

Article

Early Risk Indicators for DSM-IV Diagnoses in Adolescents and Young Adults with Intellectual Disabilities

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ABSTRACT: To identify risk indicators at ages 6–18 years that are associated with DSM-IV diagnoses in adolescents and young adults with intellectual disabilities five years later. To assess the potential health gain and efficiency of preventive interventions targeting these risk indicators. Parents reported on potential child, parental, and environmental risk indicators. Five years later, parents were interviewed using a standardised psychiatric interview schedule (DISC-IV) to assess DSM-IV diagnoses in children with ID (N = 614) at the age of 11 to 24 years. Logistic regression and linear probability models were used to test the contribution of risk indicators to the prediction of DSM-IV diagnoses. Deviant levels of internalising and externalising problems, inadequate adaptive behaviour, and parental psychopathology predicted psychiatric disorder. Children/adolescents exposed to multiple risk indicators were at greater risk of developing DSM-IV disorders. Strategies aiming for the risk reduction of psychiatric disorders in children/adolescents with ID should focus on intervening at an early age, improving psychopathology and adaptive behaviour skills of the children/adolescents, and supporting their parents.

Keywords: Intellectual Disabilities; DSM-IV diagnoses; Risk indicators; Prevention; Health gain



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1. Introduction

In the last decennia, various studies have reported high prevalence of psychiatric disorders in children/adolescents, young adults and adults with intellectual disabilities (ID) [1–13]. For example, a study by Dekker and Koot [4] showed that 21.9% of the 474 children and adolescents with ID in their sample met the DSM-IV symptom criteria for anxiety disorder, 4.4% for mood disorder, and 25.1% for disruptive disorder. More than half of those meeting the criteria for a DSM-IV disorder were severely impaired in everyday functioning, and about 37% had a comorbid disorder. Further, studies also reported high stability of psychopathology from childhood into adulthood in individuals with ID [14]. Therefore, it seems essential to prevent the development of these problems as early as possible. However, so far, only a few studies addressed risk factors for psychiatric disorders in youth with ID using a prospective design. Therefore, hardly anything is known on factors to be targeted in efforts to prevent these disorders. The goals of this study were to identify risk indicators present at ages 6–18 associated with DSM-IV diagnoses five years later and to assess the potential health gain and efficiency of preventive interventions targeting these risk indicators.

The development of psychopathology in youths with ID in the context of risk may be understood from a general developmental framework. The bioecological model of human development [15] offers a holistic approach to analysing the influences of child-related and environmental factors on the child's development. According to this model, factors at the microsystem level, including factors at the family level, have the most influence on the child, next to influences from child characteristics. We assume that mechanisms and factors explaining the development of children and adolescents from the general population apply equally to populations of youths with ID and equally apply to the

development of psychiatric problems. For example, studies on risk and adversity in developmental psychopathology [16] have identified factors like previous psychopathology of the child, (chronic) physical condition, learning problems, stressful life events, maternal psychopathology, family dysfunction, and low socioeconomic status as risk factors. Over and above these, limitations in cognitive and social skills may pose additional risk for children and adolescents with ID. This study focuses on each of these factors.

Studies in the general population have found elevated levels of emotional and behavioural problems and psychiatric disorders in childhood to be predictive of the onset and continuity of psychiatric disorders in adolescence and adulthood [17]. Besides childhood psychopathology, also gender, social competence, self-esteem, chronic physical illness of the child or parent, socio-economical status, ethnicity, early adverse life-experiences, and family structure and functioning have been found to be prospectively associated with psychiatric disorders [17–23].

While the co-occurrence of ID and mental health problems in children and adolescents has been well-documented [6,24,25], very little research has addressed the association between a range of risk factors and mental health issues in this population. Most studies address the association of specific experiences (e.g., maltreatment) with specific outcomes (e.g., depression), although some efforts are made for broader perspectives using cross-sectional study designs. For example, Weiss, Ting, and Perry [26] found that youth with a psychiatric diagnosis had higher levels of adaptive behavior and experienced more negative life events than youth without a psychiatric diagnosis, while the presence of clinically significant maladaptive behavior was related to higher levels of adaptive behavior, parents' mental health problems and lower family quality of life. Whitney et al. [27] showed that co-morbid conditions, physical and social factors, autism spectrum disorders, Down syndrome, attention-deficit/hyperactivity disorder, pain, and bully victimisation were each associated with depression and/or anxiety problems. However, very little has been done to address prospective relations between risk and mental health outcomes in this population, and in fact, many longitudinal epidemiological studies of prevalence, risk factors, and continuity/discontinuity of child and adolescent mental disorders omitted study participants with ID/IDD [28].

Only a few studies identified factors associated with psychiatric disorders among people with ID. In a study among children and adolescents with mild to moderate ID, apart from initial high levels of problem behaviour, child low social competence, inadequate daily living skills, child health problems, negative life events, parental psychopathology, and poor family functioning were found to be predictive of DSM-IV diagnoses one year later [29]. Significant associations between psychopathology and the experience of negative life events were demonstrated in cross-sectional studies among adults [30,31] and children with ID [32]. A longitudinal study by McCarthy and Boyd [33] among people with Down's syndrome, however, found no significant associations between childhood levels of adaptive behaviour, or family factors like social class or parental psychopathology, and psychiatric disorders in adulthood.

The studies described above successfully identified high-risk groups of children but also had several limitations. They predominantly focussed on clinical samples, samples of children or adults with ID in residential settings, or people with a specific syndrome, and most studies only looked at cross-sectional or short term longitudinal associations between psychopathology and child/adolescent risk factors. The first aim of this study is to identify child/adolescent risk indicators of psychiatric problems of individuals identified in adolescence or young adulthood. The study takes a prospective longitudinal approach tracking children and adolescents with ID across a five-year period. The transitional periods from childhood to adolescence and adolescence to young adulthood are of special interest as these periods put increasing demands on these individuals and their families in several important life domains, including social relations, education, and possibly entrance into the labour market. Especially normative demands involving independence and participation in society require more adaptation than may be expected given the limitations of young people with ID. Across these periods, in the general population, serious symptoms of depression and conduct tend to rise (specifically in females and males, respectively), and other problems (e.g., anxiety) tend to grow into more serious forms (e.g., from separation anxiety into social anxiety). However, the extant knowledge on the contribution of personal and family factors to the development of psychiatric problems in children and adolescents with ID across the transition to adolescence and young adulthood, respectively, is seriously wanting.

As demonstrated by several studies in clinical as well as general population samples, child/adolescent psychopathology tends to be fairly stable, and psychiatric disorders show fairly high levels of continuity, also in individuals with ID [14]. Therefore, there is a strong need to prevent the development of psychiatric problems in children/adolescents with ID and identify malleable risk factors related to this development. However, we are unaware of any study that empirically addressed that need. From a prevention perspective, several indices are informative about the potency of reduction of psychiatric problems when specific risk factors are tackled successfully. First, the *impact* of risk factors on the development of psychiatric disorders tells us how big the chance is for a child who is exposed to a

certain risk factor to develop a certain psychiatric disorder. Exposure to some risk factors might have a larger impact than exposure to other risk factors. Risk factors with a larger impact should be the focus of preventive strategies. The impact of risk factors can be described by the risk difference (RD) index, which is the difference in the risk of developing a psychiatric problem between exposed and unexposed children/adolescents. Its inverse, the number needed to be treated (NNT) indicates how many children/adolescents would have to receive a preventive intervention to avoid one new case with psychiatric problems, assuming that the adverse effect of the risk factor can be completely blocked by some preventive intervention. The NNT is thus an indicator of the maximum *efficiency* of the intervention. To assess the *potential health gain* in a population the incidence rate ratio (IRR) and the attributable fraction (AF) are useful. The IRR is the ratio of the incidence rate of psychiatric problems in a group of exposed children over the incidence rate in the unexposed group. The IRR thus captures the strength of association between an exposure and outcome, not unlike a relative risk or an odds ratio. Finally, the AF reflects the percentage by which the incidence rate of psychiatric problems can be reduced when the adverse effect of a risk factor is eliminated by a preventive intervention. Higher RDs, IRRs, and AFs, and lower NNTs point to greater utility of the studied risk indicators for preventive interventions because then we may expect that substantial health gains can be generated efficiently.

The second aim of this study is to evaluate the impact, efficiency, and possible health gain of addressing risk factors associated with psychiatric problems in adolescents and young adults with ID. This epidemiological approach recently introduced in the area of adult depression [34,35] and applied in the field of childhood internalising problems [36], was expected to throw some light on potential preventive intervention targets related to these psychiatric problems.

Not only is it necessary to identify risk indicators for the development of psychiatric disorders and their relevance in preventing psychiatric disorders when addressed, but also to know the cumulative effect of these relevant risk indicators. From studies in the general population, it is known that the accumulation of risk factors, such as poor family functioning, high levels of parental stress and low family income, is exponentially related to the development of psychiatric disorders [37,38]. The severity of and impairment from psychiatric disorders is higher in families that experience negative influences in more than one area [32]. Children with two or more co-morbid psychiatric disorders and their families are also more likely to have other associated developmental, social, and educational needs [39]. Following this lead, young people with ID might be considered as a multi-risk group with complex and interrelated mental health needs over and above their intellectual disabilities. Therefore, the third aim of this was to determine the potential cumulative effect of reduction of all significant risk indicators in preventing psychiatric disorders in adolescents and young adults with ID.

To summarise, the present longitudinal study aimed, first, to identify risk indicators at ages 6 to 18 years that are predictive of psychiatric DSM-IV disorder 5-years later. Given the age and gender-sensitive development of specific psychiatric problems across adolescence and young adulthood, these two factors were considered moderators of the effects of potential risk indicators. Second, we aimed to contribute to developing effective preventive interventions by identifying the risk indicators associated with the largest potential health gain from preventive interventions. Third, we aimed to improve knowledge about the cumulative effect of risk indicators. This approach, seeking to identify high-risk groups of children with ID where preventive interventions are most likely to generate the largest health gains for the least effort, might yield important information relevant to the prevention of psychiatric symptoms in adolescents and young adults with ID.

2. Method

2.1. Procedure and Participants

The present study used the first (indicated as Time 1, T1) and the third (Time 3, T3) parent assessments of a large random school-based study on psychopathology in Dutch children and adolescents aged 6–18 years at T1, with moderate to borderline ID. At T1 for 891 of 1396 children/adolescents eligible for this study, information for all addressed risk indicators was collected (response rate of T1 participants: 63.8%) (for more details on T1, see [4,29,40]). About 5 years after T1, 614 parents (T3 response rate: 68.8% of the participants at T1) were interviewed using the Diagnostic Interview Schedule for Children—Parent version (DISC-IV [41–43], after signing an informed consent form. For sample characteristics, see Table 1. We compared children/adolescents who participated at both times, excluding those who had incomplete data due to missing risk indicator information and/or missing DISC-IV interview data. No significant differences were found for age, sex, or deviant (i.e., above the borderline cut-off) CBCL total problem scores. However, children/adolescents who did not participate at both times more often came from families with low SES (versus medium or high SES, percentages: 69.9% vs. 47.7%, OR = 2.5, $p < 0.001$). They more often attended a school for

children/adolescents with moderate ID (versus a school for children/adolescents with mild ID, percentages: 34.4% vs. 25.7%, OR = 1.5, $p < 0.001$) than children/adolescents who participated both at T1 and T3.

2.2. Measures

2.2.1. Putative Risk Indicators

A description of putative risk indicators included in this study can be found in Table 1.

Table 1. Description of child and family risk indicators measured at T1 (N = 614).

Risk Indicators	Description of Instruments and Variables	Risk Group (Coded 1)	Cronbach's α
Child			
Sex	39.6% females, 60.4% males	Female	
Age	51.1% 6 to 12-year-olds 49.9% 13 to 18-year-olds Mean age 11.54 (SD:3.02)	6 to 12 year-olds	
Educational level	34.2% attended a school for the moderate ID (IQ < 60) 65.8% attended a school for the mild ID (IQ 60–80)	Moderate ID	
High physical symptoms	Wahler Physical Symptom Checklist (WPSI) [44]; 6-point scale (almost never to almost every day) assessing 42 physical symptoms (e.g., pains, paralysis, nausea)	1 SD above mean	0.83
Inadequate adaptive behaviour	Vineland Screener [45]; 45-item short version of the Vineland Adaptive Behaviours Scales [46]	Standard score < 85	0.87
Inadequate daily living skills			0.89
Inadequate communication			0.78
Inadequate socialization			
Emotional/ behavioural problems	Children's emotional and behavioural problems (past 6 months) were assessed using the Child Behavior Checklist (CBCL) [47,48]	CBCL <i>T</i> scores: Internalising score \geq 60, Externalising score \geq 60	0.86 0.91
Family			
Single parent	Not married/cohabiting at T1 versus non-single parent	Single parent	
Socioeconomic status	Occupational level of highest rated parent [49]; low socioeconomic status (SES) includes unemployed, unskilled workers and workers with lower vocational training; and medium/high SES includes jobs at higher levels	Medium of high SES	
Ethnicity	Both (or single) parents Dutch versus \geq 1 parent born outside the Netherlands	Dutch parents	
Family dysfunction	12-item General Functioning subscale of the McMaster Family Assessment Device [50]	Mean item score > 2.17	0.85
Parental psychopathology	Young Adult Self-Report (YASR) [51]; 29-item version [52] representing recent internalising problems of the primary caregiver.	1 SD above mean of normative Dutch sample [53]	0.92
Sibling referral to mental health care	\geq 1 sibling referred to mental health care ever	Present	
Environment			
Negative life events	Child negatively affected by \geq 1 life event in the past 2 years (e.g., parent leaving the house, 20% decline in income) using 16-item adjusted version of the Life Events Questionnaire [54]	>1 negative life events	

2.2.2. DSM-IV Disorders

To assess the presence of any DSM-IV anxiety, mood, or disruptive disorder in the past year, the main caregiver was interviewed at T3 using the Dutch version of the DISC-IV [41–43]. The impairment criteria were not taken into account. The DISC-IV was administered by trained lay interviewers.

2.2.3. Data Analysis

Univariate logistic regression analysis of each DSM-IV on each predictor was performed using SPSS version 12. Moderating effects of age and sex were tested separately by adding each variable and its interaction term with the main predictor variable to the model. If the model chi-square improved significantly, post hoc probing was performed to determine the significance and strength of the association between the main predictor variable and DSM-IV disorder within each moderator level. Next, linear probability models were tested using STATA [55] in three steps. In the first step, all predictor variables were entered into a backward stepping linear probability model. This model served to identify risk indicators with a significant contribution to the prediction of DSM-IV disorders. This step was carried out twice, first without including Internalising and Externalising, and second with Internalising and Externalising included as predictor variables. In the second step, the significant risk indicators from the previous step were used in parsimonious models to determine the potential relevance of the risk indicators in efforts to prevent DSM-IV disorders. The risk difference (RD) associated with each factor was obtained under a linear probability model and numbers to be treated (NNT) as a function of the RDs ($NNT = 1/RD$). In addition, incidence rate ratios (IRR) and attributable fractions (AF) were obtained under a Poisson regression model with the help of STATA's Aflogit procedure [56]. In the third step, the possible cumulative effect of the risk indicators was investigated by calculating a risk index. This risk index represents the sum of the risk indicators that were found to have a significant OR. Since there were few children with 5 or more risks, the presence of 5, 6, 7 or 8 risks was combined into a new category "5 or more risks present" for Any DSM-IV, Any anxiety and Any disruptive disorder. Likewise, the category "3 or more" was created for Any mood disorder. In a next step, the risk indexes were used as predictors of DSM-IV disorders. The dose-response nature of the relationship between the risk index and the outcome was investigated using Kendall's Tau.

3. Results

3.1. Association between Risk Indicators and DISC-IV Disorders

Table 2 shows the percentage of children exposed to each risk indicator, and the significant Odds Ratios within each level of the moderator variables age and sex whenever a significant interaction effect was present. All included factors except child educational level (mild vs moderate ID), single parenthood, and mental health problems in siblings were significantly related to one or more of the psychiatric disorder clusters. Only four significant interaction terms were found inconsistent across the four clusters.

Table 2. Percentages of children exposed to risk indicators at T1.

Risk indicators	Complete Sample	Any DISC-IV Disorder		Any Anxiety Disorder		Any Mood Disorder		Any Disruptive Disorder	
	% children (N = 614)	% children (N = 177)	OR (95% CI)	% children (N = 109)	OR (95% CI)	% children (N = 20)	OR (95% CI)	% children (N = 96)	OR (95% CI)
Sex (Female)	39.6	43.5		49.5	1.6 (1.1–2.5)	60.0	2.4 (1.0–5.9)	33.3	
Age (<13 years)	51.1	56.5		53.2		30.0	0.4 (0.2–1.1)	63.5	1.8 (1.2–2.9)
Educational level (Moderate ID)	34.2	37.3		41.3		30.0		35.4	
Physical symptoms	10.6	15.8	2.0 (1.2–3.4)	17.4	2.1 (1.2–3.8)	20.0		16.7	1.9 (1.0–3.5)
Inadequate Daily living skills	71.5	80.2	1.9 (1.3–2.9)	80.7	1.8 (1.1–3.1)	75.0		79.2	1.6 (1.0–2.8)
Inadequate Communication skills	88.8	93.2	2.1 (1.1–4.0)	93.6	2.0 (0.9–4.6)	90.0		93.8	
Inadequate Socialisation	74.3	81.9	1.8 (1.2–2.8)	81.7	1.7 (1.0–2.8)	75.0		84.4	2.1 (1.2–3.7)
Internalising problems	39.7	53.1	2.2 (1.5–3.1)	51.4	1.8 (1.2–2.7)	60.0	2.3 (1.0–5.8)	61.5	2.9 (1.8–4.5)
Externalising problems	38.3	54.2	2.5 (1.8–3.6)	51.4	1.9 (1.3–2.9)	50.0		65.6	3.8 (2.4–6.1)
Single parent	10.3	8.5		11.0		10.0		6.3	
Medium or high ses	51.8	58.2	1.4 (1.0–2.0)	59.6	1.5 (1.0–2.2)	55.0		56.3	2.1 ^Y (1.2–3.7)
Dutch parents	87.8	88.7	2.1 ^Y (1.0–4.6)	85.3		95.0		94.8	13.0 ^Y (1.8–96.2)
Family dysfunction	12.1	13.6		11.9		20.0		18.8	1.9 (1.1–3.4)
Parental psychopathology	12.5	14.1	2.4 ^F (1.2–4.8)	14.7		30.0	3.2 (1.2–8.5)	13.5	2.6 ^F (1.1–6.1)
Mental health problems siblings	3.4	4.0		3.7		5.0		4.2	
Negative life events >1	20.2	20.3		17.4		30.0		24.0	2.7 ^F (1.2–6.1)

Note: Only significant ($p < 0.05$) Odds Ratios (OR) are presented. Odds ratios resulting from post hoc probing of significant interaction effects ($p < 0.05$) of predictors with age (Y = 6–12 years at T1) or sex (F = female) are reported within each level of the moderator.

3.2. Parsimonious Models: Identifying the Most Relevant Risk Indicators

After entering all significant risk indicators simultaneously in regression analyses several sets of indicators appeared as significant ($p < 0.05$) but independent contributors to the prediction of each DSM-IV disorder. For any DSM-IV disorder, the independent risk indicators appearing from these analyses were internalising, externalising, inadequate communication skills, and medium or high SES. For Any anxiety disorder, female sex, externalising problems, inadequate communication skills, and medium or high SES were the most relevant risk indicators. Any mood disorder was only significantly related to deviant levels of parental psychopathology. Finally, ages 6–12, internalising and externalising problems and having Dutch parents were the most relevant risk indicators for Any disruptive disorder.

3.3. Relevance of the Risk Indicators in Preventing DSM-IV Disorders

Table 3 presents the RDs, NNTs, IRRs and AFs of the parsimonious models described above. The risk indicators of the parsimonious models resulted in total AFs of 86.1%, 69.8%, 20.0%, and 98.6%, indicating the percentage of the cases of any DSM-IV, any anxiety, any mood and any disruptive disorder, respectively, that can be identified with these risk profiles.

Higher RD, IRR and AF and lower NNT indicate that a risk indicator is important for targeting preventive interventions. We use the first row of Table 3, corresponding to inadequate level communication skills, as an illustration to explain the findings reported in Table 3. The RD suggests a difference of 12% in the likelihood of having any DSM-IV disorder at T3 between children with inadequate and children with adequate communication skills at T1. The NNT of 8.3 indicates that 9 children would have to receive an intervention blocking the adverse effect of inadequate communication skills, in order to avoid one case with DSM-IV disorders, under the presumption that the intervention is completely successful in containing the adverse effect of this risk indicator. The IRR, which can be interpreted as an odds ratio, shows the increase in the probability of any DSM-IV disorder among the exposed group (children with inadequate communication skills) relative to the unexposed group (children with adequate communication skills). The AF value of 39.7% indicates that of all adolescents/young adults with any DSM-IV disorder at T3, 39.7% can be ascribed to inadequate communication skills at T1. In other words, if all children with inadequate communication skills at T1 had received an effective intervention to improve their communication skills, the prevalence of any DSM-IV disorder at T3 could have been reduced by 39.7%. The entries for the other risk indicators in Table 3 should be interpreted similarly.

Table 3. Parsimonious models based on the variables obtained from backward-stepping linear probability model.

Parsimonious Models	RD	95% CI	NNT	IRR	95% CI	AF (%)	95% CI	
Any Disc-IV disorder								
Inadequate communication skills	0.12	0.01–0.23 *	8.3	1.74	0.96–3.15	39.7	7.4–60.8	
Internalising CBCL	0.10	0.02–0.18 *	10	1.40	0.99–1.97 *	15.3	5.3–24.2	
Externalising CBCL	0.16	0.08–0.24 ***	6.3	1.70	1.22–2.38 **	22.4	12.9–30.9	
Medium or high SES	0.10	0.03–0.17 **	10	1.42	1.05–1.92 *	17.2	6.9–26.4	
Dutch parents and age <13 years	0.08	0.01–0.15 *	12.5	1.30	0.97–1.75	12.0	3.0–20.2	
No parental psychopathology	0.20	0.05–0.34 **	5	2.13	0.99–4.60	45.6	10.9–66.8	
Deviant parental psychopathology and female sex	0.32	0.12–0.51 **	3.1	2.92	1.22–7.02 *	6.7	3.6–9.7	
						Total AF	86.1	64.5–94.6
Any Anxiety disorder								
Female sex	0.07	0.01–0.13 *	14.3	1.48	1.02–2.16 *	16.2	2.9–27.6	
Inadequate communication skills	0.09	–0.01–0.18	11.1	1.86	0.87–4.01	43.4	–6.3–69.8	
Externalising CBCL	0.10	0.04–0.16 **	10	1.72	1.18–2.52 **	21.6	8.5–32.8	
Medium or high SES	0.07	0.01–0.13 *	16.7	1.46	0.99–2.14	18.7	2.2–32.5	
						Total AF	69.8	36.0–85.8
Any Mood disorder								
Deviant parental psychopathology	0.05	0.01–0.09 *	20	2.99	1.15–7.78 *	20.0	–4.8–38.9	
						Total AF	20.0	–4.8–38.9
Any Disruptive disorder								
Male sex	0.07	0.01–0.13 *	14.3	1.63	1.01–2.64 *	25.8	5.7–41.7	
Age > 12 years	0.13	0.03–0.24 *	7.7	6.84	0.93–50.32	31.1	26.5–43.2	
Internalising CBCL	0.09	0.02–0.15 **	11.1	1.59	0.99–2.56	22.8	5.0–37.2	
Externalising CBCL	0.16	0.10–0.22 ***	6.3	2.70	1.67–4.37 ***	41.3	25.8–53.7	
Medium or high ses and age < 13 years	0.11	0.03–0.19 **	9.1	1.84	1.09–3.09 *	17.6	6.9–27.0	
Dutch parents and age <13 years	0.18	0.07–0.29 ***	5.6	8.91	1.23–64.71 *	55.5	36.1–69.0	
No parental psychopathology	0.17	0.06–0.29 **	5.9	2.92	1.05–8.11 *	56.8	15.1–78.1	
Deviant parental psychopathology and female sex	0.22	0.06–0.39 **	4.5	4.19	1.17–14.96 *	7.1	2.4–11.6	
						Total AF	98.6	86.8–99.9

Note: RD = risk difference, CI = confidence interval, NNT = numbers needed to be treated, IRR = incidence rate ratio, AF = attributable fraction, * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$.

3.4. Cumulative Effects

The probability of having any DSM-IV disorder increased with the number of joint exposures to risk indicators; from 0.0% when having 0 risk indicators to 45.3% when having 5 or more risk indicators. When the significant risk indicators for any or a specific disorder were combined in a risk index and used as a single predictor of the outcome, the AFs attributable to this index were 80.2%, 84.0%, 44.6%, and 79.9% for any DSM-IV disorder, any anxiety disorder, any mood disorder, and any disruptive disorder, respectively. This indicates that approximately 80% of most DSM-IV disorders (except mood disorders) can be prevented when the negative effect of the combined risk indicators factors could be blocked. Accumulative exposures were significantly related to an increase in the risk for disorders (Kendall's Tau = 0.25, 0.18, 0.06, and 0.17 for any DSM-IV disorder, any anxiety disorder, any mood disorder, and any disruptive disorder, respectively).

4. Discussion

The main findings of this study are threefold. First, we found that not only childhood/adolescence levels of psychopathology but also child factors like inadequate adaptive functioning and family factors like parental psychopathology are important risk indicators for DSM-IV disorders in young people with ID who all after entering adolescence or young adulthood. Second, we found disorder specific Time 1 risk profiles, to which large proportions of the psychiatric problems in this sample identified 5 years later could be attributed. Third, similar to findings from the general population of children and adolescents, we found that the accumulation of risk factors increases the chances of developing a DSM-IV disorder, in this case, 5 years later. In the following section, we will discuss these findings in more detail.

The present study showed that when children have matured into adolescence or young adulthood, their levels of risk indicators from the child and family domain assessed at childhood or adolescence are still predictively associated with developed DSM-IV disorders. This extends the findings of Dekker and Koot [29], who looked into the 1-year predictive value of these child and family factors for the development of DSM-IV disorders in children and adolescents with ID using part of this same sample. In our study, independent and important risk indicators at the child level were internalising and externalising problems, inadequate communication skills, and at the family level and parental psychopathology. The previous study showed that the strongest 1-year predictive associations with DSM-IV disorders came from child social incompetence, inadequate daily living skills, health problems, emotional and behavioral problems, parental mental health problems and parent-reported negative life events. Apparently, next to child behavior problems, child/adolescent communication skills gain importance for predictive significance over this longer period. The risk indicators for psychopathology identified in this study are quite similar to those found for children and adolescents in the general population. In addition, inadequate daily living skills, specifically, inadequate communication skills, appeared as important predictors. It seems that with the increasing demands for independent functioning encountered during adolescence, these factors come to play a significant role in the development of mental health problems.

We found that parental psychopathology was the only significant risk indicator for mood disorders and that this factor alone can identify 20% of the cases of mood disorders in young people with ID. Apart from the important risk indicators included in this study, there might be other factors (e.g., genetic vulnerability and expressed emotion in the family [31,57]) that are valuable risk indicators of psychiatric disorders too. It is quite likely that youths with ID are exposed more often to parental psychopathology than youths from the general population. We know that the likelihood of mental health problems is strongly elevated in people with ID and that a sizeable portion of parents of children with ID are affected by borderline to mild ID themselves.

Although, in accordance with what might be expected, females were more at risk for anxiety and mood disorders, and younger individuals more often had a disruptive disorder, very few moderation effects of gender and age were found. A significant finding was that females with ID were more affected by parental psychopathology and life events, specifically in the prediction of disruptive disorder.

A strong finding was that accumulation of risk factors was highly predictive of later psychiatric problems. The effect of the presence of multiple risks as opposed to a single risk in a child on the development of DSM-IV disorders had been found before in general population studies [58]. These studies showed that it is not the *nature* of risk factors but rather their *number* that is the best predictor of later outcome. More specifically, the more risk indicators present in the child and its family, the bigger the risk of psychopathology. In a study into risk indicators in children aged 2-3 years, it was shown that children who had a cumulative risk index of 3 or more risks were at a 50 times higher risk of problem behaviour [59]. Vervoort-Schel et al. [60] found that about 20% of the children with moderate and mild ID experienced

4 adverse childhood experiences or more. Many of their families faced multiple and complex problems. This pattern of cumulative incidence and effect was also found in adults with ID for the influence of life events on the development of psychopathology [32]. Some risk indicators (e.g., inadequate communication skills) are present in the majority (88.8%) of the sample. This makes the chance of having 2 or 3 risk indicators high, which might explain why the probability of disorder is especially high when the cumulative risk is 3 or more. The high percentage of multiple risk indicators in people with ID might partly explain the high prevalence of psychopathology compared to the general population, as found in many studies [1,5]. Screening for the cumulative risk status of children and adolescents with mild or moderate ID might enhance early detection of developing psychiatric problems at later ages.

The results of this study have potential meaning for the choice of targets for preventive interventions. For example, our findings on the influence of inadequate communication skills suggest that if all children with inadequate communication skills at T1 had received an effective intervention to improve their communication skills to an adequate level, the prevalence of any DSM-IV disorder at T3 could have been reduced by almost 40%. The attributable fractions (AF) for the several risk profiles established in this study (except for mood disorder) suggest almost full prevention options. However, only a few of the factors included in these profiles seem malleable and, therefore, the most likely options for targeting preventive interventions. These include child inadequate communication skills and both internalising and externalising problems next to parental psychopathology. That being said, the prediction models employed in this study are insufficient to imply interventions. Not only do the estimates in the parsimonious models fairly wide confidence intervals, but more importantly, no intervention is completely effective whatever actionable predictor is addressed. Randomized trials are indispensable to establishing true effectiveness.

4.1. Limitations

The present study is not without limitations. First, the prevalence of mood disorders was low in our sample; therefore, we should interpret the results carefully. Future studies should focus on a larger sample of young people with ID and mood disorders in order to find out if sex and internalising problems in childhood should really not be included in a risk profile. Moreover, although not uncommon for this type of study, the non-response across waves of measurement in this study was rather large, and drop-out was not random, which may have influenced the reliability of our findings. Secondly, the diagnostic interview used in this study was designed for people without ID. It might be that people with ID express symptoms of disorders in a different manner than people without ID (e.g., expressing feelings of depression requires the ability to communicate on a high level). Also, some symptoms of depressive feelings, like having trouble focusing on (school)work, are inherent to the ID and might therefore, not be reported by their parents. Thirdly, predictions as obtained in this study might be influenced by unmeasured instability of the predictors across time up until the assessment of the outcome. For example, Erickson, Sroufe, and Egeland (1985) found a relationship between quality of attachment at 12 and 18 months and behavior problems in preschool, but much more so when attachment quality was stable.

4.2. Implications

This study shows that this group is vulnerable in multiple ways. In order to be able to prevent DSM-IV disorders in young people with ID, we need to focus on all the different areas in which these children experience negative influences and not on one single problem at the time. Parents, caregivers, teachers, and health care professionals should work together to monitor young children with ID. We can conclude three things: first, prevention should start early, when the first symptoms of internalising or externalising problems appear. Second, early training in communication skills for children with ID may prepare them better to face life's challenges and provide more options for helping them reduce stress when needed. Third, parents need to be supported; when they experience mental health problems, they will have less resources to raise and support their children.

Future research is needed to replicate these findings and add risk indicators not currently included (especially in mood disorders). Also, future studies should focus on designing and evaluating prevention programs for young people with ID based on the risk profiles found in this study and in future studies.

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Author Contributions

Conceptualization, H.M.K. and M.C.D.; Methodology, H.M.K., M.C.D., K.P.d.R. and J.C.H.D.; Validation, H.M.K. and M.C.D.; Formal Analysis, H.M.K., M.C.D., K.P.d.R. and J.C.H.D.; Investigation, M.C.D., K.P.d.R. and J.C.H.D.; Data Curation, M.C.D.; Writing—Original Draft Preparation, K.P.d.R.; Writing—Review & Editing, H.M.K., M.C.D., K.P.d.R. and J.C.H.D.; Supervision, H.M.K. and M.C.D.; Project Administration, H.M.K. and M.C.D.; Funding Acquisition, H.M.K.

Ethics Statement

The study was conducted according to the guidelines of the Declaration of Helsinki, and approved by the Committee for Medical Ethics of the Erasmus MC-Sophia Children's Hospital (Approval no. MEC 95.576).

Informed Consent Statement

Informed consent was obtained from all participants involved in the study.

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Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

References

1. Borthwick-Duffy SA. Epidemiology and prevalence of psychopathology in people with mental retardation. *J. Consult Clin. Psychol.* **1994**, *62*, 17–27.
2. Chadwick O, Kusel Y, Cuddy M, Taylor E. Psychiatric diagnoses and behaviour problems from childhood to early adolescence in young people with severe intellectual disabilities. *Psychol. Med.* **2005**, *35*, 751–760.
3. De Ruiter KP, Dekker MC, Douma JCH, Verhulst FC, Koot HM. Associated change in in risk and problem behaviours in youths with intellectual disabilities. Available online: <https://research.vu.nl/ws/portalfiles/portal/75917390/abstract+english.pdf> (accessed on 28 June 2024).
4. Dekker MC, Koot HM. DSM-IV disorders in children with borderline to moderate intellectual disability. I: Prevalence and impact. *J. Am. Acad. Child Adolesc. Psychiatry* **2003**, *42*, 915–922.
5. Dykens EM. Psychopathology in children with intellectual disability. *J. Child Psychol. Psychiatry* **2000**, *41*, 407–417.
6. Einfeld SL, Ellis LA, Emerson E. Comorbidity of intellectual disability and mental disorder in children and adolescents: A systematic review. *J. Intellect. Dev. Disabil.* **2011**, *36*, 137–143.
7. Emerson E. Prevalence of psychiatric disorders in children and adolescents with and without intellectual disability. *J. Intellect. Disabil. Res.* **2003**, *47*, 51–58.
8. Erickson MF, Sroufe LA, Egeland B. The relationship between quality of attachment and behavior problems in preschool in a high-risk sample. In *Monographs of the Society for Research in Child Development: Growing Points of Attachment Theory and Research*; University of Chicago Press: Chicago, IL, USA, 1985; Volume 50, pp. 147–166.
9. Koskentausta T, Iivanainen M, Almqvist F. Psychiatric disorders in children with intellectual disability. *Nord. J. Psychiatry.* **2002**, *56*, 126–131.
10. Linna SL, Moilanen I, Ebeling H, Piha J, Kumpulainen K, Tamminen T, et al. Psychiatric symptoms in children with intellectual disability. *Eur. Child Adolesc. Psychiatry* **1999**, *8*, 77–82.
11. Moss S, Emerson E, Kiernan C, Turner S, Hatton C, Alborz A. Psychiatric symptoms in adults with learning disability and challenging behaviour. *Br. J. Psychiatry* **2000**, *177*, 452–456.
12. Sevin JA, Bowers-Stephens C, Crafton CG. Psychiatric disorders in adolescents with developmental disabilities: Longitudinal data on diagnostic disagreement in 150 clients. *Child Psychiatry Hum. Dev.* **2003**, *34*, 147–163.

13. Sturmey P. DSM-III-R and persons with dual diagnoses: Conceptual issues and strategies for future research. *J. Intellect. Disabil. Res.* **1995**, *39*, 357–364.
14. Wallander JL, Dekker MC, Koot HM. Psychopathology in children and adolescents with intellectual disability: Measurement, prevalence, course, and risk. In *International Review of Research in Mental Retardation, Vol. 26*; Glidden LM, ed.; Academic Press: San Diego, CA, USA, 2023; pp. 93–134.
15. Bronfenbrenner U. *Making Human Beings Human: Bioecological Perspectives on Human Development*; Sage: Newbury Park, CA, USA, 2005.
16. Obradović J, Shaffer A, Masten AS. Risk and adversity in developmental psychopathology: Progress and future directions. In *The Cambridge Handbook of Environment in Human Development*; Mayes LG, Lewis M, eds.; Cambridge University Press: Cambridge, UK, 2012; pp. 35–57.
17. Rutter M, Kim-Cohen J, Maughan B. Continuities and discontinuities in psychopathology between childhood and adult life. *J. Child. Psychol. Psychiatry* **2006**, *47*, 276–295.
18. Barkley RA, Fischer M, Smallish L, Fletcher K. Young adult outcome of hyperactive children: Adaptive functioning in major life activities. *J. Am. Acad. Child Adolesc. Psychiatry* **2006**, *45*, 192–202.
19. Barlow JH, Ellard DR. The psychosocial well-being of children with chronic disease, their parents and siblings: An overview of the research evidence base. *Child Care Health Dev.* **2006**, *32*, 19–31.
20. Beardslee WR, Versage EM, Gladstone TR. Children of affectively ill parents: A review of the past 10 years. *J. Am. Acad. Child Adolesc. Psychiatry* **1998**, *37*, 1134–1141.
21. Caspi A, Moffitt TE, Newman DL, Silva PA. Behavioral observations at age 3 years predict adult psychiatric disorders. Longitudinal evidence from a birth cohort. *Arch. Gen. Psychiatry* **1996**, *53*, 1033–1039.
22. Offord DR, Boyle MH, Racine YA, Fleming JE, Cadman DT, Blum HM, et al. Outcome, prognosis, and risk in a longitudinal follow-up study. *J. Am. Acad. Child Adolesc. Psychiatry* **1992**, *31*, 916–923.
23. Velez CN, Johnson J, Cohen P. A longitudinal analysis of selected risk factors for childhood psychopathology. *J. Am. Acad. Child Adolesc. Psychiatry* **1989**, *28*, 861–864.
24. Buckley N, Glasson EJ, Chen W, Epstein E, Leonard H, Skoss R, et al. Prevalence estimates of mental health problems in children and adolescents with intellectual disability: A systematic review and meta-analysis. *Aust. N. Z. J. Psychiatry* **2020**, *54*, 970–984.
25. Platt JM, Keyes KM, McLaughlin KA, Kaufman AS. Intellectual disability and mental disorders in a US population representative sample of adolescents. *Psychol. Med.* **2019**, *49*, 952–961.
26. Weiss JA, Ting V, Perry A. Psychosocial correlates of psychiatric diagnoses and maladaptive behaviour in youth with severe developmental disability. *J. Intellect. Disabil. Res.* **2016**, *60*, 583–593.
27. Whitney DG, Shapiro DN, Peterson MD, Warschausky SA. Factors associated with depression and anxiety in children with intellectual disabilities. *J. Intellect. Disabil. Res.* **2019**, *63*, 408–417.
28. Munir KM. The co-occurrence of mental disorders in children and adolescents with intellectual disability/intellectual developmental disorder. *Curr. Opin. Psychiatry* **2016**, *29*, 95–102.
29. Dekker MC, Koot HM. DSM-IV disorders in children with borderline to moderate intellectual disability. II: Child and family predictors. *J. Am. Acad. Child Adolesc. Psychiatry* **2003**, *42*, 923–931.
30. Hamilton D, Sutherland G, Iacono T. Further examination of relationships between life events and psychiatric symptoms in adults with intellectual disability. *J. Intellect. Disabil. Res.* **2005**, *49*, 839–844.
31. Hastings RP, Daley D, Burns C, Beck A. Maternal distress and expressed emotion: Cross-sectional and longitudinal relationships with behavior problems of children with intellectual disabilities. *Am. J. Ment. Retard.* **2006**, *111*, 48–61.
32. Hatton C, Emerson E. The relationship between life events and psychopathology amongst children with intellectual disabilities. *J. Appl. Res. Intellect. Disabil.* **2004**, *17*, 109–117.
33. McCarthy J, Boyd J. Psychopathology and young people with Down’s syndrome: Childhood predictors and adult outcome of disorder. *J. Intellect. Disabil.* **2001**, *45*, 99–105.
34. Smit F, Beekman A, Cuijpers P, de Graaf R, Vollebergh W. Selecting key variables for depression prevention: Results from a population-based prospective epidemiological study. *J. Affect Disord.* **2004**, *81*, 241–249.
35. Smit F, Ederveen A, Cuijpers P, Deeg D, Beekman A. Opportunities for cost-effective prevention of late-life depression: An epidemiological approach. *Arch. Gen. Psychiatry* **2006**, *63*, 290–296.
36. Ashford J, Smit F, Van Lier PA, Cuijpers P, Koot HM. Early risk indicators of internalizing problems in late childhood: A 9-year longitudinal study. *J. Child Psychol. Psychiatry* **2008**, *49*, 774–780.
37. Campbell SB, Shaw DS, Gilliom M. Early externalizing behavior problems: Toddlers and preschoolers at risk for later maladjustment. *Dev. Psychopathol.* **2000**, *12*, 467–488.
38. Sameroff AJ. Environmental risk factors in infancy. *Pediatrics* **1998**, *102*, 1287–1292.
39. Arcelus J, Vostanis P. Psychiatric comorbidity in children and adolescents. *Curr. Opin. Psychiatry* **2005**, *18*, 429–434.
40. Dekker MC, Koot HM, Van der Ende J, Verhulst FC. Emotional and behavioral problems in children and adolescents with and without intellectual disability. *J. Child Psychol. Psychiatry* **2002**, *43*, 1087–1098.

41. Ferdinand RF, van der Ende J. *DISC-IV: Diagnostic Interview Schedule for Children; Informatie Ouder (Interview over Kind of Jeugdige)*; Department of Child and Adolescent Psychiatry, Sophia Children's Hospital: Rotterdam, The Netherlands, 1998.
42. Shaffer D, Fisher P, Lucas CP, Comer J. *Scoring manual Diagnostic Interview Schedule for Children (DISC-IV)*; Columbia University: New York, NY, USA, 2000.
43. Shaffer D, Fisher P, Lucas CP, Dulcan MK, Schwab-Stone ME. NIMH Diagnostic Interview Schedule for Children Version IV (NIMH DISC-IV): Description, differences from previous versions, and reliability of some common diagnoses. *J. Am. Acad. Child Adolesc. Psychiatry* **2000**, *39*, 28–38.
44. Wahler HJ. The Physical Symptoms Inventory: Measuring levels of somatic complaining behavior. *J. Clin. Psychol.* **1968**, *24*, 207–211.
45. Sparrow SS, Balla DA, Carter AS, Cicchetti DV. Vineland Screener. Overview, reliability, validity, administration and scoring. Manuscript obtained from the authors.
46. Sparrow SS, Balla DA, Cicchetti DV. *Manual Vineland Adaptive Behavior Scales. Interview Edition*; American Guidance Service: Circle Pines, MN, USA, 1984.
47. Achenbach TM. *Manual for the Child Behavior Checklist/4-18 and 1991 Profiles*; Department of Psychiatry, University of Vermont: Burlington, VT, USA, 1991.
48. Verhulst FC, van der Ende J, Koot HM. *Handleiding voor de CBCL/4-18*; Afdeling Kinder- en jeugdpsychiatrie: Rotterdam, The Netherlands, 1996.
49. Central Bureau of Statistics. Basisonderwijs en (voortgezet) speciaal onderwijs: Leerlingen naar leeftijd 1995/1996 (primary and secondary special education: Students by age 1995/1996). Available online: <https://www.cbs.nl/nl-nl/cijfers/detail/70762ned> (accessed on 28 June 2024).
50. Byles JA, Byrne C, Boyle MH, Offord DR. Ontario Child Health Study: Reliability and validity of the general functioning subscale of the McMaster Family Assessment Device. *Fam. Process.* **1988**, *27*, 97–104.
51. Achenbach TM. *Manual for the Young Adult Self-Report and Young Adult Behavior Checklist*; Department of Psychiatry, University of Vermont: Burlington, VT, USA, 1997.
52. Wiznitzer M, Verhulst FC, van den Brink W, Koeter M, van der Ende J, Giel R, et al. Detecting psychopathology in young adults: The Young Adult Self Report, the General Health Questionnaire and the Symptom Checklist as screening instruments. *Acta. Psychiatr. Scand.* **1992**, *86*, 32–37.
53. Verhulst FC, Van der Ende J. Factors associated with child mental health service use in the community. *J. Am. Acad. Child. Adolesc. Psychiatry* **1997**, *36*, 901–909.
54. Berden GF, Althaus M, Verhulst F. Major life events and changes in the behavioural functioning of children. *J. Child Psychol. Psychiatry* **1990**, *31*, 949–959.
55. StataCorp (2004). Stata Statistical Software: Release 8.2. Available online: <https://www.stata.com/manuals/r.pdf> (accessed on 28 June 2024).
56. Greenland S, Drescher K. Maximum likelihood estimation of the attributable fraction from logistic models. *Biometrics* **1993**, *49*, 865–872.
57. Burt SA, McGue M, Krueger RF, Iacono WG. How are parent-child conflict and childhood externalizing symptoms related over time? Results from a genetically informative cross-lagged study. *Dev. Psychopathol.* **2005**, *17*, 145–165.
58. Rutter M. Protective factors in children's responses to stress and disadvantage. In *Primary Prevention of Psychopathology, Social Competence in Children*; Kent MW, Rolf JE, eds.; University Press of New England: Hanover, NH, USA, 1997.
59. Koot HM. Problem Behavior in Dutch Preschoolers. Doctoral Thesis, Erasmus University Rotterdam, Rotterdam, The Netherlands, 6 October 1993.
60. Vervoort-Schel J, Mercera G, Wissink I, Van der Helm P, Lindauer R, Moonen X. Prevalence of and relationship between adverse childhood experiences and family context risk factors among children with intellectual disabilities and borderline intellectual functioning. *Res. Dev. Disabil.* **2021**, *113*, 103935.