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Australian children of alcoholic female twins

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MARC Project 4

Australian Children of Alcoholic Female Twins
Background

• Although it has been widely embraced by the treatment community, and certainly has a great deal of intuitive appeal, it has been difficult to demonstrate empirically a (non-genetic) consequence of being reared by an alcoholic parent.

• One critical test for demonstrating an important environmental effect of being reared by an alcoholic parent is to compare the rates of adverse outcomes among the biological offspring of an alcoholic parent to the rates of adverse outcomes among the biological offspring of the unaffected monozygotic cotwin of the alcoholic parent.

• The major aim of this project is to determine whether being raised by an alcoholic parent, in particular an alcoholic mother, increases the risk of adverse outcomes in the offspring after controlling for genetic transmission, and to identify mediators and modifiers of risk-outcome relationships.
Limitations of Previous Research

- **Family studies** have demonstrated that offspring of alcoholic parents are at higher risk for adverse outcomes than offspring of nonalcoholic parents, but it is impossible to determine from such studies whether this is due to genetic or environmental transmission of risk.

- **Twin studies** have generally led to the conclusion that family environmental influences do not play a major role in the familial transmission of alcoholism risk. However, in the twin design the estimate of family environmental effects only includes those that are independent of genetic effects.

- **Adoption studies** have not consistently demonstrated that offspring of alcoholic adoptive parents are at higher risk for adverse outcomes than offspring of nonalcoholic adoptive parents. Adoption studies are ideal in theory but limited in practice due to the screening of adoptive parents, which results in a restriction in the range of environmental adversity to which adoptive offspring are exposed.

- There is a paucity of research focused on the risk of adverse outcomes for **offspring of alcoholic mothers**.
Above are pedigree diagrams of the three types of twin-families included in this study of offspring of twins (shaded circles represent female twins with a history of alcohol use disorder (alcohol dependence or alcohol abuse -- AUD): families with at least one monozygotic female twin with a history of AUD, families with at least one dizygotic female twin with a history of AUD, and monozygotic or dizygotic twin families in which both female twins are unaffected with AUDs (control families).
Above are hypothetical results of the risk of adverse outcomes among offspring of twins from different risk categories represented in the previous panel. Panel ‘A’ represents the risk to offspring when the familial transmission is solely due to genetic effects, panel ‘B’ represents the risk to offspring when the familial transmission is solely due to family environmental effects, and panel ‘C’ represents the risk to offspring when the familial transmission is largely due to genotype x family environmental effects. All panels assume statistical control for paternal psychopathology.
Data Collection

• Data collection for this project is being done at the Queensland Institute of Medical Research in Brisbane, Australia.
• Female twin pairs from the different risk categories have been identified from previous large twin interview surveys.
• Female twin pairs are administered structured psychiatric telephone interviews in which they report about themselves, their biological offspring ages 7-22, and the father of the offspring. Fathers of the offspring are administered structured psychiatric telephone interviews in which they report about themselves.
• All offspring ages 11 and older are interviewed.
• Follow-up interviews with offspring are conducted every two years for a maximum of four interviews over the entire course of the 10-year study.
Key Constructs Assessed

In addition to collecting information about alcohol use and alcohol use disorders among all participants, we are assessing constructs related to three hypothesized pathways of the genetic and environmental transmission of alcoholism risk:

- **Deviant socialization pathway**
  - impaired parenting, family disruption
  - deviant peers
  - academic failure, childhood ADHD, oppositional behavior, and conduct problems

- **Negative affect pathway**
  - childhood stressors (physical and sexual abuse, traumatic events)
  - personality trait of neuroticism
  - internalizing disorders (depression and anxiety)

- **Pharmacological vulnerability pathway**
  - initial sensitivity to alcohol
  - drinking motives, alcohol expectancies
We are currently in year 7 of this project. By the end of year 10, we expect the following sample sizes:

<table>
<thead>
<tr>
<th>Risk group</th>
<th>Mothers</th>
<th>Offspring</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 - Mother AUD</td>
<td>332</td>
<td>512</td>
</tr>
<tr>
<td>2 - Mother unaffected, MZ cotwin AUD</td>
<td>101</td>
<td>179</td>
</tr>
<tr>
<td>3 - Mother unaffected, DZ cotwin AUD</td>
<td>104</td>
<td>171</td>
</tr>
<tr>
<td>4 - Mother unaffected, cotwin unaffected</td>
<td>654</td>
<td>1023</td>
</tr>
<tr>
<td><strong>Total</strong></td>
<td><strong>1191</strong></td>
<td><strong>1879</strong></td>
</tr>
</tbody>
</table>
Preliminary Findings From Years 1-5:
Rates of ADHD among offspring from different risk groups

<table>
<thead>
<tr>
<th>Risk group</th>
<th>% with ADHD</th>
<th>Odds ratio</th>
</tr>
</thead>
<tbody>
<tr>
<td>1a – Mother alcohol dependent</td>
<td>10.1</td>
<td>2.48*</td>
</tr>
<tr>
<td>1b - Mother alcohol abuse</td>
<td>9.2</td>
<td>2.16*</td>
</tr>
<tr>
<td>2 - Mother unaffected, MZ cotwin AUD</td>
<td>11.9</td>
<td>3.04*</td>
</tr>
<tr>
<td>3 - Mother unaffected, DZ cotwin AUD</td>
<td>1.6</td>
<td>0.31</td>
</tr>
<tr>
<td>4 - Mother unaffected, cotwin unaffected</td>
<td>4.8</td>
<td>1.00</td>
</tr>
</tbody>
</table>

These results are most consistent with a genetic transmission explanation of the increased risk of ADHD among the offspring of alcoholic mothers.