

Time-to-treatment of mental disorders in a community sample of Dutch adolescents. A TRAILS study

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Aims. Timely recognition and treatment of mental disorders with an onset in childhood and adolescence is paramount, as these are characterized by greater severity and longer persistence than disorders with an onset in adulthood. Studies examining time-to-treatment, also referred to as treatment delay, duration of untreated illness or latency to treatment, and defined as the time between disorder onset and initial treatment contact, are sparse and all based on adult samples. The aim of this study was to describe time-to-treatment and its correlates for any health care professional (any care) and secondary mental health care (secondary care), for a broad range of mental disorders, in adolescents.

Methods. Data from the Dutch community-based cohort study TRacking Adolescents' Individual Lives Survey (TRAILS; $N = 2230$) were used. The Composite International Diagnostic Interview (CIDI) was administered to assess DSM-IV disorders, the age of onset, and the age of initial treatment contact with any health care professional in 1584 adolescents of 18–20 years old. In total 43% of the adolescents ($n = 675$) were diagnosed with a lifetime DSM-IV disorder. The age of initial treatment contact with secondary care was based on administrative records from 321 adolescents without a disorder onset before the age of 10. Descriptive statistics, cumulative lifetime probability plots, and Cox regression analyses were used to analyze time-to-treatment.

Results. The proportion of adolescents who reported lifetime treatment contact with any care varied from 15% for alcohol dependence to 82% for dysthymia. Regarding secondary care, proportions of lifetime treatment contact were lower for mood disorders and higher for substance dependence. Time-to-treatment for any care varied considerably between and within diagnostic classes. The probability of lifetime treatment contact for mood disorders was above 90%, whereas for other mental disorders this was substantially lower. An earlier age of onset predicted a longer, and the presence of a co-morbid mood disorder predicted a shorter time-to-treatment in general. Disorder severity predicted a shorter time-to-treatment for any care, but not for secondary care. Time-to-treatment for secondary care was shorter for adolescents from low and middle socioeconomic background than for adolescents from a high socioeconomic background.

Conclusion. Although the time-to-treatment was shorter for adolescents than for adults, it was still substantial, and the overall patterns were remarkably similar to those found in adults. Efforts to reduce time-to-treatment should therefore be aimed at children and adolescents. Future research should address mechanisms underlying time-to-treatment and its consequences for early-onset disorders in particular.

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Introduction

The prevalence of mental disorders is high (Kessler *et al.* 2005a; Moffitt *et al.* 2010; de Graaf *et al.* 2012). Although mental disorders are associated with a

tremendous disease burden (Whiteford *et al.* 2013), worldwide, no more than one-third of people with a mental disorder receive treatment (Kessler *et al.* 2005b; Thornicroft, 2012). The majority of mental disorders in adulthood have their onset in adolescence and early adulthood (Wang *et al.* 2005; Kessler *et al.* 2007; Merikangas *et al.* 2010; de Girolamo *et al.* 2012; Ormel *et al.* 2015), and interfere with key areas of development such as education, social relationships and the transition to work (Costello & Maughan, 2015). Timely recognition and treatment of such

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early-onset mental disorders is paramount, as these are characterised by greater severity and longer persistence than disorders with an onset in adulthood (Kessler *et al.* 1998; Korczak & Goldstein, 2009; Reef *et al.* 2010). However, despite the apparent need for care (Jörg *et al.* 2015), only a small proportion of youths actually receive timely treatment (Merikangas *et al.* 2011; de Girolamo *et al.* 2012; Jansen *et al.* 2013).

Studies focusing on the time between the onset of a mental disorder and initial treatment contact (time-to-treatment; also referred to as treatment delay, duration of untreated illness or latency to treatment), have mainly focused on the duration of untreated psychosis (DUP). Review studies show that a longer DUP is associated with a worse course of illness and worse outcomes (Marshall *et al.* 2005; Perkins *et al.* 2005). Studies focusing on time-to-treatment in common mental disorders are sparse (Ghio *et al.* 2014), but these also point towards poorer outcomes of disorders with longer time-to-treatment (Kisely *et al.* 2006; Dell'Osso & Altamura, 2010). Of particular interest is the finding that an earlier age of onset is associated with a longer time-to-treatment in both community (Kessler *et al.* 1998; Christiana *et al.* 2000; Wang *et al.* 2004, 2005, 2007; Bruffaerts *et al.* 2007; Korczak & Goldstein, 2009; ten Have *et al.* 2013a) and clinical samples (Altamura *et al.* 2007, 2008). Although these studies generally stress the importance of recognition and treatment of early-onset disorders in the critical age range of 10–24 years, they are all based on adult samples.

Our aim is to expand on the available literature by describing time-to-treatment and its correlates for any health care professional (hereafter referred to as *any care*) and secondary mental health care (hereafter referred to as *secondary care*), for a broad range of mental disorders, in adolescents. We will use data from the Dutch Tracking Adolescents' Individual Lives Survey (TRAILS), a large community-based cohort study in which participants were followed from childhood into emerging adulthood (Oldehinkel *et al.* 2015), to do so. The Composite International Diagnostic Interview (CIDI; Kessler & Üstün, 2004) was administered to establish age of onset of mental disorders as well as age of initial treatment contact with any care. Furthermore, data from the Psychiatric Case Register North Netherlands (PCRNN) were used to establish age of initial treatment contact with secondary care.

Methods

Sample

The data used in this study were from TRAILS, a prospective population-based cohort study aimed at explaining the development of mental health from

early adolescence into adulthood. The TRAILS sample, response rates and study contents have been described in detail elsewhere (de Winter *et al.* 2005; Nederhof *et al.* 2012; Oldehinkel *et al.* 2015). In short, after the exclusion of children whose schools refused participation ($n=338$), and children with serious mental or physical health problems or language difficulties ($n=210$), consent to participate in the study was obtained from 2230 (76.0%) out of 2935 eligible children and their parents. Teacher-reported levels of psychopathology did not differ between responding and non-responding children, but boys, children with a lower socioeconomic background, and children with relatively poorer school performance were more likely to be non-responders (de Winter *et al.* 2005).

We used data from the first and fourth assessment wave, which ran from March 2001 to July 2002 (T1), and from October 2008 to September 2010 (T4). Of the T1 participants (mean age=11.1 years, *s.d.*=0.6 years, 50.8% girls), 84.3% still participated at T4 ($n=1881$, mean age=19.1 years, *s.d.*=0.6 years, 52.3% girls). Drop-out was related to being male, low intelligence, low educational level, low parental socioeconomic position (SEP), single-parent families, being bullied and parent-reported behaviour problems (Nederhof *et al.* 2012). As part of T4, the CIDI was completed by 1584 adolescents (response rate=71.0%, mean age=19.3 years, *s.d.*=0.6 years, 54.0% girls) (Ormel *et al.* 2015).

The TRAILS data were linked to the PCRNN (hereafter referred to as the register), which covers secondary child, adolescent and adult mental health care organisations. The catchment area of the register covers 1.7 million inhabitants, and overlaps with the geographic area from which TRAILS participants were recruited. The register, which contained data from 2000 onward, did not include primary (youth) mental health care, private practices and commercial mental health care organisations. Consent to link the TRAILS database to the register was obtained from 1385 CIDI participants and their parents (87.4%). A 95% likelihood matching procedure uniquely identified 342 children with one or more records in the PCRNN (24.7%).

Measures

Lifetime prevalence, age of onset and age of initial treatment contact of DSM-IV disorders (American Psychological Association, 1994) were established using the World Mental Health Organization (WHO) CIDI version 3.0 (Kessler & Üstün, 2004), a structured diagnostic interview that can be administered by trained lay interviewers. Clinical reappraisal studies showed generally good validity of CIDI diagnoses

when compared with blinded clinical reappraisal interviews (Haro *et al.* 2006; Kessler *et al.* 2009). Disorders included in this study were mood disorders [major depressive disorder (MDD), dysthymic disorder (DYS) and bipolar disorder types I and II (BPD)]; anxiety disorders (separation anxiety disorder (SAD), agoraphobia without panic disorder (AGP), generalised anxiety disorder (GAD), obsessive-compulsive disorder (OCD), panic disorder (PDS), social phobia (SO) and specific phobia (SP)); behaviour disorders [attention deficit hyperactivity disorder (ADHD) and oppositional defiant disorder (ODD)]; and substance dependence [alcohol dependence (ALD) and drug dependence (DRD)]. Organic exclusion criteria, for disorders caused by physical illness, and diagnostic hierarchy rules, for disorders better explained by other disorders, were used where applicable.

Time-to-treatment was defined as the time in years between the age of onset, which is the age at which all DSM-IV diagnostic criteria for the index disorder were met for the first time, and the age of initial treatment contact. The age of initial treatment contact was established in two different ways: with regard to any care as assessed by the CIDI, and with regard to secondary care based on the register. In the CIDI, respondents were asked in each diagnostic section separately whether they had ever talked about the symptoms of the index disorder with a medical doctor or any other health care professional, such as psychologists, clergymen, herbalists and acupuncturists. If acknowledged, respondents were asked their age at first contact. For respondents with a record in the register, the age of initial treatment contact in secondary care was determined based on the date of the first entry in the register. Thus, in case of multiple disorders, the age of initial treatment contact could differ by disorder for any care, while it would be the same for all disorders for secondary care.

The predictor variables included sex (male; female), ethnic minority status (at least one parent born in a non-developed country; both parents born in a developed country), intelligence ($IQ < 85$; $85 \leq IQ \leq 115$; $IQ > 115$) (Silverstein, 1975), parental SEP (lowest 25%; intermediate 50%; highest 25%) (Amony-P'Olak *et al.* 2010), number of biological parents in the household (none or one; two), disorder severity (mild; severe) (Merikangas *et al.* 2010; Ormel *et al.* 2015), age at onset (1–5; 6–10; 11–15; 16–20) and presence of a co-morbid disorder from another diagnostic class (no; yes). Intelligence, parental SEP, and number of biological parents in the household were assessed at T1. A disorder was considered severe if it exceeded, at any time, the impairment or distress thresholds required for the regular CIDI DSM-IV disorders.

Co-morbidity was included as a time-varying covariate for each diagnostic class separately.

Analyses

For the analyses, only participants with a CIDI DSM-IV diagnosis were included, which amounted to 42.6% of all CIDI participants ($n=675$). Although seemingly high, according to Ormel *et al.* (2015) the prevalence rates found in TRAILS are comparable with those found in similar studies. Results from prospective studies suggest that actual lifetime prevalence rates are even higher (Moffitt *et al.* 2010), and that emotional and behavioural problems are nearly universal in nature (Angst *et al.* 2016). Of the CIDI participants with consent to link their data to the register ($n=1385$), 23.2% ($n=321$) were included because their first disorder had an onset since 2000, whereas 19.5% ($n=270$) were excluded because of a disorder onset before 2000. Cases with a disorder onset before 2000 were more often identified in the register (38 *v.* 26%, $\chi^2_1=9.7$, $p<0.002$), and more often had disorders from multiple diagnostic classes (47 *v.* 28%, $\chi^2_1=25.4$, $p<0.001$) than cases with a disorder onset since 2000.

The main analyses were divided into two parts. First, time-to-treatment was described using observed proportions of adolescents who made treatment contact at any point in their lives, subdivided into three groups: after initial symptoms and before, in, and after the year of onset of the full-blown disorder. Furthermore, cumulative probability curves of lifetime treatment contact were estimated using survival analysis. These curves were generated for each disorder separately using survival analysis, and showed the estimated cumulative proportion of cases that eventually make treatment contact. The actuarial method was used, because it is better suited than the Kaplan–Meier method for events for which the period rather than the exact date during which an event has occurred is known (c.f. Wang *et al.* 2005; Bunting *et al.* 2012). The results were weighted by sex, the Child Behavior Checklist total problems score (Achenbach & Rescorla, 2001), and parental SEP to account for selective non-response (Ormel *et al.* 2015). Second, Cox regression analyses (Kleinbaum & Klein, 2012) were used to test predictors of time-to-treatment for the four diagnostic classes, and any disorder. Cases without treatment contact were censored at the age of the interview. The analyses were performed separately for any care and for secondary care.

Additionally, we performed two sensitivity analyses on the data regarding any care. First, the Cox regression analyses were repeated while excluding cases with any disorder onset before 2000. This exclusion criterion was also used in the analyses regarding

secondary care. Second, the Cox regression analyses for disorder classes were repeated using treatment for any disorder rather than disorder-specific treatment, because the register data could not be linked to any specific disorder class.

All analyses were performed with SPSS 23.0 (IBM Corp, 2015).

Results

Table 1 shows the proportions of lifetime treatment contact, subdivided into treatment contact after initial symptoms and before, in, and after the year of onset of the full-blown disorder, for any care and secondary care, among adolescents with a mental disorder according to the CIDI. The proportion of adolescents with a mental disorder who had lifetime treatment contact with any care by the age of 18–20 varied between 15% for alcohol dependence and 82% for dysthymia. Lifetime treatment rates for secondary care were considerably lower for mood disorders, and higher for substance dependence compared with any care. Notably, for secondary care the proportions of cases with treatment contact before onset of the full-blown disorder were higher than for any care.

Figure 1 shows for each disorder separately the estimated cumulative proportions of adolescents who will eventually make treatment contact. Three observations stood out. First, the curves showed much variation. For mood (Fig. 1a) and behaviour disorders (Fig. 1c), the curves were comparable within their class, but differed markedly from the other class. The curves for anxiety disorders (Fig. 1b) showed much within-class variation. Curves for substance dependence (Fig. 1d) resembled those for behaviour disorders. Second, disorders with a high probability of treatment contact, such as major depression and generalised anxiety, typically had distinctly higher proportions of initial treatment contact in the first years after onset than disorders with a low probability of treatment contact, such as separation anxiety and specific phobia. Third, time-to-treatment was substantial. Time-to-treatment was shortest for mood disorders, yet the cumulative probability of treatment contact at 2 years after onset was only 50%. A cumulative probability of treatment contact of 50% for anxiety and behaviour disorders was only reached 17 and 12 years after onset, respectively.

Results from the Cox regression analyses predicting time-to-treatment are shown in Table 2 (any care) and 3 (secondary care). Age of onset predicted time-to-treatment for any disorder for both any care and secondary care. When a disorder had an earlier onset, the time-to-treatment was longer. Models analysing each of the disorder classes separately showed similar effects,

although the effects were mostly non-significant for secondary care. Co-morbidity predicted time-to-treatment only in six out of the 32 possible associations reported in Tables 2 and 3; a co-morbid mood disorder in particular predicted a shorter time-to-treatment. Disorder severity predicted shorter time-to-treatment with any care, while it was not associated with secondary care. The effect of parental SEP showed a trend towards shorter time-to-treatment for secondary care for adolescents with parents from a low or middle SEP compared to adolescents with parents from a high SEP.

Two sensitivity analyses for any care were performed (available as online supplementary material). When excluding adolescents with any disorder onset before 2000 (appendix table 1), age at onset effects were often no longer statistically significant, although hazard ratios remained similar, and disorder severity no longer significantly predicted a shorter time-to-treatment for anxiety and behaviour disorders. When considering any treatment contact rather than disorder-specific treatment contact (appendix table 2), co-morbidity more often predicted a shorter time-to-treatment.

Conclusion

The time-to-treatment with any care for adolescents varied considerably across disorders, but was substantial even for mood disorders, which in general showed the shortest time-to-treatment. Cox regression analyses showed that the time-to-treatment was longer as the onset was earlier. Furthermore, the time-to-treatment was shorter for severe compared with mild disorders, and for disorders with a co-morbid mood disorder. These results were replicated for secondary care, with the exception that disorder severity was not related to time-to-treatment.

Limitations

The results need to be interpreted considering three limitations. The first limitation is recall bias (e.g., Wang *et al.* 2005; Altamura *et al.* 2010). Respondents may forget about or downplay mental health problems for which they did not seek treatment, which would lead to overestimated proportions of treatment contact. Recall bias may also cause respondents to remember past events as more recent than they actually took place (telescoping). Since onset usually occurs years before initial treatment contact, the probability of telescoping is likely larger for age of onset than for age of initial treatment contact. The time-to-treatment is therefore possibly underestimated. Our study, however, had two advantages over previous studies (Wang *et al.* 2004; Bruffaerts *et al.* 2007; Bunting *et al.* 2012; ten Have *et al.* 2013a), which probably limited recall bias. First, the

Table 1. Weighted^a lifetime treatment contact, subdivided into initial treatment contact after initial symptoms and before, in, or after the year of onset of the full-blown disorder. The left part of the table shows treatment contact with any health care professional^b for all adolescents with a DSM-IV disorder. The right part of the table shows treatment contact with secondary mental health care^c for adolescents with a DSM-IV disorder but without any disorder onset before 2000

	Initial treatment contact with any health care professional for all adolescents with a full-blown DSM-IV disorder ^b					Initial treatment contact with secondary mental health care for adolescents with a full-blown DSM-IV disorder but without any disorder onset before 2000 ^c				
	Lifetime		Before year of onset % (S.E.)	In year of onset % (S.E.)	After year of onset % (S.E.)	Lifetime		Before year of onset % (S.E.)	In year of onset % (S.E.)	After year of onset % (S.E.)
	<i>n</i>	% (S.E.)				<i>n</i>	% (S.E.)			
Mood disorders										
Bipolar disorder	13	55.4 (10.5)	4.3 (4.3)	21.9 (8.7)	29.1 (9.6)	3	39.5 (19.2)	27.4 (17.5)	0.0 (0.0)	12.1 (12.9)
Major depressive disorder	159	64.5 (3.1)	7.1 (1.6)	23.4 (2.7)	33.9 (3.0)	48	34.5 (4.0)	10.7 (2.6)	4.6 (1.8)	19.3 (3.3)
Dysthymia	22	81.7 (7.5)	14.6 (6.9)	26.3 (8.6)	40.9 (9.6)	1	14.1 (14.4)	0.0 (0.0)	0.0 (0.0)	14.1 (14.4)
Any mood disorder	175	63.8 (2.9)	7.2 (1.6)	23.4 (2.6)	33.3 (2.9)	51	34.5 (3.9)	11.5 (2.6)	4.3 (1.7)	18.8 (3.2)
Anxiety disorders										
Separation anxiety disorder	10	22.5 (6.2)	0.0 (0.0)	10.5 (4.5)	12.0 (4.8)	5	39.4 (15.0)	21.9 (12.7)	8.8 (8.7)	8.8 (8.7)
Agoraphobia without panic disorder	6	37.8 (12.8)	0.0 (0.0)	18.5 (10.3)	19.3 (10.4)	2	65.9 (34.2)	0.0 (0.0)	34.9 (34.4)	31.0 (33.4)
Generalised anxiety disorder	27	58.1 (7.4)	6.5 (3.7)	21.3 (6.1)	30.3 (6.9)	9	43.9 (11.5)	19.8 (9.2)	14.3 (8.1)	9.8 (6.9)
Obsessive-compulsive disorder	28	30.2 (4.8)	0.0 (0.0)	7.6 (2.7)	22.6 (4.3)	11	28.9 (7.5)	13.7 (5.7)	2.4 (2.5)	12.8 (5.5)
Panic disorder	11	43.0 (9.9)	0.0 (0.0)	35.5 (9.6)	7.6 (5.3)	4	45.7 (17.8)	22.9 (15.0)	0.0 (0.0)	22.8 (15.0)
Social phobia	68	34.6 (3.4)	0.0 (0.0)	4.7 (1.5)	29.9 (3.3)	21	22.3 (4.3)	5.0 (2.3)	0.9 (1.0)	16.3 (3.9)
Specific phobia	44	24.1 (3.2)	0.0 (0.0)	2.8 (1.2)	21.4 (3.1)	10	31.2 (8.4)	2.6 (2.9)	3.1 (3.2)	25.5 (7.9)
Any anxiety disorder	152	34.6 (2.3)	0.0 (0.0)	5.7 (1.1)	28.9 (2.2)	46	27.3 (3.4)	8.9 (2.2)	1.6 (1.0)	16.8 (2.9)
Behaviour disorders										
Attention deficit disorder	37	56.5 (6.2)	3.4 (2.3)	6.3 (3.1)	46.8 (6.2)	2	100.0 (0.0)	62.8 (40.4)	0.0 (0.0)	37.2 (40.4)
Oppositional defiant disorder	58	41.7 (4.2)	5.0 (1.9)	12.1 (2.8)	24.6 (3.7)	17	38.3 (7.4)	7.0 (3.9)	5.5 (3.5)	25.8 (6.7)
Any behaviour disorder	82	45.9 (3.7)	4.6 (1.6)	9.9 (2.2)	31.4 (3.5)	19	41.5 (7.3)	9.9 (4.4)	5.3 (3.3)	26.4 (6.5)
Substance dependence										
Alcohol dependence	8	15.3 (5.1)	3.7 (2.7)	6.1 (3.4)	5.5 (3.3)	4	18.4 (8.1)	14.6 (7.4)	3.8 (4.0)	0.0 (0.0)
Drug dependence	18	25.5 (5.2)	6.6 (2.9)	4.0 (2.3)	14.9 (4.2)	13	37.5 (8.2)	21.0 (6.9)	2.8 (2.8)	13.6 (5.8)
Any substance dependence	23	20.3 (3.8)	4.9 (2.1)	3.3 (1.7)	12.0 (3.1)	17	30.0 (6.2)	17.8 (5.2)	3.4 (2.5)	8.7 (3.8)
Total										
Any disorder	328	48.3 (1.9)	2.1 (0.5)	9.1 (1.1)	37.2 (1.9)	87	27.6 (2.5)	9.2 (1.6)	1.9 (0.8)	16.5 (2.1)

^aWeighted by sex, Child Behavior Checklist cut-offs (normal *v.* borderline clinical/clinical) and parental SEP. Cases with missing values were assigned the weight 1.

^bInitial treatment contact for any health care professional based on the Composite International Diagnostic Interview.

^cInitial treatment contact for secondary mental health care based on the Psychiatric Case Register North Netherlands. Adolescents with any disorder onset before 2000 were excluded because register data were not available before 2000.

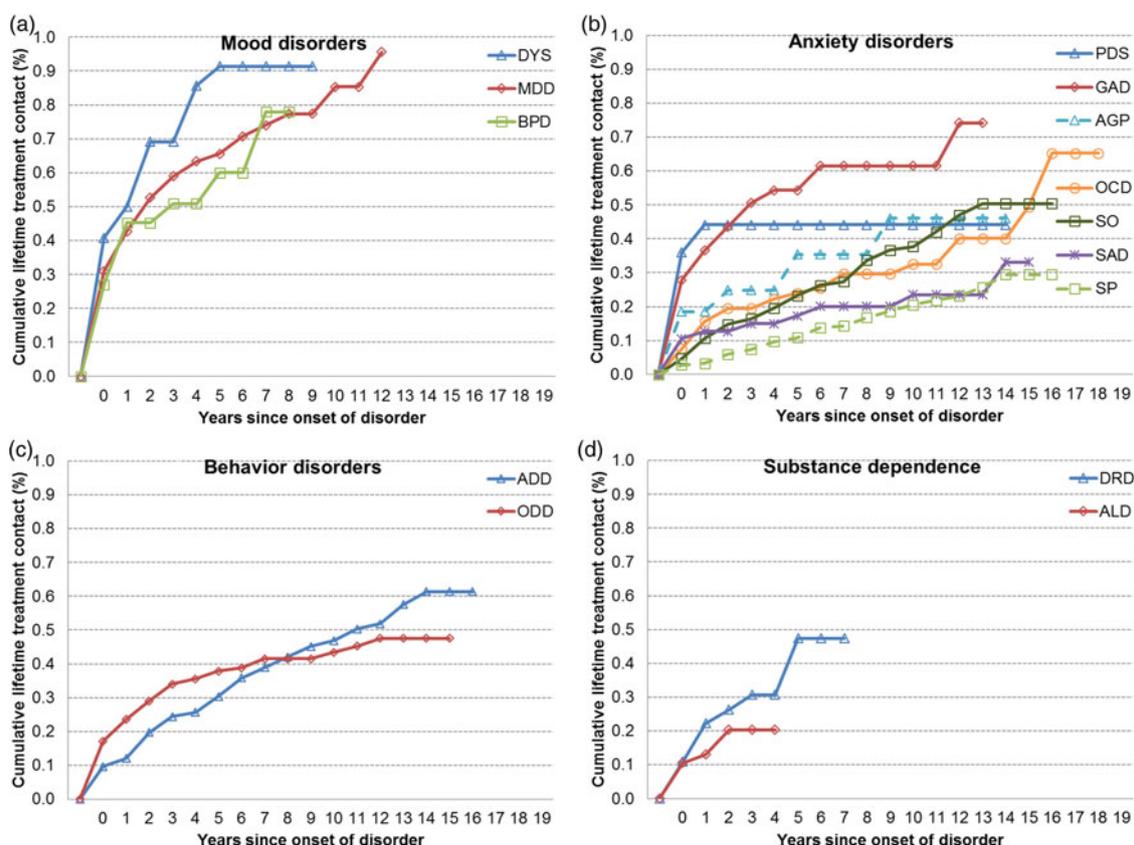


Fig. 1. Weighted cumulative lifetime treatment probabilities with any health care professional for DSM-IV mood disorders (a), anxiety disorders (b), behavior disorders (c), and substance dependence (d). Notes: Weighted by sex, Child Behavior Checklist cut-offs (normal v. borderline clinical/clinical) and parental SEP. Cases with missing values were assigned the weight 1. Probabilities based on life tables using the Actuarial method. Time-to-treatment for disorders with initial treatment contact after initial symptoms and before the year of onset of the respective full-blown disorder set to 0. DYS: dysthymia; MDD: major depressive disorder; BPD: bipolar disorder types I and II; PDS: panic disorder; GAD: generalized anxiety disorder; AGP: agoraphobia; OCD: obsessive-compulsive disorder; SO: social phobia; SAD: separation anxiety disorder; SP: specific phobia; ADD: attention deficit hyperactivity disorder; ODD: oppositional defiant disorder; DRD: drug dependence; ALD: alcohol dependence.

diagnostic interview was administered at the age of 18–20 years rather than up to 60 years and older, so the recall period was much shorter than in previous studies. Second, administrative records are considered more reliable than self-reported treatment seeking (Wang *et al.* 2004; Olfson *et al.* 2012).

The second limitation is that most predictors of time-to-treatment were not assessed over time. Intelligence, parental SEP, and the number of biological parents in the household were only assessed when the participants were 10–12 years old. For the majority of adolescents, however, these are likely to be stable factors. Furthermore, disorder severity could only be assessed lifetime, rather than at the moment of initial treatment contact. Therefore, assuming symptom recognition and treatment seeking are more likely when disorders are severe than when they are mild (Merikangas *et al.* 2011; ten Have *et al.* 2013b) the effect

of disorder severity on time-to-treatment could have been underestimated.

The third limitation of this study concerns the coverage of the PCRNN. First, this register does not cover primary (youth) mental health care, private practices, and commercial mental health care organisations. Nevertheless, the register still covers an estimated 75% of all mental health treatment trajectories for children and adolescents in secondary care (Jörg *et al.* 2015). Second, this register does not include data prior to 2000, which corresponds approximately to the age of ten in our sample. Both lead to an underestimation of the proportion of secondary care users. Additionally, time-to-treatment for secondary care is likely to be underestimated considerably, because cases with early-onset disorders, who typically had the longest time-to-treatment, had to be excluded from the secondary care analyses.

Table 2. Cox regression analyses predicting time-to-treatment^a with any health care professional^b for DSM-IV disorders by disorder class and any disorder

	Any health care professional ^b				
	Mood disorders HR (95 %CI)	Anxiety disorders HR (95 %CI)	Behaviour disorders HR (95 %CI)	Substance dependence HR (95 %CI)	Any disorder HR (95 %CI)
Male (ref = female)	0.74 (0.51–1.06)	0.90 (0.61–1.31)	1.64 (0.97–2.77)	0.96 (0.39–2.36)	0.88 (0.69–1.13)
Ethnic minority (ref = majority)	1.01 (0.61–1.69)	0.50 (0.24–1.01)	0.82 (0.29–2.27)	– ^c	0.90 (0.59–1.36)
Low IQ (ref = high)	1.46 (0.74–2.85)	1.72 (0.88–3.39)	1.52 (0.51–4.53)	3.27 (0.58–18.43)	1.37 (0.85–2.21)
Middle IQ (ref = high)	1.29 (0.74–2.23)	1.36 (0.77–2.40)	1.73 (0.67–4.48)	1.74 (0.37–8.20)	1.31 (0.88–1.93)
Low parental SEP (ref = high)	1.20 (0.74–1.94)	1.08 (0.64–1.80)	1.02 (0.53–1.98)	1.31 (0.42–4.10)	1.27 (0.91–1.77)
Middle parental SEP (ref = high)	1.39 (0.94–2.06)	1.04 (0.69–1.58)	0.64 (0.34–1.18)	1.30 (0.46–3.70)	1.18 (0.90–1.57)
0 or 1 biological parents (ref = both)	0.97 (0.68–1.37)	1.32 (0.93–1.88)	1.03 (0.61–1.75)	0.94 (0.36–2.45)	1.28* (1.00–1.62)
Severe disorder (ref = mild) ^d	1.73*** (1.26–2.36)	1.94*** (1.33–2.85)	1.98** (1.19–3.29)	– ^d	1.57*** (1.22–2.03)
Age at onset 1–5 (ref = 16–20)	– ^e	0.15*** (0.07–0.32)	0.67 (0.35–1.30)	– ^c	0.20*** (0.12–0.32)
Age at onset 6–10 (ref = 16–20)	0.38** (0.20–0.73)	0.23*** (0.11–0.47)	0.41** (0.21–0.80)	– ^f	0.26*** (0.17–0.42)
Age at onset 11–15 (ref = 16–20)	0.79 (0.55–1.13)	0.48* (0.24–0.93)	– ^g	1.78 (0.73–4.32)	0.58** (0.39–0.87)
Co-morbid mood disorder (ref = no) ^h	– ⁱ	2.79*** (1.90–4.09)	1.82 (0.93–3.56)	2.43 (0.98–6.03)	3.09*** (2.35–4.06)
Co-morbid anxiety disorder (ref = no) ^h	1.01 (0.73–1.39)	– ⁱ	0.88 (0.53–1.49)	2.43 (0.98–6.03)	1.09 (0.83–1.42)
Co-morbid behavior disorder (ref = no) ^h	0.85 (0.57–1.29)	1.49 (0.98–2.25)	– ⁱ	0.60 (0.22–1.60)	1.49** (1.11–2.01)
Co-morbid substance dependence (ref = no) ^h	1.91* (1.12–3.27)	0.81 (0.38–1.73)	1.04 (0.35–3.09)	– ⁱ	1.06 (0.65–1.73)
Chi-square	36.3 (13)***	125.9 (14)***	24.9 (13)*	14.9 (10)	276.2 (15)***
<i>n</i> events	172	145	72	23	314
<i>n</i> censored	96	284	93	86	347
<i>n</i> total	268	429	165	109	661

DSM-IV, Diagnostic and Statistical Manual of Mental Disorders, fourth edition; HR, Hazard ratio; 95% CI, 95% confidence interval; IQ, intelligence quotient; SEP, socio-economic position.

^aTime-to-treatment for disorders with initial treatment contact after initial symptoms and before the year of onset of the respective full-blown disorder set to 0.

^bLifetime treatment contact and age of initial treatment contact for any health care professional based on the Composite International Diagnostic Interview.

^cCovariate excluded due to insufficient cases.

^dAll substance dependence diagnoses were considered severe.

^eCombined with age of onset 6–10 years due to insufficient cases.

^fCombined with age of onset 11–15 years due to insufficient cases.

^gCombined with age of onset 16–20 years (reference category) due to insufficient cases in the reference category.

^hTime-dependent covariate.

ⁱDisorder class is the dependent variable.

* $p < 0.05$ ** $p < 0.01$ *** $p < 0.001$.

Table 3. Cox regression analyses predicting time-to-treatment^a with secondary mental health care^b for DSM-IV disorders by disorder class and any disorder for adolescents without any disorder onset before 2000

	Secondary mental health care ^b				
	Mood disorders HR (95% CI)	Anxiety disorders HR (95% CI)	Behaviour disorders HR (95% CI)	Substance dependence HR (95% CI)	Any disorder HR (95% CI)
Male (ref = female)	1.31 (0.66–2.63)	0.73 (0.33–1.62)	1.54 (0.35–6.73)	1.81 (0.49–6.76)	1.09 (0.67–1.77)
Ethnic minority (ref = majority)	0.53 (0.15–1.91)	– ^c	23.12* (1.32–403.92)	– ^c	0.89 (0.35–2.30)
Low IQ (ref = high)	1.84 (0.59–5.74)	1.12 (0.38–3.29)	1.19 (0.17–8.17)	0.90 (0.12–6.78)	1.81 (0.77–4.25)
Middle IQ (ref = high)	1.15 (0.43–3.05)	0.96 (0.38–2.40)	0.51 (0.09–2.75)	0.47 (0.10–2.13)	1.12 (0.55–2.29)
Low parental SEP (ref = high)	1.71 (0.65–4.48)	3.08 (0.98–9.70)	6.82 (0.69–67.49)	3.32 (0.63–17.45)	1.66 (0.81–3.41)
Middle parental SEP (ref = high)	2.09 (0.96–4.53)	3.49* (1.25–9.74)	5.49 (0.59–51.45)	1.47 (0.39–5.53)	1.85* (1.03–3.32)
0 or 1 biological parents (ref = both)	1.50 (0.78–2.90)	1.54 (0.78–3.05)	1.55 (0.40–5.99)	0.61 (0.18–2.14)	1.42 (0.88–2.28)
Severe disorder (ref = mild) ^d	1.39 (0.78–2.50)	1.44 (0.69–3.03)	0.70 (0.21–2.38)	– ^d	1.20 (0.71–2.02)
Age at onset 1–5 (ref = 16–20)	– ^e	– ^e	– ^e	– ^e	– ^e
Age at onset 6–10 (ref = 16–20)	0.17 (0.02–1.33)	0.41 (0.12–1.45)	0.54 (0.11–2.59)	– ^f	0.34** (0.15–0.77)
Age at onset 11–15 (ref = 16–20)	0.71 (0.37–1.37)	0.54 (0.20–1.47)	– ^g	0.54 (0.13–2.30)	0.58 (0.31–1.09)
Co-morbid mood disorder (ref = no) ^h	– ⁱ	4.30*** (2.23–8.28)	1.65 (0.38–7.04)	2.35 (0.72–7.60)	2.13** (1.24–3.67)
Co-morbid anxiety disorder (ref = no) ^h	1.82 (0.99–3.33)	– ⁱ	1.09 (0.29–4.06)	1.60 (0.45–5.67)	1.18 (0.69–2.04)
Co-morbid behavior disorder (ref = no) ^h	1.54 (0.65–3.67)	1.54 (0.63–3.75)	– ⁱ	2.15 (0.60–7.70)	1.60 (0.79–3.23)
Co-morbid substance dependence (ref = no) ^h	1.63 (0.59–4.47)	1.89 (0.64–5.59)	1.31 (0.17–10.12)	– ⁱ	1.40 (0.62–3.14)
Chi-square	20.3 (13)	63.1 (12)***	14.9 (12)	10.0 (10)	41.3 (14)***
<i>n</i> events	50	44	17	16	83
<i>n</i> censored	89	117	27	37	216
<i>n</i> total	139	161	44	53	299

DSM-IV, Diagnostic and Statistical Manual of Mental Disorders, fourth edition; HR, Hazard ratio; 95%CI, 95% confidence interval; IQ, intelligence quotient; SEP, socio-economic position.

^aTime-to-treatment for disorders with initial treatment contact before the year of onset of the respective disorder set to 0.

^bLifetime treatment contact and age of initial treatment contact for secondary mental health care based on the Psychiatric Case Register North Netherlands.

^cCovariate excluded due to insufficient cases.

^dAll substance dependence diagnoses were considered severe.

^eNo cases, because adolescents with any disorder onset before 2000 were excluded.

^fCombined with age of onset 11–15 years due to insufficient cases.

^gCombined with age of onset 16–20 years (reference category) due to insufficient cases in the reference category.

^hTime-dependent covariate.

ⁱDisorder class is the dependent variable.

* $p < 0.05$ ** $p < 0.01$ *** $p < 0.001$.

Time-to-treatment

A comparison between our findings on time-to-treatment for mental disorders in adolescents and prior studies conducted in adults yields two main observations. First, the time-to-treatment was shorter in our adolescent sample than in comparable adult samples (Wang *et al.* 2004, 2005, 2007; Bruffaerts *et al.* 2007; ten Have *et al.* 2013a). However, estimates of time-to-treatment are highly dependent on follow-up time, which was substantially shorter in our sample of adolescents than in the adult samples. Second, the patterns of time-to-treatment in adolescents are remarkably similar to those found in Dutch (ten Have *et al.* 2013a), Belgian (Bruffaerts *et al.* 2007), Northern Irish (Bunting *et al.* 2012), and American (Wang *et al.* 2005) adults. For instance, mood disorders are characterised by high proportions of lifetime treatment contact and a relatively short time-to-treatment in all ages, while within the class of anxiety disorders initial treatment contact is made most and fastest for panic disorder, and least and slowest for specific phobia. Although differences among countries do exist (Wang *et al.* 2007), we expect to find highly similar patterns of time-to-treatment in adolescents from other countries as well.

Predictors of time-to-treatment

Following studies using adult community samples (Kessler *et al.* 1998; Wang *et al.* 2004, 2005, 2007; Bruffaerts *et al.* 2007; Korczak & Goldstein, 2009; ten Have *et al.* 2013a), we found that time-to-treatment is longer when disorders have an onset earlier in life. We were able to confirm a robust association between earlier onset and a longer time-to-treatment. An important reason for this age of onset-effect is that children's access to mental health care depends on recognition and help-seeking by their parents or teachers (Wang *et al.* 2005; Bruffaerts *et al.* 2007; ten Have *et al.* 2013a). Symptoms from early-onset disorders may not be recognised because they may be considered as being a part of a child's identity, are not severe enough, or are not disturbing enough to the social environment (Wang *et al.* 2005; ten Have *et al.* 2013a; Jörg *et al.* 2015), potentially resulting in unmet need. The development of coping strategies may mitigate or even eliminate the need for treatment, at least in the short term, until adolescents enter a life phase during which they have to be more self-reliant. In adults, early-onset disorders have been associated with a time-to-treatment of decades (Kessler *et al.* 1998; Christiana *et al.* 2000; Wang *et al.* 2004, 2005; Bunting *et al.* 2012). This at least raises the questions of how potentially harmful a long time-to-treatment for

early-onset disorders is, and whether prevention and early intervention programmes aimed specifically at children and adolescents should be employed (Merikangas *et al.* 2010; Bunting *et al.* 2012; de Girolamo *et al.* 2012; Thornicroft, 2012; Ghio *et al.* 2014).

The current study added to the literature by including severity and co-morbidity as predictors of time-to-treatment. Co-morbid mood disorders most often predicted shorter time-to-treatment, but co-morbidity from other classes was mostly unrelated to time-to-treatment for any care. Maybe only co-morbid disorders with a short time-to-treatment themselves, such as dysthymia and panic disorder, accelerate the time-to-treatment for other disorders, as opposed to for instance social and specific phobia (Olfson *et al.* 2012). Alternatively, perhaps the onset of a co-morbid disorder prompts treatment seeking for the co-morbid disorder, rather than for the index disorder (Chapman *et al.* 2015). That co-morbidity tended to be a stronger predictor for secondary care than for any care, was probably because treatment contact in secondary care could not be attributed to any disorder in particular.

Disorder severity, operationalised in this study as high levels of impairment or distress, predicted shorter time-to-treatment for any care. This is largely in line with a previous finding that symptoms of functional impairment predicted shorter time-to-treatment for alcohol dependence, whereas the number of dependence symptoms did not (Chapman *et al.* 2015). Unexpectedly, disorder severity was not associated with time-to-treatment for secondary care. Adolescents whose first disorder had an onset approximately before the age of 10 did not have a severe disorder more often than did adolescents whose first disorder had an onset later in life. They did have treatment contact with secondary care more often, and they showed more signs of multimorbidity. This might indicate that the time-to-treatment with secondary care is reduced by the complexity of psychopathology, rather than the levels of impairment or distress.

Secondary care

The results for second care were largely similar to those for any care. The sensitivity analyses for any care shared many characteristics with the Cox regression analyses for secondary care, and lead to the same substantive conclusions. We therefore think that the latter suffered from reduced statistical power, but not reduced precision of estimates.

An interesting finding was that time-to-treatment for secondary care was shorter for adolescents from a low or middle than for adolescents from a high socio-economic background, while no such pattern was

found for any care. As a high socioeconomic background has been associated with more parent reported specialist mental health care use (Amone-P'Olak et al. 2010), parents from a high socioeconomic background may prefer to send their children to other types of care, such as private practices.

Concluding remark

This study is, as far as we know, the first to describe time-to-treatment and its correlates for lifetime mental disorders in a large cohort of adolescents. The differentiation between any care and secondary care, and the inclusion of disorder severity and co-morbidity as predictors of time-to-treatment add further relevance to this study. Time-to-treatment is already substantial in adolescence, and shows patterns highly similar to those observed in adult samples, which confirms the importance of focusing on childhood and adolescence for the reduction of time-to-treatment. Next to age of onset, only disorder severity and co-morbidity are consistently related to time-to-treatment. This suggests that the characteristics of psychopathology are more important correlates of time-to-treatment than the background variables that are generally included yet hardly produce consistent findings, such as family characteristics. For a better comprehension of time-to-treatment, future studies should ideally address theoretical explanations of time-to-treatment, such as parental recognition, coping and unmet need, as well as the outcomes of time-to-treatment, such as social functioning and educational attainment.

Supplementary material

The supplementary material for this article can be found at <http://dx.doi.org/10.1017/S2045796016000226>.

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Conflict of Interest

None.

Ethical Standard

The authors assert that all procedures contributing to this work comply with the ethical standards of the relevant national and institutional committees on human experimentation and with the Helsinki Declaration of 1975, as revised in 2008.

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