Genetic and Familial Environmental Influences on the Risk for Drug Abuse

A National Swedish Adoption Study

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**Context:** Prior research suggests that drug abuse (DA) is strongly influenced by both genetic and familial environmental factors. No large-scale adoption study has previously attempted to verify and integrate these findings.

**Objective:** To determine how genetic and environmental factors contribute to the risk for DA.

**Design:** Follow-up in 9 public databases (1961-2009) of adopted children and their biological and adoptive relatives.

**Setting:** Sweden.

**Participants:** The study included 18 115 adopted children born between 1950 and 1993; 78 079 biological parents and siblings; and 51 208 adoptive parents and siblings.

**Main Outcome Measures:** Drug abuse recorded in medical, legal, or pharmacy registry records.

**Results:** Risk for DA was significantly elevated in the adopted offspring of biological parents with DA (odds ratio, 2.09; 95% CI, 1.66-2.62), in biological full and half siblings of adopted children with DA (odds ratio, 1.84; 95% CI, 1.28-2.64; and odds ratio, 1.41; 95% CI, 1.19-1.67, respectively), and in adoptive siblings of adopted children with DA (odds ratio, 1.95; 95% CI, 1.43-2.65). A genetic risk index (including biological parental or sibling history of DA, criminal activity, and psychiatric or alcohol problems) and an environmental risk index (including adoptive parental history of divorce, death, criminal activity, and alcohol problems, as well as an adoptive sibling history of DA and psychiatric or alcohol problems) both strongly predicted the risk for DA. Including both indices along with sex and age at adoption in a predictive model revealed a significant positive interaction between the genetic and environmental risk indices.

**Conclusions:** Drug abuse is an etiologically complex syndrome strongly influenced by a diverse set of genetic risk factors reflecting a specific liability to DA, by a vulnerability to other externalizing disorders, and by a range of environmental factors reflecting marital instability, as well as psychopathology and criminal behavior in the adoptive home. Adverse environmental effects on DA are more pathogenic in individuals with high levels of genetic risk. These results should be interpreted in the context of limitations of the diagnosis of DA from registries.

Arch Gen Psychiatry. 2012;69(7):690-697. Published online March 5, 2012. doi:10.1001/archgenpsychiatry.2011.2112
ture of the genetic and familial environmental risk factors for DA and the interaction between them.

METHODS

To assess DA among Swedish adopted children and their relatives, we linked comprehensive registry and health care data from multiple nationwide sources to form a DA database. This linking was based on the Swedish unique individual 10-digit personal identification number that is nearly 100% complete and assigned at birth for all Swedish residents for their lifetime. This number was replaced by a serial number to provide anonymity. Our database contained 9 sources:

1. The Swedish Hospital Discharge Register included all hospitalizations (including for DA) for all people in Sweden from 1964 to 2009. Every record has the main discharge diagnosis and 8 secondary diagnoses.
2. The Swedish Prescribed Drug Register included all prescriptions in Sweden picked up by patients from 2005 to 2009. It is complete as all prescriptions are registered at the National Board of Health and Welfare.
3. The Swedish Mortality Register contained all causes of death and time of death from 1961 to 2009.
5. The Total Population Registry included annual data on education and marital status from 1990 to 2009.
6. The Multi-Generation Register provided information on family relationships from 1932 to 2009 including all adoptions and adoptive and biological parents and siblings. Biological siblings reared with the adopted child were excluded.
7. The Outpatient Care Register included information from all outpatient clinics in Sweden from 2001 to 2009.
8. The Primary Health Care Register included outpatient care data on diagnoses and time for diagnoses from 2001 to 2007 for 1 million patients from Stockholm and middle Sweden.
9. The Swedish Crime Register included national complete data on all convictions including those for DA from 1973 to 2007.

This study was approved by the ethics committee of Lund University in Malmo, Sweden.

SAMPLE

The study population consisted of individuals born between 1950 and 1993 who had been adopted, with information available on both adoptive parents and at least 1 biological parent. Individuals adopted by biological relatives or by an adoptive parent living with a biological parent were excluded. Age at formal adoption was not available in national records until 1991. Therefore, we estimated age at first cohabitation with adoptive parents (AFCAP) from census data including individual addresses available every 5 years (eg, 1950, 1955, 1960, etc.) For an adopted child born in 1961 and living with adoptive parents in the 1965 census, AFCAP was calculated as 4 years, although it could have been from 0 to 4 years. Thus, AFCAP represents an upper limit of true age at adoption. We constructed 2 cohorts with maximal AFCAPs of 5 years (n=12,783) and 10 years (n=18,115), and we performed analyses in parallel. To maximize statistical power, we present results on the entire cohort.

OUTCOME VARIABLE

Drug abuse was operationalized in this study using a range of Swedish registries. We identified DA in the Swedish hospital discharge, mortality, primary care, and outpatient care registries with the following codes from the eighth, ninth, and tenth revisions of the International Classification of Disease: ICD-8 codes for drug dependence (304); ICD-9 codes for drug psychoses (292) and drug dependence (304); and ICD-10 codes for mental and behavioral disorders due to psychoactive substance use (F10-F19), except those due to alcohol (F10) or tobacco (F17).

Drug abuse was also identified in the Crime Register by codes 5011 and 5012, which reflect crimes related to DA. Crimes related only to alcohol abuse or to trafficking in or possession of drugs of abuse were excluded. Plea bargains in Sweden only occur with minor offenses. An arrest for serious offenses including drug-related crimes routinely leads to prosecution. The Crime Register includes only criminal convictions that occur in general courts in Sweden. Minor offenses that typically do not go to these courts such as traffic and parking violations were not included.

Drug abuse was identified in individuals in the Prescribed Drug Register who had retrieved (on average) more than 4 daily doses for 12 months of hypnotics and sedatives (Anatomical Therapeutic Chemical Classification System codes N05C and N05BA) or opioids (Anatomical Therapeutic Chemical Classification System code N02A). Patients with cancer were excluded.

The 820 unique cases of DA in our cohort came from the following registries: Discharge (n=527), Crime (n=313), Outpatient (n=264), Prescribed Drugs (n=118), and Primary Health Care (n=8). No unique cases of DA were identified through the Mortality Register. Table 1 shows the odds ratios (ORs) and 95% confidence intervals for the ascertainment of DA among these 5 sources. Odds ratios ranged from a low of 3.6 between the Prescription and Crime Registers to a high of 118.0 between the Discharge and Outpatient Registers.

Misidentification of paternity was expected to be rare in our sample. Since 1938, all children born in Sweden must have a registered father. A public declaration of paternity is required and if there is uncertainty, blood tests for traditional genotyping (or after 1993, DNA testing) are required.

GENETIC RISK SCORE

We used the following variables in biological parents and/or biological siblings, measured during the entire life course: DA, hospitalization owing to alcoholism (identified in the Swedish

<table>
<thead>
<tr>
<th></th>
<th>Hospital Discharge</th>
<th>Outpatient</th>
<th>Primary Health Care</th>
<th>Drug Prescription</th>
</tr>
</thead>
<tbody>
<tr>
<td>Crime</td>
<td>32.9 (32.2-33.4)</td>
<td>65.2 (63.9-66.5)</td>
<td>47.4 (41.8-53.7)</td>
<td>5.6 (5.3-5.9)</td>
</tr>
<tr>
<td>Hospital discharge</td>
<td>118.0 (115.7-120.4)</td>
<td>69.8 (61.8-78.7)</td>
<td>20.9 (20.2-21.7)</td>
<td>29.6 (28.5-30.8)</td>
</tr>
<tr>
<td>Outpatient</td>
<td>94.4 (83.5-108.6)</td>
<td></td>
<td>37.9 (30.9-46.4)</td>
<td></td>
</tr>
</tbody>
</table>

Table 1. Registration of Drug Abuse Between the 5 Registers Used in the Study
enings in the household to deviant peer networks, were consid-
erginal family dysfunction and possible exposure for the sib-
the time of adoption.
also considered the age of the adoptive mother
omic status, using the highest education of the adoptive mother
ished among adoptive siblings and were measured during their entire life: DA; hospitalization owing to alcoholism, psychiatric illness, or other medical problems; and criminality. We created a score weighted according to the number of siblings.

ENVIRONMENTAL AND GENETIC RISK SCORES

We performed a logistic regression on the entire sample of 18 115 adopted children and modeled DA as a function of numerous factors associated with an increased risk for DA. All variables linked with biological parents/siblings and associated with DA (P < .10) in univariate analyses were included in the genetic risk score. When 2 predictors were strongly correlated, 1 was excluded from the model. Finally, we obtained the predicted probabilities (ie, genetic risk scores) for each adopted child and categorized them into 10 groups by deciles and used these variables as continuous variables in the final analysis. The same procedure was performed for all variables linked with adoptive parents/siblings to create an environmental risk score.

STATISTICAL ANALYSIS

Our key dependent variable, DA, is dichotomous. We initially used logistic regression and modeled DA as a function of the genetic risk score, the environmental risk score, sex of the adopted child, and AFCAP. However, a key a priori goal of these analyses was to determine whether genetic and environmental risk factors interacted in the etiology of DA. We have previously argued that the scale of raw probabilities, rather than the logistic scale, is more appropriate for such analyses. Therefore, for our analyses of gene × environment interaction, we used SAS Proc GENMOD (SAS Institute) with the identity link and we specified the variance to be binomial. We specified the effects of the explanatory variables (and the interaction term) to be additive on the scale of probabilities. All P values are reported as 2-tailed.

RESULTS

GENERAL DESCRIPTION

The general characteristics of our study sample are outlined in Table 2. In the 18 115 adopted children, whose average (SD) age at last available information was 46.2
4.5% (95% CI, 4.2-4.8) compared with 2.9% in all of Sweden from the same birth years. Compared with their adoptive parents, the biological parents had higher rates of DA, hospitalization for psychiatric illness and alcoholism, criminal convictions, low education, and divorce. Similar differences were seen between the adoptive and biological full and half siblings of the adopted children. The mean AFCAP was 4.4 years.

We examined the prevalence of DA as defined in this study by decade of birth (ie, 1950-59, 1960-69, 1970-79, and 1980-1993). In the entire Swedish population, this varied slightly, with no monotonic trends, between 2.8% and 3.1%. However, among the adopted children, prevalence rates of DA increased consistently across these 4 cohorts: 4.1%, 4.6%, 5.0%, and 6.5%. Therefore, we controlled for year of birth in all subsequent analyses.

Table 3. Rates of Drug Abuse in Adopted Children of Biological Parents, Adoptive Offspring of Adoptive Parents, and Biological and Adoptive Siblings of Adopted Children With and Without Drug Abuse

<table>
<thead>
<tr>
<th>Relative</th>
<th>Rate of Drug Abuse in Relative When Proband Does Not Have Drug Abuse</th>
<th>Rate of Drug Abuse in Relative When Proband Has Drug Abuse</th>
</tr>
</thead>
<tbody>
<tr>
<td>Adopted away biological offspring</td>
<td>8.6 (7.0-10.4)</td>
<td>4.2 (4.0-4.6)</td>
</tr>
<tr>
<td>Adopted away biological offspring</td>
<td>9.9 (7.2-12.1)</td>
<td>4.4 (4.1-4.7)</td>
</tr>
<tr>
<td>Adopted away biological offspring</td>
<td>8.2 (6.3-10.5)</td>
<td>4.4 (4.1-4.7)</td>
</tr>
<tr>
<td>Biological offspring</td>
<td>6.8 (3.8-11.6)</td>
<td>4.5 (4.2-4.8)</td>
</tr>
<tr>
<td>Biological full sibling</td>
<td>10.7 (7.6-14.5)</td>
<td>6.1 (5.6-6.7)</td>
</tr>
<tr>
<td>Biological half sibling</td>
<td>7.4 (6.3-8.5)</td>
<td>5.3 (5.1-5.5)</td>
</tr>
<tr>
<td>Adoptive sibling</td>
<td>6.2 (4.5-8.1)</td>
<td>3.2 (3.0-3.5)</td>
</tr>
</tbody>
</table>

The risk for DA among children given for adoption by biological parents, at least 1 of whom had DA, was 8.6%, which was substantially and significantly elevated over that seen in children given for adoption when neither biological parent had DA (4.2%) (Table 3) (OR, 2.09; 95% CI, 1.66-2.62; \( \chi^2 = 39.4; P \leq .001 \)). Among adopted children, those who had none, 1, and both biological parents with DA had a prevalence of DA of 4.2%, 8.2%, and 11.9%, respectively (\( \chi^2 = 44.0; P \leq .001 \)). The risk for DA among adopted children of biological fathers and mothers with DA was 9.9% and 8.2%, respectively. The risk for DA was increased in both biological full siblings and biological half siblings of adopted children with DA (10.7% and 7.4%, respectively) vs without DA (6.1% and 5.3%, respectively) (OR, 1.84; 95% CI, 1.28-2.64; \( \chi^2 = 11.0; P \leq .001 \)); and OR, 1.41; 95% CI, 1.19-1.67; \( \chi^2 = 9.7; P \leq .001 \)), respectively.

The risk for DA in adopted children raised by 1 or more adoptive parents with vs without DA was not significantly increased (6.8% vs 4.5%; OR, 1.55; 95% CI, 0.86-2.80; \( \chi^2 = 2.13; P = .15 \)). However, adoptive siblings of adopted children with vs without DA had a significantly increased risk for DA (6.2% vs 3.2%; OR, 1.95; 95% CI, 1.43-2.65; \( \chi^2 = 13.9; P \leq .001 \)).

Predictors of Genetic and Environmental Risk for DA and Their Interrelationship

We created indices of genetic and environmental risks for DA from available characteristics of the adopted children’s biological parents and siblings and their adoptive parents and siblings. As seen in Table 2, genetic risk for DA in the adopted child was indexed by a range of features including biological parental low-educational attainment and divorce, and a parental or sibling history of DA, criminal activity, and treatment for psychiatric or alcohol problems. Environmental risk for DA was predicted by a diverse set of characteristics including an adoptive parental history of divorce, premature death, criminal activity, and hospitalization for medical or alcohol problems, and an adoptive sibling history of DA and hospitalization for psychiatric, alcohol, or medical problems.

Examined individually in logistic regression (Table 4), where the genetic and environmental risk indices were divided into 10 deciles, both risk scores were strongly predictive of DA: OR (per decile), 1.13 and 1.10, respectively. Thus, the ORs for DA between individuals at the lowest and highest decile of genetic and environmental risk equal 1.1310 or 3.39 and 1.1010 or 2.59, respectively. The correlation between risk scores was small (plus 0.11) but significant (\( P < .001 \)). Examined individually, DA in adopted children was also significantly predicted by male sex, younger AFCAP, and a later birth year.

Next, we examined the genetic and environmental indices jointly with sex, AFCAP, and birth year (Table 5). The predictive power of both indices declined slightly as would be expected given their low intercorrelation. Sex remained predictive but not AFCAP or birth year.

We then explored whether the sexes differed in their sensitivity to genetic or environmental risk factors. Adding these interactions one at a time to the model depicted in Table 4 showed no significant interaction between sex and environmental risk (\( P = .91 \)), but there was modest evidence that males were more sensitive to the impact of genetic risk factors for DA (\( P = .03 \)).

Finally, we analyzed these predictor variables on the scale of raw probabilities without and with an interaction between the genetic and environmental risk scores.
Table 4. Creation of Genetic and Environmental Risk Scores for Adopted Children

<table>
<thead>
<tr>
<th></th>
<th>Adoptive Relations (Environmental Risk Score)</th>
<th>Biological Relations (Genetic Risk Score)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Univariate Analyses</td>
<td>Multivariate Analysis</td>
</tr>
<tr>
<td>Parents</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Drug abuse</td>
<td>1.55 (0.86-2.80)</td>
<td>2.09 (1.66-2.63)(a)</td>
</tr>
<tr>
<td>Psychiatric disease</td>
<td>1.25 (0.86-1.80)</td>
<td>1.58 (1.34-1.85)(a)</td>
</tr>
<tr>
<td>Alcoholism</td>
<td>1.89 (1.09-3.29)(c)</td>
<td>1.76 (1.49-2.08)(a)</td>
</tr>
<tr>
<td>Convictions</td>
<td>1.42 (1.15-1.76)(b)</td>
<td>1.49 (1.30-1.72)(a)</td>
</tr>
<tr>
<td>Maternal age</td>
<td>0.97 (0.96-0.98)(c)</td>
<td>0.99 (0.98-1.00)(a)</td>
</tr>
<tr>
<td>Maternal divorce</td>
<td>1.51 (1.19-1.92)(b)</td>
<td>1.24 (1.08-1.43)(b)</td>
</tr>
<tr>
<td>Education (low vs high)</td>
<td>0.92 (0.80-1.06)</td>
<td>1.29 (1.08-1.54)(b)</td>
</tr>
<tr>
<td>Medical hospitalization</td>
<td>1.30 (1.13-1.50)(d)</td>
<td>1.17 (1.01-1.35)(c)</td>
</tr>
<tr>
<td>Death</td>
<td>1.29 (1.02-1.63)(c)</td>
<td>1.37 (1.08-1.73)(b)</td>
</tr>
<tr>
<td>Siblings</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Drug abuse</td>
<td>2.29 (1.61-3.27)(a)</td>
<td>1.76 (1.16-2.67)(b)</td>
</tr>
<tr>
<td>Psychiatric disease</td>
<td>1.57 (1.16-2.11)(b)</td>
<td>1.16 (0.83-1.62)(b)</td>
</tr>
<tr>
<td>Alcoholism</td>
<td>1.89 (1.26-2.84)(b)</td>
<td>1.27 (0.80-2.01)</td>
</tr>
<tr>
<td>Convictions</td>
<td>0.99 (0.85-1.14)(b)</td>
<td>1.21 (0.95-1.55)(a)</td>
</tr>
<tr>
<td>Medical hospitalization</td>
<td>1.25 (1.09-1.44)(b)</td>
<td>1.12 (0.97-1.30)(a)</td>
</tr>
</tbody>
</table>

\(a\) \(P < .001.\)
\(b\) \(P < .01.\)
\(c\) \(P < .05.\)
\(d\) \(P < .10.\)

Table 5. Prediction of Drug Abuse in Adopted Children From the Genetic and Environmental Risk Scores by Logistic Regression Analysis

<table>
<thead>
<tr>
<th></th>
<th>Odds Ratio (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Univariate Analysis</td>
</tr>
<tr>
<td>Genetic risk score</td>
<td>1.13 (1.10-1.16)(b)</td>
</tr>
<tr>
<td>Environmental risk score</td>
<td>1.10 (1.07-1.12)(b)</td>
</tr>
<tr>
<td>Sex of adopted child (male vs female)</td>
<td>1.45 (1.26-1.67)(b)</td>
</tr>
<tr>
<td>AFCAP from census data</td>
<td>0.96 (0.93-0.98)(c)</td>
</tr>
<tr>
<td>Birth year of adopted child</td>
<td>1.02 (1.01-1.02)(c)</td>
</tr>
</tbody>
</table>

Abbreviation: AFCAP, age at first cohabitation with adoptive parents.

\(a\) Multivariate model includes adopted children’s genetic and environmental risk scores, sex, AFCAP, and birth year.
\(b\) \(P < .001.\)
\(c\) \(P < .01.\)

Table 6. The interaction was significant, showing that the impact of an adverse environment on risk for DA was substantially greater in those with a high vs low genetic liability to DA (Figure).

From these analyses, 5 results are noteworthy. First, we replicated in a large national full-adoption design results from 4 large-sample general population twin studies performed in Australia and the United States\(7,9\) showing a strong contribution of genetic factors to the etiology of DA. While our study used registry data, diagnoses of DA in these twin studies were all based on personal interviews. Our results are also consistent with findings from a small-sample US adoption study that examined 197 adopted children. The interaction was significant, showing that the impact of an adverse environment on risk for DA was substantially greater in those with a high vs low genetic liability to DA (Figure).

COMMENT

From these analyses, 5 results are noteworthy. First, we replicated in a large national full-adoption design results from 4 large-sample general population twin studies performed in Australia and the United States\(7,9\) showing a strong contribution of genetic factors to the etiology of DA. While our study used registry data, diagnoses of DA in these twin studies were all based on personal interviews. Our results are also consistent with findings from a small-sample US adoption study that examined 197 adopted children. The interaction was significant, showing that the impact of an adverse environment on risk for DA was substantially greater in those with a high vs low genetic liability to DA (Figure).
parent-child bonds through death or divorce, alcohol problems in the adoptive parents or siblings, and crimi-
unal behavior and medical hospitalization in the adopt-
ive parents. We would speculate that the causal pro-
cesses here might include increased marital tension, poorer parent-child relationships, and reduced parental moni-
toring. Interestingly, risk for DA in adopted children was more strongly predicted by DA in the adoptive siblings than adoptive parents. Our results suggest that social in-
fluences (eg, peer deviance and drug availability) shared with adoptive siblings are more potent environmental risks for DA than direct psychological transmission of DA from parent to child.24 Evidence from this study for robust shared environmental effects on DA is at variance with results from some6,7,9 but not all5,8 adult twin studies ex-
amining lifetime drug abuse/dependence. Prior twin studies of drug use typically found strong shared environ-
mental effects in adolescence25,26 that largely disappeared in adulthood. However, our findings confirm prior re-
results from a US study that found significant resem-
lance in 246 adoptive adolescent same-sex sibling pairs.

Fifth, we found evidence for gene-environment in-
teraction in the etiology of DA. Adopted children at high genetic risk were more sensitive to the pathogenic ef-
facts of adverse family environments than those at low genetic risk. In other words, genetic effects on DA were less potent in low-risk than in high-risk environments. These results are consistent with prior twin studies sug-
gesting that genetic influences on psychoactive sub-
stance use in adolescence are enhanced in high-risk en-
vironments characterized by low parental monitoring,28 easy substance availability,29 and the presence of sub-
stance-using friends,30,31 as well as consistent with mol-
ecular genetic studies showing that risk variants may in-
teract positively with familial environmental adversity in the prediction of DA.32

**METHODOLOGICAL LIMITATIONS**

These results should be interpreted in the context of 3 po-
tentially important methodologic limitations. First, we
identified subjects with DA from medical, legal, and phar-
acy records. While this method has the important ad-
vantage of not requiring accurate respondent recall and

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**Table 6. Prediction of Drug Abuse Among Adopted Children From the Genetic and Environmental Risk Scores and Their Interaction on the Scale of Raw Probabilities**

<table>
<thead>
<tr>
<th>Genetic risk score</th>
<th>Multivariate Analysis Without an Interaction Term</th>
<th>Multivariate Analysis With an Interaction Term</th>
</tr>
</thead>
<tbody>
<tr>
<td>Environmental risk score</td>
<td>0.0049 (0.0038 to 0.0060) ( a )</td>
<td>0.0028 (0.0010 to 0.0048) ( b )</td>
</tr>
<tr>
<td>Sex adoptive child (male vs female)</td>
<td>0.0036 (0.0024 to 0.0048) ( a )</td>
<td>0.0015 (−0.0006 to 0.0036)</td>
</tr>
<tr>
<td>AFCAP from census data</td>
<td>0.0151 (0.0091 to 0.0212) ( a )</td>
<td>0.0152 (0.0091 to 0.0213) ( a )</td>
</tr>
<tr>
<td>Birth year of adopted child</td>
<td>−0.0003 (−0.0017 to 0.0010)</td>
<td>−0.0005 (−0.0018 to 0.0009)</td>
</tr>
<tr>
<td>Interaction term (genetic × environmental risk scores)</td>
<td>−0.0004 (−0.0008 to 0.0001)</td>
<td>−0.0004 (−0.0008 to 0.0001)</td>
</tr>
</tbody>
</table>

Abbreviation: AFCAP, age at first cohabitation with adoptive parents.

\( a P < .001. \)

\( b P < .05. \)
0.11) but the magnitude was too small to substantially influence our findings.

Third, bias can also arise in adoption studies from extensive contact between the adopted children and biological parents prior to adoption. We had to approximate AFCAP from census records available every 5 years. We know that at least 70.9% of adopted children were living with their adoptive parents by age 5 years. However, during the years of our study, adopted children were typically removed shortly after birth from the biological mother and placed in a special nursery home.³⁹ ⁴⁰ Often prior to formal adoption, the child would be placed for trial periods in foster homes. Thus, prior to placement in the adoptive home, adopted children in our sample were more likely to be in the special nursery or foster homes than with their biological parents.

We could assess this possible bias in 3 ways. First, if sustained contact with biological parents occurred and increased the risk for DA in the adopted child, then AFCAP should be significant and positively associated with DA. Instead, the correlation was negative. Second, delayed placement in the adoptive home could attenuate the impact of the adoptive environment. This would predict that AFCAP should negatively interact with the environmental risk score to predict DA. No such interaction was found (P = .88). Finally, we repeated our analyses in the subsample (n = 12,783) where AFCAP was 5 years or less. The broad pattern of results was unchanged.

CONCLUSIONS

To our knowledge, this is the first large-scale, comprehensive adoption study of DA that has confirmed and extended prior studies showing that DA is etiologically complex with important genetic and shared-environmental influences. Both the genetic and the familial environmental influences on DA are themselves multifaceted. Risk for DA in adopted children is increased by a history in biological parents and siblings not only of DA but also of alcoholism, major psychiatric illness, and criminal convictions. Risk for DA in adopted children is increased by disruption in the adoptive parent–adopted child bond by death or divorce but also by a range of indices of a disturbed adoptive home environment and deviant peer influences such as parental alcoholism and sibling drug abuse, respectively. Finally, the genetic factors that influence liability to DA act not only by directly increasing the risk for illness but also by increasing the pathogenic effects of adverse environmental experiences.

Submitted for Publication: August 1, 2011; final revision received October 13, 2011; accepted December 7, 2011.

Published Online: March 5, 2012. doi:10.1001/archgenspsychiatry.2011.2112

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Financial Disclosure: None reported.

Funding/Support: This study was funded by grant RO1 DA030005 from the National Institute of Drug Abuse, grants 2008-3110 and 2008-2638 from the Swedish Research Council, and a project grant from ALF, Lund, Sweden.

Role of the Sponsors: The sponsors were not involved in the design and conduct of the study; collection, management, analysis, and interpretation of the data; and preparation, review or approval of the manuscript.

REFERENCES


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