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Stability in symptoms of anxiety and depression as a function of genotype and environment: a longitudinal twin study from ages 3 to 63 years

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Background. The influence of genetic factors on major depressive disorder is lower than on other psychiatric disorders. Heritability estimates mainly derive from cross-sectional studies, and knowledge on the longitudinal aetiology of symptoms of anxiety and depression (SxAnxDep) across the lifespan is limited. We aimed to assess phenotypic, genetic and environmental stability in SxAnxDep between ages 3 and 63 years.

Method. We used a cohort-sequential design combining data from 49 524 twins followed from birth to age ≥ 20 years, and from adolescence into adulthood. SxAnxDep were assessed repeatedly with a maximum of eight assessments over a 25-year period. Data were ordered in 30 age groups and analysed with longitudinal genetic models.

Results. Over age, there was a significant increase during adolescence in mean scores with sex differences (women > men) emerging. Heritability was high in childhood and decreased to 30–40% during adulthood. This decrease in heritability was due to an increase in environmental variance. Phenotypic stability was moderate in children (correlations across ages ~ 0.5) and high in adolescents ($r = 0.6$), young adults ($r = 0.7$), and adults ($r = 0.8$). Longitudinal stability was mostly attributable to genetic factors. During childhood and adolescence there was also significant genetic innovation, which was absent in adults. Environmental effects contributed to short-term stability.

Conclusions. The substantial stability in SxAnxDep is mainly due to genetic effects. The importance of environmental effects increases with age and explains the relatively low heritability of depression in adults. The environmental effects are transient, but the contribution to stability increases with age.

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Key words: Anxiety symptoms, depression symptoms, longitudinal genetic analysis.

Introduction

Insight into the course and the aetiology of variation in trajectories of psychopathology from childhood into adolescence and adulthood is required to address questions concerning origins and prognosis of psychopathology. In this paper we aim to unravel the causes of variation in trajectories of symptoms of anxiety and

depression (SxAnxDep) between ages 3 and 63 years, and specifically to assess the extent to which such variation is caused by genetic factors. A large prospective cohort study would be the optimal design for identifying the importance of genetic aetiological factors, but would require following the same subjects for ≥ 50 years. Here we made use of the long-term data collection in nearly 50 000 twins from The Netherlands Twin Register (NTR) over the past 25 years, and analysed data on SxAnxDep reordered according to a cohort-sequential design.

Stability in symptoms and diagnoses of anxiety and depression is evident throughout the lifespan; stability is lowest between childhood and adolescence, and

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increases from adolescence into adulthood (Hofstra *et al.* 2000; Richards, 2011; Beesdo-Baum & Knappe, 2012). During adolescence, there is a rise in the prevalence of anxious and depressive symptoms and diagnoses, especially in women (with the exception of separation anxiety disorder and specific phobia), and there is a marked continuity of symptoms into adulthood (Costello *et al.* 2003; Beesdo-Baum & Knappe, 2012). When considering stability in longitudinal studies, outcomes depend in part on how anxiety and depression are measured. Greater stability of depression has been observed in continuous measures (e.g. symptom counts) than in binary measures indicating that individuals, who no longer satisfy the criteria for a diagnosis, may still have residual symptoms (Ormel *et al.* 1993).

A case has been made that studies of the genetic and environmental contributions to individual differences in health and disease are important to the understanding of illness (Collins, 2004). In addition, such studies inform molecular genetics studies (van Dongen *et al.* 2012). Knowledge regarding the contributions of genetic and environmental factors to the long-term course of SxAnxDep is still lacking, as most studies using genetically informative subjects have been limited to cross-sectional or short-term follow-up analyses. Moreover, in these short-term follow-up analyses, the age range of the participants at the start of a longitudinal study is often greater than the duration of the follow-up. As a consequence, the age ranges of the participants at the different assessments overlap and age-specific explanations of variation over time might be missed.

The empirical evidence suggests that between ages 3 and 12 years genetic effects are important, and relatively stable, while adolescence is characterized by genetic innovations, i.e. the emergence of novel genetic effects. In an overview of cross-sectional genetic studies of young and adolescent twins, heritability estimates for anxiety, depression and internalizing symptoms range from 0% to 74%, but the majority is over 30% (Rice, 2009). The great variability over studies is attributed to rater and age effects; in most studies heritability is higher for parental reports than for self-reports, and heritability is higher at adolescence than in childhood (Rice & Thapar, 2009). Longitudinal studies of children in The Netherlands and in the UK showed that stability in SxAnxDep between ages 3 and 12 years is mainly attributable to stable genetic effects (Boomsma *et al.* 2008; Trzaskowski *et al.* 2012), although genetic innovation was also evident in this age range. The period of transition from childhood to adolescence was investigated in Swedish twins with SxAnxDep, assessed repeatedly between ages 8 and 20 years (Kendler *et al.* 2008a–c). Stability was partly

explained by genetic factors. However, the influence of genetic innovation was large, and the contribution of genetic factors expressed during childhood declined in adolescence. Thus, genetic effects on SxAnxDep were developmentally dynamic from middle childhood to young adulthood.

In adulthood, heritability of anxiety disorders and major depression was estimated around 40% (Sullivan *et al.* 2000; Middeldorp *et al.* 2005; Kendler *et al.* 2006). For depression in middle age (50–70 years) heritability estimates ranged from 20% to 50% (Franz *et al.* 2011). In one longitudinal twin study, participants aged 20–70 years rated themselves on SxAnxDep two or three times with an interval of 10 or 20 years. Genetic effects on symptom scores showed a large degree of stability, with some evidence in females for new genetic effects on anxiety and depression in mid-life and later-life, respectively (Gillespie *et al.* 2004).

The role of environmental effects on stability has been found to be smaller than the role of genetic factors. Environmental influences shared by members of the same family, often referred to as the common environment, contributed to stability in SxAnxDep during childhood (Boomsma *et al.* 2008; Trzaskowski *et al.* 2012), but effects were small and waned from around 15% explained variance in childhood to zero in adolescence and adulthood (Rapee *et al.* 2009). Environmental factors that are not shared by family members, referred to as unique environment, mainly seemed to have short-term effects on SxAnxDep. The impact of life events, for example, on the risk for major depression has been shown to last for 1–3 months (Kendler *et al.* 1998). However, two twin studies found enduring unique environmental effects from adolescence into adulthood and beyond (Gillespie *et al.* 2004; Kendler *et al.* 2008c). This was confirmed in a meta-analysis of longitudinally assessed SxAnxDep in eight samples of monozygotic twins, spanning an age range of 10–66 years (Kendler *et al.* 2011). Within-pair differences between MZ twins in SxAnxDep increased from childhood into late adulthood. By middle adulthood environmental factors contributed substantially to stable and predictable individual differences in SxAnxDep.

The aim of the present study was to gain insight into the genetic architecture of SxAnxDep across the life-span by analysing SxAnxDep assessed with a standardized instrument in twins aged 3–63 years. Data collection spanned a period of 25 years, and entry into the study was at different ages, running from birth to old age. The data were reorganized according to a cohort-sequential longitudinal design, covering the entire age range from age 3 to age 63 years with a maximum follow-up time of 25 years.

Method

Subjects

Longitudinal survey data were collected in twins registered with the Netherlands Twin Register (NTR), which includes the Young NTR (YNTR; van Beijsterveldt *et al.* 2013) and the Adult NTR (ANTR; Willemsen *et al.* 2013). In the YNTR, twins have been registered at birth by their parents since 1987 (Bartels *et al.* 2007). Maternal ratings at ages 3, 7, 10, and 12, and self-ratings at ages 12, 14, 16 and 18 years were included in the analysis. When young twins reach age 18, they are enrolled in the ANTR. The ANTR includes adolescent and adult twins, who were recruited through city councils and other means (Willemsen *et al.* 2013). The twins completed the SxAnxDep subscale in 1991, 1995, 1997, 2000, 2002, and 2009. All twins between the ages of 12 and 63 years were included in the current study. The total dataset comprised 49 524 twins, including 7863 monozygotic (MZ) and 15 815 same-sex and opposite-sex dizygotic (DZ) complete twin pairs. The majority (60%) participated in more than one survey; with 10% taking part ≥ 5 times (see online Supplementary Table S1). To analyse the data as a function of age, data were reordered into age bins spanning 2 years. For instance, a 21-year-old twin in the 1991 survey and a 21-year-old twin in the 2003 survey were both included in the 20–21 years age group. Up to age 30, all age bins included more than 1000 observations. Across the entire dataset, no 2-year age bin included fewer than 130 observations.

Phenotypes

SxAnxDep scores were obtained from the ‘anxious-depressed’ subscale from the age-appropriate questionnaires from the Achenbach System of Empirically Based Assessment (ASEBA): the Child Behavior Checklist CBCL/1.5-5 (Achenbach & Rescorla, 2000) and CBCL/4-18 (Achenbach & Rescorla, 2001), the Youth Self-Report (YSR; Achenbach, 1990), and the Adult Self-Report (ASR; Achenbach & Rescorla, 2003). The instruments were designed to measure comparable constructs over the ages, and are similar in item content. Mothers were asked to rate the extent to which a statement described their child on a 3-point scale (‘not true’, ‘somewhat or sometimes true’, ‘very or often true’). An example item of the mother rating SxAnxDep scale is ‘Unhappy, Sad or Depressed’. In the self-rating scales, used from age 12 onwards, the item is phrased as ‘I am Unhappy, Sad or Depressed’, and the response format was the same. Similar strong parallels exist for the other items for mother-rated and self-rated SxAnxDep. Full sample questionnaires can be found

at <http://www.aseba.com>. At all ages, except age 3 years, the number of items was the same (nine items at age 3, 16 items at all other ages). The instrument has been found to be measurement invariant in adolescence across age and sex (Fonseca-Pedrero *et al.* 2012).

While the CBCL, YSR and ASR ‘anxious depressed’ scale scores are a good predictor of anxiety disorders and depression diagnosis, a high score is not equivalent to a diagnosis (Bird *et al.* 1991). Using composite international diagnostic interviews (CIDI), DSM-IV anxiety and depressive diagnoses were assessed in 1331 (345 cases, 986 controls) adults from the present sample in 1997 and 2007. We compared the SxAnxDep scores of these subjects with their CIDI diagnoses using a ROC curve analysis. The area under the curve for all anxiety and mood disorders was fair at 0.76. For major depressive disorder, the area under the curve was 0.75, and for generalized anxiety disorder 0.78. The use of a continuous measure is consistent with the dimensional view of psychopathology, and with diagnoses based on continuous measures (Markon *et al.* 2011).

Statistical model

Structural equation modelling was employed to analyse the mean trend across age, sex differences in the mean trend, and the covariance structure of SxAnxDep across age. Estimates of the mean trend for men and women are further analysed using weighted least squares (WLS) (see online Supplementary material). A genetic simplex model was chosen to analyse the longitudinal data (Boomsma & Molenaar, 1987; Neale *et al.* 2006) and to estimate heritability at each age as well as the phenotypic, genetic, and environmental stability across the lifespan. Fig. 1 provides a graphical description of the model, which is further detailed in the online Supplementary material. In brief, the model allows partitioning the variance in the observed data into variance due to additive genetic factors (A), unique environment (E, not shared between twins) and a common environment (C, shared between twins within a family). The model further allows estimation of the stability of the effects of genetic and environmental factors over age, and to establish the extent to which new effects, called innovations, come into play (Gillespie *et al.* 2004). From the longitudinal model we derived heritability estimates at each age and calculated genetic and environmental correlations between the ages. For model details, and identification, see the online Supplementary material.

Results

Fig. 2 shows means for men and women and the *post-hoc* fitted mean trends. The mean in males for

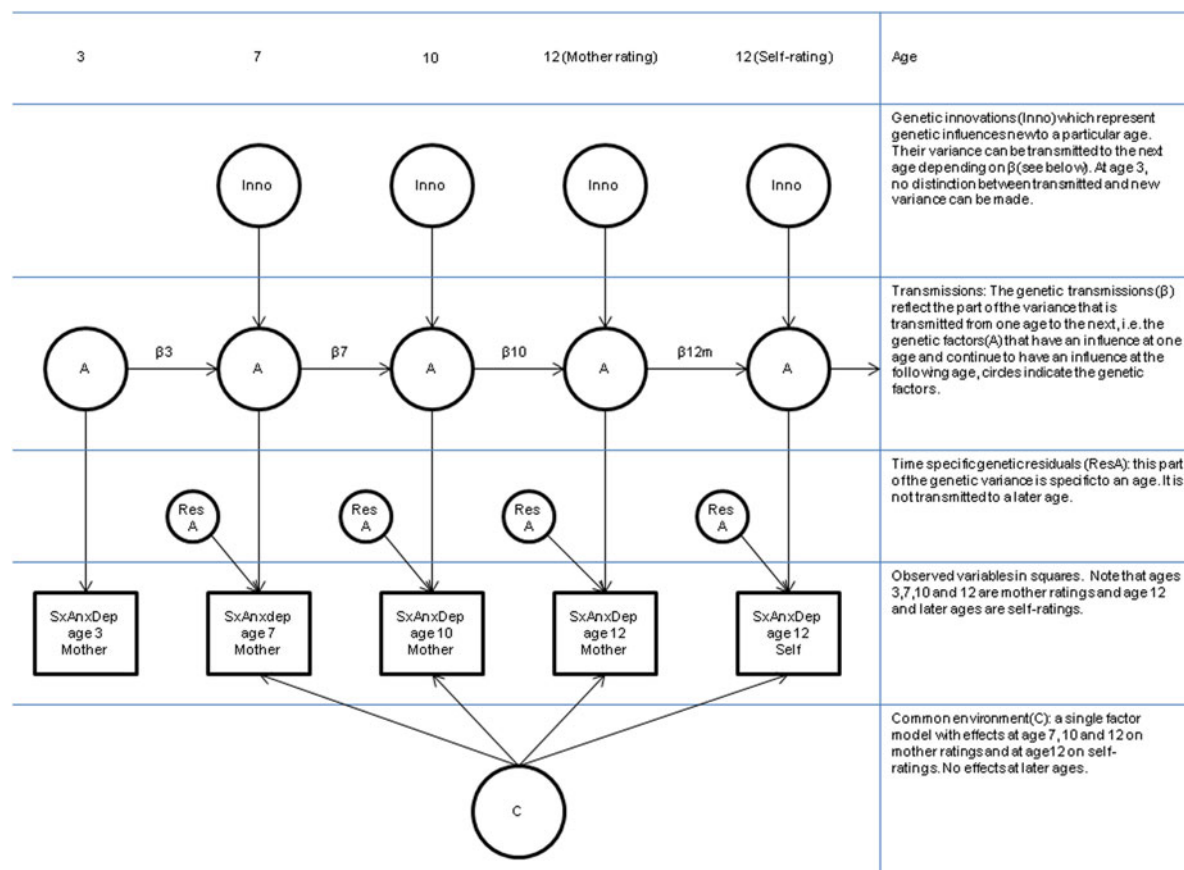


Fig. 1. Path model representing the genetic simplex model. The simplex model describes an autoregressive process, in which latent genetic values (A) at age t are regressed (β) on previous latent values ($t - 1$). In addition, at each age, novel genetic influences, called innovations (Inno) may come into play. Genotypic values thus consist of a part that is transmitted and a part that is innovation (except at the first age at which data are observed, where such a distinction cannot be made). The residual variance in the phenotype may also be influenced by additive genetic (r_A) factors. The unique environmental (E) process is not depicted to avoid clutter but is structurally identical to the genetic process. The common environment shared between twins is modeled as a single factor, loading on observed symptoms of anxiety and depression (SxAnxDep) scores at ages 7, 10, 12 (mother) and age 12 (self-) ratings.

SxAnxDep in childhood was 2.79. Between ages 12 and 28 years the mean SxAnxDep scores significantly increased ($b = 0.19$, $t = 5.568$, $df = 1$, $p < 0.001$). After age 28, the means in males no longer significantly changed ($b = -0.06$, $t = -1.922$, $df = 1$, $p = 0.062$). In females, the mean SxAnxDep scores in childhood was 2.97. Between the ages of 12 and 28 years the mean SxAnxDep scores significantly increased ($b = 0.68$, $t = 14.35$, $df = 1$, $p < 0.001$). After age 28 the SxAnxDep scores in females significantly declined ($b = -0.18$, $t = -5.477$, $df = 1$, $p < 0.001$). The sex difference in SxAnxDep score in childhood was not significant ($b = 0.17$, $df = 1$, $t = 0.911$, $p = 0.36$), while the increase between ages 12 and 28 years was significantly steeper for females than for males ($b = 0.43$, $t = 6.665$, $p < 0.001$). In this trends analysis means were weighted for sample size (see online Supplementary material for more details).

Fig. 2b depicts the heritability estimates by sex and age for seven broad age intervals, with their 99% confidence intervals (CIs). The proportion of variance attributable to genetic factors (i.e. heritability) in males and females differed little, except in the 36–43 years groups.

The observed MZ and DZ twin correlations (presented in online Supplementary Table S2) and the cross-twin cross-age correlations (online Supplementary Fig. S1) suggested a genetic simplex model as longitudinal correlations decreased over time (Boomsma & Molenaar, 1987) and MZ correlations were higher than DZ correlations. Based on previous analyses in this sample (Boomsma et al. 2008) we expected the presence of common environmental effects in childhood.

Having fitted the genetic simplex model to the data (model fits online Supplementary Table S3), we obtained heritability estimates for SxAnxDep at each

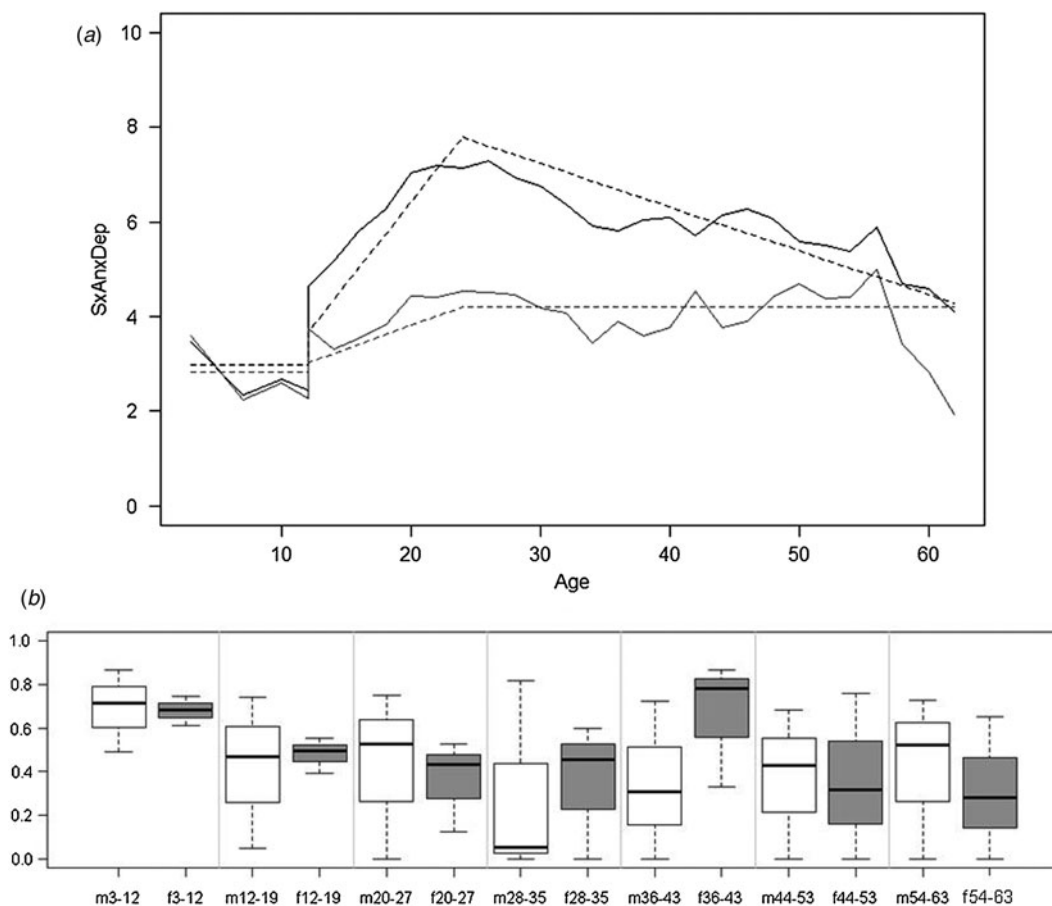


Fig. 2. (a) Mean trends in symptoms of anxiety and depression (SxAnxDep) across the lifespan for females (black) and males (grey). Solid lines are raw means. Dashed lines are fitted trend lines. (b) Univariate heritability estimates across seven broad ages (maternal ratings 3–12, self-report ratings 12–19, 20–27, 28–35, 36–43, 44–53, 54–63 years) separately by sex (m, males; f, females).

2-year age bin between ages 3 and 63 years. The estimates are given in Fig. 3a, which shows that heritability (h^2 , Fig. 3a) declined from childhood (around 60–70%) to adulthood (around 40–50%). This decrease in heritability was due to an increase in environmental variance (V_E , Fig. 3b), and not to a decrease in genetic variance (V_A , Fig. 3b), as can be seen in the plot of the absolute variance components estimates in Fig. 3b. Part of the increase in the unique environmental variance is associated with the switch from mother ratings to self-ratings (see the ‘jump’ in Fig. 3b at age 12 years). However, Fig. 3b shows that the unique environmental variance continues to rise after the switch from mother ratings to self-ratings.

At age 12, we looked at the correlations between self-ratings and maternal ratings and found that the agreement was moderate (phenotypic correlation 0.35). The genetic and environmental contributions to this correlation were 56% (genetic), 27% (unique environmental), and 17% (shared environmental).

Based on the longitudinal model (Table 1), we derived the phenotypic, genetic, and environmental correlations between SxAnxDep across different ages and visualized these in heat maps (Fig. 4 and online Supplementary Fig. S1). The phenotypic correlations (i.e. observed stability) between two successive ages ranged from 0.29 to 0.63 during childhood (up to age 12), from 0.48 to 0.70 during adolescence (age 12–18), from 0.64 to 0.77 between ages 18 and 32, and from 0.45 and 0.86 from age 32 years onwards. Correlations between successive ages did not differ between males and females ($t = -0.57$, $df = 814$, $p = 0.57$). The genetic correlations between subsequent ages during childhood (mean $r = 0.71$, range 0.4–0.94) and adulthood (mean $r = 0.92$, range 0.85–1) were large (Fig. 4). Environmental correlations between subsequent ages were clearly lower than genetic correlations in childhood (mean $r = 0.31$, range 0.03–0.47), and adulthood (mean $r = 0.60$, range 0.47–0.73), although an increase was observed with age. The

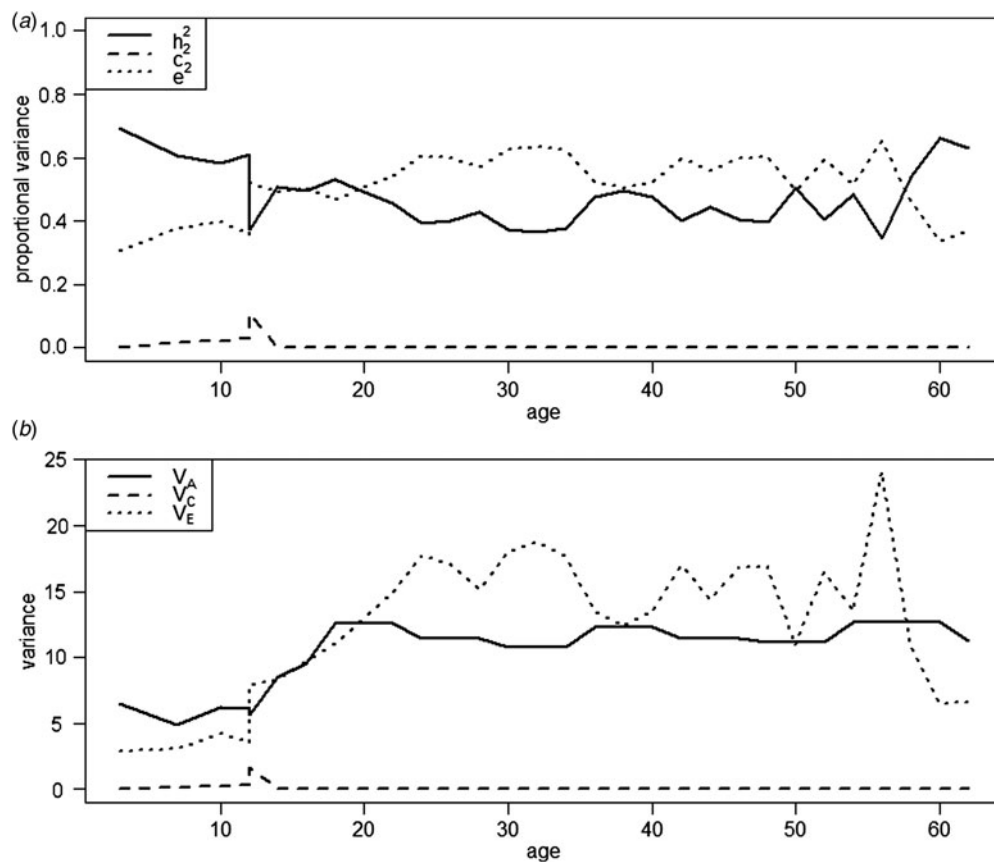


Fig. 3. (a) Proportions of variance explained by genetic factors (h^2 , —), common environment (c^2 , - - -), and unique environment (e^2 , ···) at each age as derived from the model. (b) Variance components V_A , V_C and V_E at each age as derived from the model.

10-year lag genetic correlations (online Supplementary Fig. S2) were large (mean 0.92, range 0.89–0.97), while 10-year lag environmental correlations were moderate (mean 0.32, range 0.18–0.66, online Supplementary Fig. S2) due to the relatively large proportion of environmental variance attributable to innovation. All together, stability was largely attributable to genetic effects, while the increase in stability from childhood to adulthood was due to an increase in environmental correlations with age.

Discussion

The aim of this study was to gain insight into the aetiology of the variation in SxAnxDep across the lifespan. Overall, genetic effects explained around 40% of the phenotypic variance at each age beyond age 7 years, and contributed greatly to the stability in SxAnxDep across age. We observed important differences between childhood and adulthood. Specifically, genetic innovation was observed during childhood and adolescence, but not after age 18. So after age 18, genetic effects are highly stable. This differs markedly from

the results obtained for the unique environmental influences, which showed innovations at each age, and less stability. Unique environmental factors contributed primarily to short-time stability, but with increasing age, the contribution to stability of these environmental factors increased. This resulted in increasing stability in SxAnxDep with age corroborating previous findings by Kendler *et al.* (2011). This comprehensive picture of the aetiology of the course of SxAnxDep is in line with previous studies of SxAnxDep in either childhood/adolescence or in adulthood (Gillespie *et al.* 2004; Boomsma *et al.* 2008; Kendler *et al.* 2008b, c; Trzaskowski *et al.* 2012).

When analysing data from children (maternal ratings) and adolescents and adults (self-ratings), the question arises if the results are influenced by the different raters. The availability of self- and maternal reports of SxAnxDep at age 12 allowed us to address this question. We found a moderate correlation between mother and child ratings (0.35). Moderate correlations among different raters have been observed before (e.g. Ferdinand *et al.* 2004). In the current dataset, we could establish that the correlation was largely

Table 1. Parameter estimates and variance components from the best longitudinal model

t	Age (years)	Transmission (regression) (β)		Innovation (variance) (ζ)		Residual (variance) (r)		C-factor loadings C	Variance decomposition		
		A	E	A	E	A	E		h^2	c^2	e^2
1	3	0.35	0.03	6.51	2.87	0	0		0.69		0.31
2	7	1	0.59	3.15	2.05	0.97	1.01	0.38	0.61	0.02	0.38
3	10	0.87	0.57	1.26	2.48	0.97	1.01	0.48	0.58	0.02	0.4
4	12(m)	0.48	0.46	1.22	1.53	0.97	1.01	0.58	0.61	0.03	0.36
5	12(s)	1.22	0.63	4.17	3.09	0.34	4.28	1.28	0.37	0.11	0.52
6	14	0.9	0.73	0.25	2.56	0.34	4.28		0.51		0.49
7	16	0.93	0.86	2.58	3.37	0.34	4.28		0.5		0.5
8	18	1	0.61	2.7	3.58	1.81	3.43		0.53		0.47
9	20	1	0.85	0	6.68	1.81	3.43		0.49		0.51
10	22	1	0.83	0	4.71	1.81	3.43		0.46		0.54
11	24	1	0.76	0	5.11	0.67	4.72		0.39		0.61
12	26	1	0.78	0	4.87	0.67	4.72		0.4		0.6
13	28	1	0.97	0	2.93	0.67	4.72		0.43		0.57
14	30	1	0.87	0	1.76	0	6.55		0.37		0.63
15	32	1	0.72	0	3.54	0	6.55		0.36		0.64
16	34	1	0.93	0	4.69	0	6.55		0.38		0.62
17	36	1	0.94	0	0.07	1.55	3.88		0.48		0.52
18	38	1	0.81	0	0	1.55	3.88		0.5		0.5
19	40	1	0.76	0	4.01	1.55	3.88		0.48		0.52
20	42	1	0.75	0	7.63	0.65	3.8		0.4		0.6
21	44	1	0.74	0	3.23	0.65	3.8		0.44		0.56
22	46	1	0.87	0	7.32	0.65	3.8		0.4		0.6
23	48	1	0.56	0	6.93	0.4	0		0.4		0.6
24	50	1	1.4	0	0.53	0.4	5.22		0.5		0.5
25	52	1	0.81	0	0	0.4	5.22		0.4		0.6
26	54	1	1.26	0	2.98	1.93	3.25		0.48		0.52
27	56	1	0.57	0	4.36	1.93	3.25		0.35		0.65
28	58	1	0.52	0	0.86	1.93	3.25		0.54		0.46
29	60	1	1.38	0	1.2	1.93	3.25		0.66		0.34
30	62			0	0.44	0	0		0.63		0.37

A, Genetic factors; E, unique environment; C, common environment; h^2 , proportion of variance explained by genetic factors = heritability; c^2 , proportion of variance explained by common environmental factors; e^2 , proportion of variance explained by unique environmental factors; m, mother; s, self. See Fig. 1 for an explanation of the terms transmission, innovation and residual variance.

attributable to genetic effects (56%), indicating that different raters to a large extent seem to agree on the genetically influenced SxAnxDep phenotype.

An important result from the current study is the clear explanation of the decline in heritability, which coincides with the switch from maternal to self-ratings. Heritability was highest in childhood (70–50%), and dropped in adulthood (50–35%), and remained stable to the age of 63. These heritability estimates are largely in line with those found in earlier studies (Sullivan *et al.* 2000; Johnson *et al.* 2002; Kendler *et al.* 2006; Rapee *et al.* 2009; Rice, 2009; Franz *et al.* 2011; Sakolsky *et al.* 2012). The longitudinal

data indicate that a decrease in heritability is not due to a decrease in genetic variance, but that there is an increase in environmental variance (see Fig. 3), which leads to a *relatively* lower influence of genetic factors. Common environmental effects were only present at ages 7–12 and explained a relatively small proportion of the variance (~2–11%). Mean sex differences in SxAnxDep emerged after age 12, around the same time when the increase in environmental variance is seen, peaked at age 28, and decreased thereafter. Figure 2b shows that there were no large sex differences in the estimates of genetic and environmental influences.

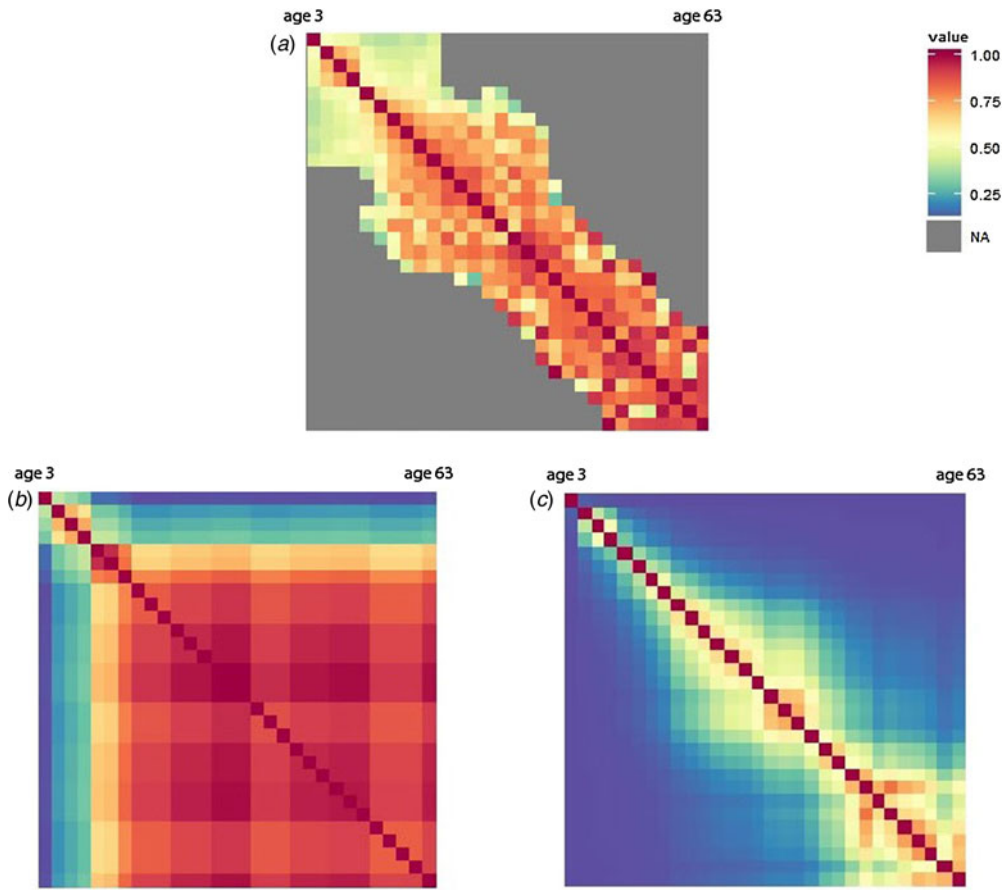


Fig. 4. Correlation heat maps to represent (a) the observed correlations between ages 3 and 63 years, (b) the model implied genetic correlations and (c) the model implied unique environmental correlations.

For clinical practice, it is important to note that environmental effects contribute to change, but also to short-term stability. This suggests that an improvement in SxAnxDep can be accomplished by positive environmental experiences, such as beneficial therapy or positive life events, and that increases of SxAnxDep can be caused by negative experiences, such as adverse life events. Importantly, these effects can endure for several years. An earlier study in part of this sample showed that SxAnxDep increase after negative life events, but that higher scores on SxAnxDep also precede negative life events (Middeldorp *et al.* 2008). Thus, individuals already suffering from anxiety or depression are at increased risk of experiencing negative events that can exacerbate their symptoms over an extended period of time. These results further underline the importance of addressing the environment in therapy (e.g. increasing social support or involving significant others). They can also suggest a possible interdependency between the individual and the environment, which may give rise to genotype-environment covariance (Dolan *et al.* 2014), but it was shown that the association

between life events and SxAnxDep was not explained by a shared genetic background (Middeldorp *et al.* 2008).

The results may have several implications for future research. The stability of genetic effects from childhood into adulthood is important as it indicates that genetic vulnerability is present from early onwards and remains a risk factor throughout life. These results suggest that gene-finding studies may include adults aged between 18 and 63 years, as we have observed little age-related heterogeneity in genetic effects. However, we recognize that high stability in polygenic effects does not rule out age effects at the level of a single causal genetic variant.

Rapee *et al.* (2009) have already pointed out that with respect to childhood anxiety disorders ‘current knowledge of the role and mechanisms of environmental factors is especially poor’ (page 331). The increase in environmental stability over time is an intriguing finding that warrants further investigation. The question is whether new environmental effects on SxAnxDep, decrease with age, or whether the impact of events lasts longer as people age. Our results

suggest the latter given that the environmental innovation parameters do not decrease with age, while the transmission parameters increase between ages 3 and 16 years and then remain at a similar level. It is well known that childhood and especially adolescence are characterized by large developmental changes in the brain (Blakemore, 2008). Possibly, the maturation of the brain is accompanied by a more enduring effect of the environment. Kendler *et al.* (2011) also found that the environmental effects on long-term stability of SxAnxDep reach a plateau after adolescence.

We have shown that even during childhood and adolescence, part of the genetic and unique environmental innovations that appear at each age are transmitted to other ages. This signifies that there is a group of children and adolescents with a risk of enduring symptoms. Given the high disease burden for these children and society it is important to identify risk and protective factors that lead to stability over ages.

Finally, the current results for SxAnxDep are partly different from the results in a similar analysis of the aetiology of stability in attention problems (Kan *et al.* 2013). For attention problems, the genetic variance decreases from childhood to adulthood, while for SxAnxDep the genetic variance increases. Moreover, the increase in stability due to environmental effects that is observed for SxAnxDep after age 18 is not present in attention problems until the age of 30. Although it is of note that, for both phenotypes, genetic factors are most important in explaining stability, these differences indicate that the pattern of genetic and unique environmental factors throughout life do not need to be similar across psychiatric phenotypes. Our results and those for attention problems (Kan *et al.* 2013) suggest similar longitudinal genetic analysis are warranted for other psychiatric phenotypes.

Longitudinal results are essential to increase our understanding of the development and genetic architecture of psychiatric phenotypes. We made use of 25 years of data collection in subjects who entered the study at different ages, and tried to inform both clinical and molecular genetics research in psychiatry, using developmental and etiological informative models.

Supplementary material

For supplementary material accompanying this paper visit <http://dx.doi.org/10.1017/S003329171400213X>.

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Declaration of Interest

None.

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