. the relative risks). Third, despite the large sample size, some results were probably non-significant due to insufficient statistical power to detect aggregation of uncommon events, such as completed suicides. Nevertheless, sensitivity analyses using a broader measure of suicidal behaviour including attempted suicides leading to hospitalization generally supported the main findings. Fourth, the matching procedure did not include socio-economic status. Fifth, familial risks were not easily adjusted for presence of mental disorder in relatives. Only data on hospitalization due to mental disorder were available. However, since mental disorders are often subclinical or treated in out-patient care and suicidality generally increases the probability of being hospitalized for a mental disorder, this could have induced bias.

The familial aggregation of completed suicide was influenced by substantial genetic and also shared risk were confirmed by maternal half-siblings having significantly higher suicide risk than paternal half-siblings (OR 1.5, 95% CI 1.3–1.7 v. OR 1.1, 95% CI 0.9–1.2). Finally, genetic and shared environmental effects were both statistically supported by adopted-away biological children and adopted (non-biological) children having higher risks than controls (OR 1.6, 95% CI 1.2–2.0 for both comparisons).
environmental factors. But what are the possible mechanisms? The shared genetic component could involve liability to impulsive aggression, a propensity to react with aggression or hostility when frustrated or provoked (Turecki, 2001; Brent & Mann, 2006; McGirr & Turecki, 2007; Brent & Melhem, 2008). This should also be reflected in the familial aggregation of violence directed towards others, such as violent crime (Frisell et al. 2011). Personality traits including neuroticism or neurodevelopmental vulnerabilities involving impaired working memory or executive functioning resulting in poor problem solving could also be involved (Baud, 2005; Brent & Melhem, 2008; Mann et al. 2009). Gene variants related, for example, to the serotonergic-, noradrenergic- and dopaminergic neurotransmitter systems appear to be associated with the risk of suicidal behaviour (Brezo et al. 2008; Currier & Mann, 2008; Ernst et al. 2009); interaction patterns (gene–gene, gene–development, gene–environment), however, are largely unknown (Rujescu et al. 2007; Brezo et al. 2008, 2010; Currier & Mann, 2008; Ernst et al. 2009; Roy et al. 2009; Wasserman et al. 2010; Fergusson et al. 2011). The genetic effect probably involves a large number of single genes or alleles, similar to what has been suggested for schizophrenia and bipolar disorder (Purcell et al. 2009). However, it is important to note that such genetic contributions do not rule out that well-designed, environment-based interventions might reduce suicide risk.

On the other hand, our findings also suggested shared environmental effects on familial clustering of completed suicide. This could involve social learning of inadequate handling or communication of frustration or other negative affects, including those following bereavement of a spouse or relative from suicide (Agerbo, 2005; Brent & Mann, 2006). Transmission of adverse rearing environments could also be involved in the aggregation of suicide among parents and their children (Dinwiddie et al. 2000; Roy, 2002; Brent & Melhem, 2008). Direct role modelling or imitation of self-destructive behaviours, for instance, between siblings, is also possible although previous studies suggest that this effect is unlikely to be substantial (Statham et al. 1998; Brent & Melhem, 2008; Burke et al. 2010).

Regarding future research, large epidemiological studies with good statistical power are clearly needed, since suicide is a rare event, even among relatives of suicide victims. Various strategies to gain more specific knowledge on how gene variants, development and environment are associated with suicidal behaviour have been proposed. These include: (a) refining the phenotype for suicidal behaviour; (b) considering different possible pathways to suicidal behaviour. Concerning the former, endophenotypes (intermediate phenotypes between genes and a more overt phenotype or outcome), such as certain personality traits or altered neurocognitive function, should be characterized for suicidal behaviour (Brezo et al. 2008; Mann et al. 2009). Concerning the latter, several possible aetiological pathways for suicidal behaviour have been suggested, for instance, direct effects from psychiatric disorders, effects of early adversity, gene–environment correlation (selection into adversity), interactions between culture and genes and causal loops from genes to environment and back again (Kendler, 2010). These two overlapping approaches, as well as other strategies, are likely to require further development of technology (e.g. neuroimaging and -physiology) for more fine-grained assessment of biological mechanisms (Wasserman et al. 2010).

To conclude, this study provided strong evidence on the familial aetiology of completed suicide, primarily caused by genetic but also by shared environmental factors. The results confirm the importance of considering the family history of suicide when assessing suicide risk in clinical practice or when designing and administering preventive interventions.

Acknowledgments
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Declaration of Interest
None.

References


Mittendorfer-Rutz E, Rasmussen F, Wasserman D (2004). Restricted fetal growth and adverse maternal psychosocial and socioeconomic conditions as risk factors for suicidal


Appendix. Relative risks of suicidal behaviour (completed suicide or suicide attempt leading to hospital care) in relatives of all probands who committed suicide in Sweden during 1952–2003 (n = 83 951) compared with relatives of matched controls, in a total population cohort of 11 384 649 individuals.

<table>
<thead>
<tr>
<th>Relation to proband</th>
<th>Number of dyads</th>
<th>Concordant pairs</th>
<th>Matched odds ratio (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>First-degree relatives</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Child</td>
<td>13 714 253</td>
<td>4689</td>
<td>2.1 (2.0–2.1)</td>
</tr>
<tr>
<td>Adopted-away child</td>
<td>66 564</td>
<td>87</td>
<td>1.6 (1.2–2.0)</td>
</tr>
<tr>
<td>Second-degree relatives</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Maternal half-sibling</td>
<td>1 121 888</td>
<td>246</td>
<td>1.5 (1.3–1.7)</td>
</tr>
<tr>
<td>Paternal half-sibling</td>
<td>1 482 456</td>
<td>219</td>
<td>1.1 (0.9–1.2)</td>
</tr>
<tr>
<td>Non-biological relatives</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Adopted child</td>
<td>206 692</td>
<td>75</td>
<td>1.6 (1.2–2.0)</td>
</tr>
<tr>
<td>Twins</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Twin (monozygotic)</td>
<td>14 018</td>
<td>12</td>
<td>12.3 (6.5–23.2)</td>
</tr>
<tr>
<td>Twin (dizygotic, same-gender)</td>
<td>19 664</td>
<td>7</td>
<td>2.3 (1.1–5.0)</td>
</tr>
</tbody>
</table>

CI, Confidence interval. Relative risks in this table were calculated by the same procedure as those in Table 1.