School success in childhood and subsequent prodromal symptoms and psychoses in the Northern Finland Birth Cohort 1986

Lassila M^{1,2}, Nordström T^{1,2}, Hurtig T^{3,4,5}, Mäki P^{2,3,6,7}, Jääskeläinen E^{1,2,6}, Oinas E¹, Miettunen J^{1,2}

- 1) Center for Life Course Health Research, University of Oulu, Oulu, Finland
- 2) Medical Research Center Oulu, Oulu University Hospital and University of Oulu, Oulu, Finland
- 3) Department of Psychiatry, Research Unit of Clinical Neuroscience, University of Oulu, Oulu, Finland
- 4) PEDEGO Research Unit, Child Psychiatry, University of Oulu, Oulu, Finland
- 5) Clinic of Child Psychiatry, Oulu University Hospital, Oulu, Finland
- 6) Department of Psychiatry, Oulu University Hospital, Oulu, Finland
- 7) Department of Psychiatry, Länsi-Pohja healthcare district; Department of Psychiatry, the Middle Ostrobothnia Central Hospital, Soite; Mental health services, Joint Municipal Authority of Wellbeing in Raahe District; Mental health services and Basic Health Care District of Kallio, Finland; Department of Psychiatry, Kainuu Central Hospital, Kainuu social and healthcare district, Finland

Abstract

Background: Low IQ is a risk factor for psychosis, but the effect of high IQ is more controversial. The aim was to explore the association of childhood school success with prodromal symptoms in adolescence and psychoses in adulthood.

Methods: In the general population-based Northern Finland Birth Cohort 1986 (n=8 229), we studied the relationship between teacher-assessed learning deficits, special talents and general school success at age 8 years and both prodromal symptoms (PROD-screen) at age 15–16 years and the occurrence of psychoses by age 30 years.

Results: More prodromal symptoms were experienced by those talented in oral presentation (boys: adjusted Odds Ratio 1.49; 95% confidence interval 1.14-1.96; girls: 1.23; 1.00–1.52) or drawing (boys: 1.44; 1.10–1.87). Conversely, being talented in athletics decreased the probability of psychotic-like symptoms (boys: OR 0.72; 0.58–0.90). School success below average predicted less prodromal symptoms with boys (OR 0.68; 0.48–0.97), whereas above-average success predicted more prodromal symptoms with girls (OR 1.22; 1.03–

1.44). The occurrence of psychoses was not affected. Learning deficits did not associate with prodromal symptoms or psychoses.

Conclusions: Learning deficits in childhood did not increase the risk of prodromal symptoms in adolescence or later psychosis in this large birth cohort. Learning deficits are not always associated with increased risk of psychosis, which might be due to e.g. special support given in schools. The higher prevalence of prodromal symptoms in talented children may reflect a different kind of relationship of school success with prodromal symptoms compared to full psychoses.

Keywords: adolescents, children, creativity, learning deficits, psychosis, school

1. Introduction

Early school success may be used to predict individuals' later life. School success is affected by many factors, for instance motivation, attention or memory, learning deficits or IQ. Special talents in certain subjects require also creativity. The onset of psychosis is often in early adolescence, when major physical and behavioral changes occur (Paus *et al.* 2008). Males have higher risk of onset of schizophrenia in early adulthood (van der Werf *et al.* 2014). Onset of psychosis is often preceded with premorbid prodromal symptoms. Changes in cognition or IQ manifesting already in childhood or adolescence before the prodrome have been reported (Davidson *et al.* 1999, Kremen *et al.* 1998). Information on school success could be used to search these predictors: such information can easily be obtained on population level and is thus applicable in large epidemiological studies.

Sometimes poor school success may indicate cognitive impairment characteristic for the trajectory to psychosis. Low and especially deteriorating premorbid IQ in childhood or puberty has been seen to precede the appearance of psychotic symptoms in adulthood. (Kremen et al. 1998, Fuller et al. 2002, Matheson et al. 2011). Some studies show a linear dose-response relationship between decreasing IQ and increased risk of psychosis (David et al. 1997, Davidson et al. 1999, Khandaker et al. 2011), whereas others suggest that IQ influences the risk of psychosis most strongly at the lowest range of IQ distribution (Schulz et al. 2014). The relationship between lower than average IQ and psychosis might depend on a third factor, such as obstetric complications (Di Prinzio et al. 2018, Sussmann et al. 2019).

Bipolar disorder has been associated with excellent school performance (MacCabe *et al.* 2010). Excellent grades are not solely due to high IQ but also motivation and productivity, which is also higher in hypomania. Less evidence exists about good school performance and the risk of schizophrenia and other psychoses (Isohanni *et al.* 1999, MacCabe *et al.* 2008). Occurrence of different prodromal symptoms and types of psychosis seems to differ by gender. Gender differences in cognition in premorbid phase and during illness have been suggested but less studied (Barajas *et al.* 2015). One study suggested that male with first-episode psychosis had higher IQ than women (Hui *et al.* 2016). High IQ might affect the course of illness, e.g. suicide risk in psychotic disorder have been found to be higher in those with good school success (Alaräisänen *et al.* 2006, Nordentoft 2007). However, high childhood IQ predicts a better outcome in subjects who later develop schizophrenia (Munro *et al.* 2002).

In order to obtain more information about the association between school success and psychoses, it is worthwhile to explore both superior and inferior performers. The aim of this study was to study how teacher-assessed learning deficits or special talents in certain school subjects are associated with later self-reported psychotic-like symptoms and with the occurrence of psychoses in the Northern Finland Birth Cohort 1986 among boys and girls. We hypothesized that learning deficits would associate with more psychotic-like symptoms and with psychosis, whereas special talents would not associate with the outcomes.

2. Methods and material

2.1 Study population

The Northern Finland Birth Cohort 1986 is a longitudinal birth cohort including 9 362 mothers and their 9 432 live-born children with expected date of birth between July 1st 1985 and June 30th 1986 in Northern Finland (Järvelin *et al.* 1993). Subjects who denied the use of their data (n=256) and those with a known diagnosis of intellectual disability (n=115) were excluded. This was done based on knowledge concerning the association between intellectual disability and psychoses reported previously (Myrbakk *et al.* 2008).

2.2 Learning deficits and special talents in childhood

At the age of 8 years, a postal questionnaire concerning school behavior and performance was sent to the teachers with permission from the parents (Taanila *et al.* 2004). Teachers replied with information on learning deficits and special talents about 4 193 (86.0%) of the boys and 4 036 (88.9%) of the girls. School success was assessed based on this questionnaire, and the final sample included 8 229 individuals.

Learning deficits and special talents of the subjects were assessed on the basis of no/yes questions. Learning deficits included difficulties in reading, writing or mathematics. Special talents were assessed both in academic and non-academic subjects. Special talents included creativity related talents: music, drawing, athletics, craft, oral presentation and acting and written presentation. Information about mathematical talent was gathered from answers to open questions about pupils' talents. The general success in school was graded to be either below average, average or above average.

2.3. Prodromal symptoms in adolescence

A questionnaire about subjects' lifestyles and habits was given to them during their clinical 16-year examination in 2001–2002 (Mäki *et al.* 2014).

Prodromal symptoms include both mild psychotic-like and unspecific psychiatric symptoms, which are associated with increased risk of transition to psychosis. The lack of specificity makes it challenging to separate the pre-psychotic state from some other mental disorders. The PROD-screen is an easily applicable tool for assessing the prodromal symptoms in clinical work and also for screening healthy individuals for prospective population-based study. The simplicity of the PROD-screen is however to the detriment of its power to detect specific pre-psychotic symptoms. The questionnaire included the PROD-screen, which consists of 21 questions (no/yes) screening psychotic-like (prodromal) symptoms, of which 12 are considered as specific symptoms for psychosis (Heinimaa et al. 2003). Others include symptoms such as depression and anxiety that are also associated with pre-psychotic states, although they are less specific (Therman et al. 2011). The total score of the specific symptoms was used in the current study the score was dichotomized (0-3 points vs. over 3 points), based on previous research. About half of the future psychotic cases scored over 3 points in the PROD-screen, odds ratio (95% confidence interval) for psychosis risk being 1.55 (OR 2.36 (95% CI 1.68-3.31) (unpublished result)). PROD-screen has been found to classify correctly 77% of prodromal cases (Heinimaa et al. 2003). The 12 specific items have Cronbach alpha value of 0.69 in the current sample. The missing information due to replying

only to some of the PROD-screen questions was taken into consideration in the analyses by assuming that the subject replied to the missing items with the same yes/no percentages as in the filled part of the questionnaire. Ruling out the subjects (0.5%) who did not reply to more than 2 specific items or to 3 of the total items, the screening information was available from 3 225 (68.7%) boys and 3 341 (75.7%) girls.

2.4. Psychoses in adolescence and early adulthood by 30 years of age

The information about psychoses were derived from multiple sources of register data: the Care Register for Healthcare (1998–2015), Primary Healthcare Outpatient Registers (2011–2015), Specialized Care Outpatient Registers (1998–2015), register for disability pensions from the Finnish Centre for Pensions (1998–2013) and register for reimbursable medication from the Social Insurance Institute (1998–2005).

2.5. Possible confounding factors

The distribution of family background factors by gender is seen in Table 1. The presented data consists of possible confounding factors which could affect the rates of prodromal symptoms or psychoses, these were selected based on earlier studies. These items were included in the regression models as covariates. The data included gender, parental psychosis (no/yes) (Keskinen *et al.* 2015), family type (married/cohabiting or single/divorced) (Chen *et al.* 2014) and parental education (less than 9 years of comprehensive school/comprehensive or elementary school/matriculation examination) (Frissen *et al.* 2015). Family type information was collected from a questionnaire made during pregnancy by the 24th gestational week. Information about parental education was collected from a questionnaire delivered to the parents when the subjects were 16 years old. Information on parental psychosis was collected from different registers (Care Register for Healthcare (1998–2015), Primary Healthcare Outpatient Registers (2011–2015), Specialized Care Outpatient Registers (1998-2015) and Finnish Center for Pensions (1998–2013)).

2.6. Statistical methods

The statistical difference between genders regarding family background factors and the prevalence of learning deficits and special talents was assessed by using Pearson chi-square tests. Odds ratios (OR) were calculated using binary logistic regression for having more than three specific PROD-screen items and for having psychoses for those with

learning deficits or special talents. Hazard ratios (HR) were calculated using Cox regression models with time of death and emigration as censoring points. Adjusted odds and hazard ratios were calculated using parental marital status, basic education, psychosis and family's place of residence as covariates. All results are presented by gender, because school variables and outcome variables differed by gender. IBM SPSS Statistics 24.0 was used in the analyses. P-values less than 0.05 were considered as statistically significant.

2.7. Attrition analysis

Attrition in the questionnaire to teachers (pupils' age 8) and the self-reported PROD-screen (pupils' age 15–16) was assessed regarding family type (marital status of the mother), type of residence (urban/rural), mother's and father's education (less than 9 years/9–12 years/more than 12 years and parental psychosis (no/yes). Teachers' assessment was available for 91.0% of the boys and 92.4% of the girls. PROD-screen was available for 70.4% of the boys and 76.7% of the girls. Regarding boys, participants and non-participants differed significantly in most family background factors in both follow-up surveys, whereas among girls the only significant finding was that those with married mothers participated more at age 15–16 years. Differences between the groups of participants and non-participants can be seen in Supplement Tables 1 and 2.

3 Results

3.1 Prevalence of special talents and learning deficits

The prevalence of learning deficits and special talents by gender is presented in Table 2. Overall, boys had more learning deficits and less special talents than girls. The prevalence of different learning deficits differed between 9.2% and 22.2% in boys and between 8.8% and 12.3% in girls. Regarding special talents, the range was between 6.9% and 27.4% in boys and between 3.2% and 27.1% in girls. In athletics and mathematics boys were more frequently talented than girls. However, boys were assessed to perform generally less well at school than girls. Of the boys, 13.1% were assessed to be below average while the percentage for girls was 6.9%. Above average performance was registered in 30.4% of boys and 42.4% of girls.

3.2 Prodromal symptoms in adolescence

The number of subjects who had at least three specific PROD-screen items with unadjusted and adjusted odds ratios according to childhood school variables are presented in Table 3. Learning deficits in childhood were not associated with having a PROD-score above 3 reported in adolescence. Those with special talents in oral presentation, written presentation and drawing had higher ORs for having a PROD-score above 3. After controlling for confounding factors, the association remained with oral presentation for boys and girls and drawing for boys. The adjusted ORs for oral presentation were 1.49 (95% CI 1.14–1.96) for boys and 1.23 (1.00–1.52) for girls and for drawing 1.44 (1.10–1.87) for boys. Talent in athletics appeared to be protecting for boys lowering the adjusted OR to 0.72 (0.58–0.90). The results with general school success showed the same trend; the prevalence of prodromal symptoms decreased with below average boys and increased with above average girls: the adjusted ORs were 0.68 (0.48–0.97) for boys and 1.22 (1.03–1.44) for girls.

3.3. Psychoses in adulthood

Table 4 shows the number of psychoses experienced by age 30 with unadjusted and adjusted hazard ratios according to school variables. In total 2.4% of boys and 2.1% of girls received a psychosis diagnosis during the follow-up until age of 30 years. Of the 214 cases with psychoses, 69 (46 boys) had schizophrenia, 49 (18 boys) affective psychosis and 96 (55 boys) other psychoses. Learning deficits and special talents in childhood were not associated significantly with psychoses in adulthood (until age of 30 years). However, some gender differences were observed. In girls, a higher number of psychoses was observed with succeeding below average at school, having problems in reading and being talented in oral presentation. With boys the trend was opposite; the same variables were associated with lower psychosis rates.

4 Discussion

4.1 Main findings

Surprisingly, learning deficits in childhood were not associated with prodromal symptoms in adolescence or with subsequent psychoses in the Northern Finland Birth Cohort 1986 sample. Prodromal symptoms appeared to be associated only with good school success

(such as talent in oral presentation and drawing). Special talent in athletics decreased the prevalence of psychotic-like symptoms in boys.

4.2. Learning deficits

In this general population based birth cohort, about every 10th child or up to every 5th boy and every 8th girl had experienced some learning deficits at the age of 8 years. None of the learning deficits were associated with prodromal symptoms in adolescence or with later psychoses. Sometimes learning deficits reflect low IQ, but social, motivational, attention problems and personality-related factors are also important aspects affecting learning deficits. Similarly, in the meta-analysis of premorbid intelligence and schizophrenia, some studies have not found a significant connection (Walker *et al.* 2002, Khandaker *et al.* 2011). Consistently but not significantly lower prodromal symptom scores among subjects with learning deficits and respectively high scores among talented subjects show a trend that may arise from the subjects' ability to describe their symptoms.

In the Finnish school system, many different forms of special support are directed towards children with problems in learning and could possibly act as a protecting factor (Ström & Hannus-Gullmets 2015). In personalized curriculums learning goals are set to be realistic concerning the students' abilities and restrictions. Special support can give students with learning deficits the experience of succeeding, and thus could prevent social isolation which could be assumed to be positive for mental health. Unfortunately, studies about the possible connections between learning deficits, special education and mental health are scarce.

Ruling out subjects with known intellectual disability from the analyses could explain the results of this study, this was done based on the significant number of intellectually disabled subjects in the learning deficits group and on evidence showing that intellectual disability is a risk factor in itself (Jacobson 1990). In studies including intellectually disabled subjects, the risk-increasing effect of learning deficits may be overestimated.

In this study, we investigated early learning deficits detected at the age of 8 years. Early learning deficits were chosen to reveal their effect on psychosis risk, avoiding the confounding effect of cognitive deterioration characteristics on the trajectory of the illness itself.

4.3 Special talents

Several special talents in childhood were associated with subsequent increased prodromal symptoms in adolescence. These talents (oral and written presentation and drawing) can be representations of intelligence but also of creativity. Higher performance in verbal learning and fluency is associated with a liability to bipolar disorder, but not to schizophrenia. Verbal fluency could have improved evolutionary fitness and its association with bipolar disorder could offer one explanation of why bipolar disorder has persisted in the population (Higier *et al.* 2014). In one study, subjects who later developed schizophrenia excelled in drawing and arts at the age of 12 (Helling *et al.* 2003).

The link between creativity and mental disorders is supported by many studies, suggesting that the link is strongest with affective disorders (Andreasen *et al.* 1987). A particular link to affective disorders could explain why only prodromal symptoms but not psychoses were affected. Because of the non-specific character of the PROD-screen discussed later in this article, subjects with high scores also include those experiencing affective rather than psychotic symptoms. Different special talents (or types of creativity) are suggested to have separate links to bipolarity and schizotypy-schizophrenia spectrum disorders (Richards 2001). Intelligent and verbally talented individuals can also be thought to be more able to describe their sensations leading to higher self-reporting of symptoms. The self-reporting of symptoms has found to be affected by cognitive ability and personality factors (Enns *et al.* 2000).

Most previous studies about creativity and mental disorders have been in relation to adults, showing that some mental disorders are more frequent in artists (Andreasen 1987, Jamison *et al.* 1989, Kyaga *et al.* 2011). In a recent large Swedish register study, it was found that artistic creativity at high school or university associated with later mental health problems (MacCabe *et al.* 2018). Interestingly, they found visual arts to associate with psychosis as we did in the current study. Special talents emerging during childhood could allow the assessment of causality but have been less studied. Our result suggests that special talents emerging already during childhood could relate to mental health in adolescence.

Some theories have been suggested to explain the mechanism underlying the link between creativity (or special talents) and mental disorder, one being the shared genetic vulnerability model. This model suggests that some hereditary traits such as reduced latent inhibition are common for creativity and psychopathology, increasing the amount of information processed consciously (Baruch *et al.* 1988, Carson 2011). Simultaneous manifestation of

protective factors, such as high IQ, controls the processing of this information increasing creativity and protecting against the formation of mental disorders (Carson 2011). Horwood *et al.* (2008) found in their ALSPAC birth cohort study that low IQ measured at the age of 8 years was associated with increased psychotic-like symptoms at the age of 12 years, but a weaker association was also observed with high IQ. However, the risk of schizophrenia declines with increasing IQ (Zammit *et al.* 2004). In our study the increase was also observed only in psychotic-like symptoms, not in actual psychoses.

The status of athletics as a protecting factor could be explained by the known positive effect of physical activities on mood and mental health (Griffiths *et al.* 2014, Wiles *et al.* 2008) or as selection process relating to minor motor difficulties associating also with psychosis risk (Filatova *et al.* 2018). Physical activity has been linked with psychosis risk also in the current birth cohort (Koivukangas *et al.* 2010).

4.4. Psychotic-like symptoms

Despite its simplicity, PROD-screen has been shown to be useful in evaluating psychosis risk (Heinimaa *et al.* 2003). The four prodromal symptoms regarding social withdrawal are more frequent in patients with first-episode psychosis than in patients with non-psychotic disorders and in controls. Although the patients had experienced less of these symptoms compared to patients with psychosis, those with non-psychotic disorder had experienced more social withdrawal associated symptoms in adolescence than the controls (Mäki *et al.* 2014).

The selection of the cut-off point of 3/12 specific symptoms can be justified based on the proven consistency with the more complex SIPS screening tool. However, the screen is designed for clinical settings, and it remains unclear what should be the cut-off point when the screen is applied to epidemiological settings.

4.5 Psychoses

To our surprise, we did not find any school variables in childhood to correlate with increase in later psychoses despite the results concerning the prodromal symptoms. One explaining factor could be the young age of our sample, in which not all the vulnerable individuals have yet become ill. Symptoms screened in the PROD-screen are also much more common in the population than psychosis diagnoses consistently with the continuum hypothesis of psychotic symptoms. According to this hypothesis, the occurrence of psychotic symptoms

is continuous in the population, varying from mild forms to more severe ones (Verdoux 2002). The increase in PROD-score but not in the psychosis rate could be a manifestation of milder, but still clinically relevant symptomatology. It is also possible that other factors such as student's temperament affected teachers' ratings (Mullola *et al.* 2012). Regarding talented individuals, it has been found that creative individuals have more some positive schizotypal symptoms more frequently than others (Nelson *et al.* 2010).

In our study, all psychosis diagnoses were included, not only schizophrenia. With regard to premorbid intelligence as a risk factor, some studies have shown that low premorbid intelligence is a risk factor only for schizophrenia, not for affective psychoses (Agnew-Blais *et al.* 2017). In general, it has been found that schizophrenia and affective psychoses have slightly different risk profiles (see *e.g.* Jääskeläinen *et al.* 2017). Although most risk factors overlap, the level of the risk increasing effect is often found to be greater with schizophrenia. Furthermore, less evidence is available concerning antecedents for affective psychoses (Laurens *et al.* 2015). This could partly explain why early learning deficits, most probably also associated with lower intelligence, did not predict psychoses in our sample.

4.6. Strengths and limitations

One strength of our study is the extensive data, the included survey data have good participation rates and the follow-up for psychosis diagnoses was based on nationwide registers with no attrition. Also the prospective design of the study is an advantage, information about the large general population-based study sample has been gathered starting from birth, resulting in an extensive follow-up time of 30 years. The study has very good participation rates both in childhood and adolescent surveys and we were able to use extensive nationwide registers to detect psychoses. Teachers' are in general well trained and competent to evaluate pupils' school performance such as talents (Bracken & Brown 2006), although there can be factors such as gender and student's temperament which may affect the ratings (Mullola et al. 2012).

Limitations of our study include the young age of the sample which explains the small number of psychosis cases. Evaluating the risk by PROD-screen also imposes some limitations. These are the PROD-screens' non-specificity in predicting psychoses and its use as a screening tool rather than in the clinical setting for which it was originally designed. Screening these symptoms on a population level does not have as strong predictive power for psychoses as using the screen in clinical settings and with chosen subjects. It is also

possible that response bias may explain some of the results as those children are more diligent (and thus rated highly by teachers) may be more likely to report psychotic experiences.

4.7. Conclusion

Surprisingly, learning deficits in childhood did not increase psychotic-like symptoms in adolescence or later psychosis rate in this large general population-based birth cohort, unlike in most previous studies. Learning problems are not always associated with increased risk of psychosis, which might be due to the special support given in school, such as remedial instruction, studying in small groups or guidance of a school helper, designing personalized curriculums or giving challenged students more time to finish compulsory education. The higher prevalence of psychotic-like symptoms in talented children may reflect a different kind of relationship of school success with psychotic-like symptoms compared to full psychoses.

Acknowledgements

We thank all cohort members and researchers who have participated in the study.

Financial support

This work was supported by the Brain and Behavior Research Fund, the Academy of Finland (grant number 268336, 278286); EU QLG1-CT-2000-01643 (EUROBLCS) (grant number E51560); NorFA (grant number 731, 20056, 30167); and USA / NIHH 2000 G DF682 (grant number 50945).

Conflict of interest

None.

References

Agnew-Blais J, Seidman LJ, Fitzmaurice GM, Smoller JW, Goldstein JM, Buka SL (2017). The interplay of childhood behavior problems and IQ in the development of later schizophrenia and affective psychoses. *Schizophrenia Research* **184**, 45-51.

Alaräisänen A, Miettunen J, Lauronen E, Räsänen P, Isohanni M (2006). Good school performance is a risk factor of suicide in psychoses: A 35-year follow up of the Northern Finland 1966 Birth Cohort. *Acta Psychiatrica Scandinavica* **114**(5), 357-362.

Andreasen NC (1987). Creativity and mental illness: Prevalence rates in writers and their first-degree relatives. *The American Journal of Psychiatry* **144**(10), 1288-1292.

Barajas A, Ochoa S, Obiols JE, Lalucat-Jo L (2015). Gender differences in individuals at high-risk of psychosis: A comprehensive literature review. *The Scientific World Journal* **2015**, 10.1155/2015/430735.

Baruch I, Hemsley DR, Gray JA (1988). Differential performance of acute and chronic schizophrenics in a latent inhibition task. *The Journal of Nervous and Mental Disease* **176**(10), 598-606.

Bracken, B.A. & Brown, E.F. (2006). Behavioral identification and assessment of gifted and talented students. Journal of psychoeducational assessment **24**(2), 112–122.**Carson SH** (2011). Creativity and psychopathology: a shared vulnerability model. *Canadian Journal of Psychiatry. Revue Canadienne De Psychiatrie* **56**(3), 144-153.

Chen F, Wang L, Heeramun-Aubeeluck A, Wang J, Shi J, Yuan J, Zhao X (2014). Identification and characterization of college students with attenuated psychosis syndrome in China. *Psychiatry Research* **216**(3), 346-350.

David AS, Malmberg A, Brandt L, Allebeck P, Lewis G (1997). IQ and risk for schizophrenia: a population-based cohort study. *Psychological Medicine* **27**(6), 1311-1323.

Davidson M, Reichenberg A, Rabinowitz J, Weiser M, Kaplan Z, Mark M (1999). Behavioral and intellectual markers for schizophrenia in apparently healthy male adolescents. *The American Journal of Psychiatry* **156**(9), 1328-1335.

Di Prinzio P, Morgan VA, Björk J, Croft M, Lin A, Jablensky A, McNeil TF (2018). Intellectual disability and psychotic disorders in children: association with maternal severe mental illness and exposure to obstetric complications in a whole-population cohort. *American Journal of Psychiatry*, 2018 Oct 3. doi: 10.1176/appi.ajp.2018.17101153.

Enns MW, Larsen DK, Cox BJ (2000). Discrepancies between self and observer ratings of depression. The relationship to demographic, clinical and personality variables. *Journal of Affective Disorders* **60**(1), 33-41.

Filatova S, Koivumaa-Honkanen H, Khandaker GM, Lowry E, Nordström T, Hurtig T, Moilanen K, Miettunen J (2018). Early motor developmental milestones and schizotypy in the Northern Finland Birth Cohort Study 1966. *Schizophrenia Bulletin* **44**(5), 1151-1158.

Fink A, Weber B, Koschutnig K, Mathias Benedek, Reishofer G, Ebner F, Papousek I, Weiss EM (2014). Creativity and schizotypy from the neuroscience perspective. *Cognitive, Affective & Behavioral Neuroscience* **14**(1), 378-387.

Frissen A, Lieverse R, Marcelis M, Drukker M, Delespaul P, GROUP Investigators (2015). Psychotic disorder and educational achievement: a family-based analysis. *Social Psychiatry and Psychiatric Epidemiology* **50**(10), 1511-1518.

Fuller R, Nopoulos P, Arndt S, O'Leary D, Ho BC, Andreasen NC (2002). Longitudinal assessment of premorbid cognitive functioning in patients with schizophrenia through examination of standardized scholastic test performance. *The American Journal of Psychiatry* **159**(7), 1183-1189.

Griffiths A, Kouvonen A, Pentti J, Oksanen T, Virtanen M, Salo P, Väänänen A, Kivimäki M, Vahtera J (2014). Association of physical activity with future mental health in older, mid-life and younger women. *European Journal of Public Health* **24**(5), 813-818.

Heinimaa M, Salokangas RK, Ristkari T, Plathin M, Huttunen J, Ilonen T, Suomela T, Korkeila J, McGlashan TH (2003). PROD-screen - a screen for prodromal symptoms of psychosis. *International Journal of Methods in Psychiatric Research* **12**(2), 92-104.

Helling I, Ohman A, Hultman CM (2003). School achievements and schizophrenia: A case-control study. *Acta Psychiatrica Scandinavica* **108**(5), 381-386.

Higier RG, Jimenez AM, Hultman CM, Borg J, Roman C, Kizling I, Larsson H, Cannon TD (2014). Enhanced neurocognitive functioning and positive temperament in twins discordant for bipolar disorder. *The American Journal of Psychiatry* **171**(11), 1191-1198.

Horwood J, Salvi G, Thomas K, Duffy L, Gunnell D, Hollis C, Lewis G, Menezes P, Thompson A, Wolke D, Zammit S, Harrison G (2008). IQ and non-clinical psychotic symptoms in 12-year-olds: Results from the ALSPAC birth cohort. *The British Journal of Psychiatry: The Journal of Mental Science* **193**(3), 185-191.

Hui CL, Leung CM, Chang WC, Chan SK, Lee EH, Chen EY (2016) Examining gender difference in adult-onset psychosis in Hong Kong. *Early Intervention in Psychiatry* **10**(4), 324-333.

Isohanni I, Isohanni M (1998). Education and mental disorders. *International Journal of Circumpolar Health* **57**(2-3), 188-194.

Isohanni I, Järvelin MR, Jones P, Jokelainen J, Isohanni M (1999). Can excellent school performance be a precursor of schizophrenia? A 28-year follow-up in the Northern Finland 1966 Birth Cohort. *Acta Psychiatrica Scandinavica* **100**(1), 17-26.

Jääskeläinen E, Juola T, Korpela H, Lehtiniemi H, Nietola M, Korkeila J, Miettunen J (2017) Epidemiology of psychotic depression - systematic review and meta-analysis. *Psychological Medicine* **48**(6), 905-918.

Jacobson JW (1990). Do some mental disorders occur less frequently among persons with mental retardation? *American Journal of Mental Retardation* **94**(6), 596-602.

Jamison KR (1989). Mood disorders and patterns of creativity in British writers and artists. *Psychiatry* **52**(2), 125-134.

Järvelin MR, Hartikainen-Sorri AL, Rantakallio P (1993). Labour induction policy in hospitals of different levels of specialisation. *British Journal of Obstetrics and Gynaecology* **100**(4), 310-315.

Karlsson JL (1999). Relation of mathematical ability to psychosis in Iceland. *Clinical Genetics* **56**(6), 447-449.

Khandaker GM, Barnett JH, White IR, Jones PB (2011). A quantitative meta-analysis of population-based studies of premorbid intelligence and schizophrenia. *Schizophrenia Research* **132**(2-3), 220-227.

Koivukangas J, Tammelin T, Kaakinen M, Mäki P, Moilanen I, Taanila A, Veijola J (2010). Physical activity and fitness in adolescents at risk for psychosis within the Northern Finland 1986 Birth Cohort. *Schizophrenia Research* **116**(2-3), 152-158.

Keskinen E, Marttila A, Marttila R, Jones PB, Murray GK, Moilanen K, Koivumaa-Honkanen H, Mäki P, Isohanni M, Jääskeläinen E, Miettunen J (2015). Interaction between parental psychosis and early motor development and the risk of schizophrenia in a general population birth cohort. *European Psychiatry* **30**(6), 719-727.

Kremen WS, Buka SL, Seidman LJ, Goldstein JM, Koren D, Tsuang MT (1998). IQ decline during childhood and adult psychotic symptoms in a community sample: A 19-year longitudinal study. *The American Journal of Psychiatry* **155**(5), 672-677.

Kyaga S, Lichtenstein P, Boman M, Hultman C, Långström N, Landen M (2011). Creativity and mental disorder: family study of 300,000 people with severe mental disorder. *The British Journal of Psychiatry* **199**(5), 373-379.

Laurens KR, Luo L, Matheson SL, Carr VJ, Raudino A, Harris F, Green MJ (2015). Common or distinct pathways to psychosis? A systematic review of evidence from prospective studies for developmental risk factors and antecedents of the schizophrenia spectrum disorders and affective psychoses. *BMC Psychiatry* **15**(1), 205.

MacCabe JH, Lambe MP, Cnattingius S, Torrång A, Björk C, Sham PC, David AS, Murray RM, Hultman CM (2008). Scholastic achievement at age 16 and risk of schizophrenia and other psychoses: a national cohort study. *Psychological Medicine* **38**(8), 1133-1140.

MacCabe JH, Lambe MP, Cnattingius S, Sham PC, David AS, Reichenberg A, Murray RM, Hultman CM (2010). Excellent school performance at age 16 and risk of adult bipolar disorder: national cohort study. *The British Journal of Psychiatry* **196**(2), 109-115.

MacCabe JH, Sariaslan A, Almqvist C, Lichtenstein P, Larsson H, Kyaga S (2018). Artistic creativity and risk for schizophrenia, bipolar disorder and unipolar depression: a Swedish population-based case-control study and sib-pair analysis. *The British Journal of Psychiatry* **212**(6), 370-376.

Mäki P, Koskela S, Murray GK, Nordström T, Miettunen J, Jääskeläinen E, Veijola JM (2014). Difficulty in making contact with others and social withdrawal as early signs of psychosis in adolescents - the Northern Finland Birth Cohort 1986. *European Psychiatry* **29**(6), 345-351.

Matheson SL, Shepherd AM, Laurens KR, Carr VJ (2011). A systematic meta-review grading the evidence for non-genetic risk factors and putative antecedents of schizophrenia. *Schizophrenia Research* **133**(1-3), 133-142.

Mullola S, Ravaja N, Lipsanen J, Alatupa S, Hintsanen M, Jokela M, Keltikangas-Järvinen L (2012). Gender differences in teachers' perceptions of students' temperament, educational competence, and teachability. *The British Journal of Educational Psychology* **82**(Pt 2), 185-206.

Munro JC, Russell AJ, Murray RM, Kerwin RW, Jones PB (2002). IQ in childhood psychiatric attendees predicts outcome of later schizophrenia at 21 year follow-up. *Acta Psychiatrica Scandinavica* **106**(2), 139-142.

Myrbakk E, von Tetzchner S (2008). Psychiatric disorders and behavior problems in people with intellectual disability. *Research in Developmental Disabilities* **29**(4), 316-332.

Nelson B, Rawlings D (2010). Relating schizotypy and personality to the phenomenology of creativity. *Schizophrenia Bulletin* **36**(2), 388-399.

Nordentoft M (2007). Prevention of suicide and attempted suicide in Denmark. Epidemiological studies of suicide and intervention studies in selected risk groups. *Danish Medical Bulletin* **54**(4), 306-369.

Paus T, Keshavan M, Giedd JN (2008). Why do many psychiatric disorders emerge during adolescence? *Nature Reviews Neuroscience* **9**(12), 947-957.

Richards R (2001). Creativity and the schizophrenia spectrum: More and more interesting. *Creativity Research Journal* **13**(1), 111-132.

Schulz J, Sundin J, Leask S, Done DJ (2014). Risk of adult schizophrenia and its relationship to childhood IQ in the 1958 British birth cohort. *Schizophrenia Bulletin* **40**(1), 143-151.

Ström K, Hannus-Gullmets B (2015). From special (class) teacher to special educator – the Finnish case. In *Transitions in the field of special education. Theoretical perspectives and implications for practice* (ed. D. L. Cameron and R. Thygesen). Waxmann: Münster.

Sussmann JE, McIntosh AM, Lawrie SM, Johnstone EC (2009). Obstetric complications and mild to moderate intellectual disability. *The British Journal of Psychiatry* **194**(3), 224-8.

Taanila A, Ebeling H, Kotimaa A, Moilanen I, Järvelin MR (2004). Is a large family a protective factor against behavioural and emotional problems at the age of 8 years? *Acta Paediatrica* **93**(4), 508-517.

Therman S, Heinimaa M, Miettunen J, Joukamaa M, Moilanen I, Mäki P, Veijola J (2011). Symptoms associated with psychosis risk in an adolescent birth cohort: Improving questionnaire utility with a multidimensional approach. *Early Intervention in Psychiatry* **5**(4), 343-348.

Verdoux H, van Os J (2002). Psychotic symptoms in non-clinical populations and the continuum of psychosis. *Schizophrenia Research* **54**(1-2), 59-65.

Walker NP, McConville PM, Hunter D, Deary IJ, Whalley LJ (2002). Childhood mental ability and lifetime psychiatric contact: A 66-year follow-up study of the 1932 Scottish mental ability survey. *Intelligence* **30**(3), 233-245.

van der Werf M, Hanssen M, Köhler S, Verkaaik M, Verhey FR, RISE Investigators, van Winkel R, van Os J, Allardyce J (2014). Systematic review and collaborative recalculation of 133,693 incident cases of schizophrenia. *Psychological Medicine* **44**(1), 9–16.

Wiles NJ, Jones GT, Haase AM, Lawlor DA, Macfarlane GJ, Lewis G (2008). Physical activity and emotional problems amongst adolescents: A longitudinal study. *Social Psychiatry and Psychiatric Epidemiology* **43**(10), 765-772.

Zammit S, Allebeck P, David AS, Dalman C, Hemmingsson T, Lundberg I, Lewis G (2004). A longitudinal study of premorbid IQ Score and risk of developing schizophrenia, bipolar disorder, severe depression, and other nonaffective psychoses. *Archives of General Psychiatry* **61**, 354-360.

Table 1. Family characteristics by gender in the Northern Finland Birth Cohort 1986.

	Gende		
	Boys (n=4692)	Girls (n=4415)	p-value ¹
Parental marital status (n=9027)			0.215
- married/cohabiting	4419 (95.2)	4145 (94.6)	
- single/widow/divorced	225 (4.8)	238 (5.4)	
Type of place of residence (n=8990)			0.394
- town	1986 (42.9)	1915 (43.9)	
- village	2638 (57.1)	2451 (56.1)	
Mothers' basic education (n=6511)			0.070
- less than 9 years of comprehensive school	186 (5.7)	153 (4.7)	
- comprehensive school/ elementary school	2036 (62.7)	2018 (61.8)	
- matriculation examination	1026 (31.6)	1092 (33.5)	
Fathers' basic education (n=6213)			0.759
- less than 9 years of comprehensive school	284 (9.2)	297 (9.5)	
- comprehensive school/ elementary school	2204 (71.4)	2247 (71.8)	
- matriculation examination	597 (19.4)	5864 (18.7)	
Parental psychosis (n=9107)			
- no	4515 (96.2)	4254 (96.4)	0.787
- yes	177 (3.8)	161 (3.6)	

¹calculated using Pearson's χ²-test.

Table 2. Learning deficits, special talents and general school success at the age of 8 years by gender in the Northern Finland Birth Cohort 1986.

	Gender				
	Boys	Girls	p-value ¹		
	n (%)	n (%)			
Learning deficits					
- reading (n=8128)	666 (16.1)	352(8.8)	<0.001		
- writing (n=8128)	918 (22.2)	490 (12.3)	<0.001		
- mathematics (n=8123)	380 (9.2)	354 (8.9)	0.603		
Special talents					
- oral presentation,	360 (8.6)	680 (16.9)	<0.001		
acting (n=8212)					
- written presentation	458 (10.9)	895 (22.2)	<0.001		
(n=8212)					
- mathematics (n=8214)	318 (7.6)	127 (3.2)	<0.001		
- athletics (n=8212)	1149 (27.4)	636 (15.8)	<0.001		
- craft (n=8212)	462 (11.0)	996 (24.7)	<0.001		
- drawing (n=8212)	547 (13.1)	1092 (27.1)	< 0.001		
- music (n=8212)	290 (6.9)	872 (21.7)	<0.001		
General school success			0.004		
(n=8173)			<0.001		
- below average	547 (13.1)	278 (6.9)			
- average	2349 (56.4)	2034 (50.7)			
- above average	1266 (30.4)	1699 (42.4)			

¹calculated using Pearson's χ²-test

Table 3. The psychotic-like symptoms (PROD-screen) at the age of 15-16 years according to scholastic traits at age 8 years in the Northern Finland Birth Cohort 1986.

Tallo Horallolli i ililalla Bila			Over 3/12 specific	symptoms in the	PROD screen	
		Boys			Girls	
		Unadjusted	Adjusted OR (95%		Unadjusted	Adjusted OR (95%
	n (%)	OR (95% CI) ¹	CI) ^{1,2}	n (%)	OR (95% CI) ¹	CI) ^{1,2}
Learning deficits						
- reading	89 (20.7)	0.85 (0.66-1.09)	0.85 (0.63-1.14)	85 (33.3)	0.82 (0.63-1.08)	0.83 (0.60-1.13)
- writing	127 (21.1)	0.87 (0.70-1.08)	0.89 (0.69-1.14)	124 (34.9)	0.89 (0.70-1.12)	0.89 (0.68-1.16)
- mathematics	44 (19.5)	0.78 (0.56-1.10)	0.70 (0.46-1.06)	81 (33.8)	0.84 (0.64-1.11)	0.87 (0.63-1.21)
Special talents						
- oral presentation,	83 (30.3)	1.49 (1.14-1.96)	1.49 (1.09-2.03)	225 (42.4)	1.28 (1.06-1.55)	1.23 (1.00-1.52)
acting						
- written presentation	97 (28.0)	1.33 (1.03-1.71)	1.30 (0.98-1.73)	292 (41.2)	1.23 (1.04-1.46)	1.18 (0.97-1.42)
- mathematics	56 (23.3)	1.00 (0.74-1.37)	1.05 (0.75-1.48)	44 (42.7)	1.26 (0.84-1.87)	1.37 (0.89-2.11)
- athletics	176 (20.5)	0.80 (0.66-0.97)	0.72 (0.58-0.90)	193 (38.2)	1.04 (0.86-1.27)	0.98 (0.79-1.22)
- craft	73 (22.3)	0.94 (0.71-1.24)	0.83 (0.60-1.15)	293 (36.5)	0.95 (0.80-1.12)	0.85 (0.71-1.02)
- drawing	115 (28.9)	1.41 (1.11-1.78)	1.44 (1.10-1.87)	348 (40.6)	1.20 (1.02-1.41)	1.13 (0.95-1.36)
- music	57 (25.0)	1.11 (0.81-1.52)	1.14 (0.80-1.62)	272 (38.5)	1.06 (0.89-1.26)	1.06 (0.88-1.29)
General school success						
- below average	65 (19.1)	0.76 (0.57-1.02)	0.68 (0.48-0.97)	63 (34.2)	0.93 (0.67-1.23)	1.04 (0.71-1.52)
- average	390 (23.6)	1.00	1.00	561 (36.0)	1.00	1.00
- above average	231 (24.2)	1.03 (0.86-1.25)	0.96 (0.78-1.19)	531 (39.3)	1.15 (0.99-1.34)	1.22 (1.03-1.44)

¹ Calculated by using binary logistic regression model. CI = confidence interval, OR = Odds Ratio. ²Adjusted for parental marital status, basic education, psychosis and family's place of residence as covariates.

Table 4. The occurrence of psychoses by the age of 30 years according to scholastic traits at age 8 years in the Northern Finland Birth Cohort 1986.

	Boys			Girls			
		Unadjusted	Adjusted		Unadjusted	Adjusted	
	Psychoses	hazard estimate	hazard	Psychoses	hazard estimate	hazard estimate	
			estimate				
	n (%)	HR (95% CI) ¹	HR (95% CI) ^{1,2}	n (%)	HR (95% CI) ¹	HR (95% CI) ^{1,2}	
	113 (2.4)			91 (2.1)			
Learning deficits							
- reading	11 (1.7)	0.68 (0.36-1.27)	0.32 (0.99-	12 (3.4)	1.72 (0.94-3.17)	2.08 (0.97-4.43)	
			1.02)				
- writing	22 (2.4)	1.07 (0.66-1.72)	0.63 (0.30-	14 (2.9)	1.43 (0.81-2.54)	1.44 (0.68-3.08)	
			1.34)				
- mathematics	10 (2.6)	1.14 (0.59-2.19)	0.67 (0.21-	8 (2.3)	1.08 (0.52-2.23)	0.98 (0.35-2.73)	
			2.16)				
Special talents							
- oral presentation.	5 (1.4)	0.56 (0.23-1.39)	0.45 (0.14-	17 (2.5)	1.25 (0.73-2.13)	1.73 (0.94-3.20)	
	3 (1.4)	0.30 (0.23-1.39)	1.44)	17 (2.3)	1.23 (0.73-2.13)	1.73 (0.94-3.20)	
acting	11 (2.4)	1 02 (0 54 1 00)	•	26 (2.0)	1 57 (0 00 2 50)	1 25 (0 69 2 22)	
- written	11 (2.4)	1.02 (0.54-1.90)	0.93 (0.44-	26 (2.9)	1.57 (0.99-2.50)	1.25 (0.68-2.33)	
presentation	F (1.6)	0.65 (0.36.4.50)	1.97)	2 (2 4)	1 14 (0 26 2 60)	1 16 (0 00 1 76)	
- mathematics	5 (1.6)	0.65 (0.26-1.59)	0.53 (0.17-	3 (2.4)	1.14 (0.36-3.60)	1.16 (0.28-4.76)	
- athletics	24 (4.9)	0.74 (0.44.4.44)	1.68)	11 (1 7)	0.00 (0.42.4.54)	0.04 (0.44.2.04)	
- amencs	21 (1.8)	0.71 (0.44-1.14)	0.77 (0.43-	11 (1.7)	0.80 (0.43-1.51)	0.94 (0.44-2.01)	
oroft	10 (2.2)	0.04 (0.07.4.76)	1.37)	20 (2.0)	0.76 (0.40.4.27)	0.72 (0.25 4.52)	
- craft	10 (2.2)	0.81 (0.37-1.76)	1.18 (0.50-	20 (2.0)	0.76 (0.42-1.37)	0.73 (0.35-1.53)	
duranta a	47 (0.4)	4 00 (0 00 0 00)	2.80)	00 (0.0)	4.05 (0.00.0.40)	0.00 (0.54.4.04)	
- drawing	17 (3.1)	1.38 (0.82-2.32)	1.28 (0.67-	28 (2.6)	1.35 (0.86-2.12)	0.99 (0.54-1.84)	
	5 /4 3\	0.74 (0.00 4.75)	2.45)	40 (0.0)	4 00 (0 00 4 70)	4 00 (0 75 0 50)	
- music	5 (1.7)	0.71 (0.30-1.75)	0.98 (0.39-	19 (2.2)	1.06 (0.63-1.76)	1.38 (0.75-2.53)	
			2.46)				
General school	00 (4 7)			70 (4.0)			
success	80 (1.7)			70 (1.6)			
- below average	11 (2.0)	0.88 (0.46-1.68)	0.74 (0.29-	13 (4.7)	2.19 (1.18-4.06)	1.80 (0.74-4.35)	
			1.90)				
- average	54 (2.3)	1.00	1.00	44 (2.2)	1.00	1.00	
- above average	32 (2.5)	1.10 (0.71-1.70)	0.97 (0.56-	26 (1.5)	0.70 (0.43-1.14)	0.60 (0.32-1.11)	
			1.68)				

¹Calculated by using Cox regression model. CI = confidence interval, HR = Hazard Ratio. ²Adjusted for parental marital status, basic education, psychosis and family's place of residence as covariates.

Supplement Table 1 Availability of teacher assessments at age 8 years according to family background factors² in the Northern Finland Birth Cohort 1986.

	Availability of teacher assessment at age 7-8 years					
	Boys			Girls		
	Available	Missing	p-value ¹	Available	Missing	p-value ¹
	n (%)	n (%)	-	n (%)	n (%)	
Family characteristics						
Parental marital status (n=8955)			0.015			>0.999
- married/cohabiting	3981 (91.3)	379 (8.7)		3808 (91.9)	337 (8.1)	
- single/widow/divorced	183 (86.3)	29 (13.7)		219 (92.0)	19 (8.0)	
Type of place of residence (n=8919)			0.013			0.371
- town	1757 (90.0)	195 (10.0)		1752 (91.5)	163 (8.5)	
- village	2396 (92.1)	205 (7.9)		2261 (92.2)	190 (7.8)	
Mothers' basic education			0.341			0.091
(n=6510)						
 less than 9 years of comprehensive school 	173 (93.0)	13 (7.0)		143 (93.5)	10 (6.5)	
 comprehensive school/ elementary school 	1896 (93.2)	139 (6.8)		1872 (92.8)	146 (7.2)	
- matriculation examination	941 (91.7)	85 (8.3)		1035 (94.8)	57 (5.2)	
Fathers' basic education (n=6211)			0.010			0.928
- less than 9 years of comprehensive school	266 (93.7)	18 (6.3)		279 (93.9)	18 (6.1)	
 comprehensive school/ elementary school 	2056 (93.3)	147 (6.7)		2105 (93.7)	142 (6.3)	
- matriculation examination	535 (89.8)	61 (10.2)		545 (93.3)	39 (6.7)	
Parental psychosis (n=9009)	145 (85.8)	24 (14.2)	0.020	147 (91.3)	14 (8.7)	>0.999

¹ Calculated using Pearson's χ²-test,

² Subjects who moved or died before the assessment not included

Supplement Table 2. Availability of PROD-screen results at age 15-16 years according to family characteristics² in the Northern Finland Birth Cohort 1986.

	Availability of PROD-screen by gender					
	Boys			Girls		
	Available	Missing	p-value ¹	Available	Missing	p-value ¹
	n (%)	n (%)	_	n (%)	n (%)	
Family characteristics						
Parental marital status			0.011			0.006
(n=8955)						
- married/cohabiting	3093 (70.9)	1267 (29.1)		3181 (76.7)	964 (23.3)	
- single/widow/divorced	133 (62.7)	79 (37.3)		164 (68.9)	74 (31.1)	
Type of place of residence (n=8919)			0.008			0.053
- town	1337 (68.5)	615 (31.5)		1434 (74.9)	481 (25.1)	
- village	1876 (72.1)	725 (27.9)		1898 (77.4)	553 (22.6)	
Mothers' basic education			0.017			0.422
(n=6510)			0.017			0.123
- less than 9 years of comprehensive school	148 (79.6)	38 (20.4)		135 (88.2)	18 (11.8)	
- comprehensive school/ elementary school	1705 (83.8)	330 (16.2)		1755 (87.0)	263 (13.0)	
- matriculation examination	890 (86.7)	136 (13.3)		977 (89.5)	115 (10.5)	
Fathers' basic education			0.095			0.902
(n=6211)						
 less than 9 years of comprehensive school 	231 (81.3)	53 (18.7)		262 (88.2)	35 (11.8)	
 comprehensive school/ elementary school 	1869 (84.8)	334 (15.2)		1985 (88.3)	262 (1170)	
- matriculation examination	518 (86.9)	78 (13.1)		512 (87.7)	72 (12.3)	
Parental psychosis (n=9009)	100 (59.2)	69 (40.8)	0.001	116 (72.0)	45 (28.0)	0.260

¹Calculated using Pearson's χ²-test,

² Subjects who moved or died before the assessment not included