Commonly occurring adversities in families as risk factors for developing psychosocial and psychiatric morbidities: evidence from general practice

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Background
Childhood adversity may lead to mental and somatic complications throughout life. General practitioners are equipped to identify and manage adverse events in households. The relationship between adversities and psychiatric symptoms has not been studied in primary care.

Aims
We investigated the relationship of common adversities in families with respect to subsequent development of psychosocial and psychiatric problems in young children.

Method
We analysed data from seven general practices, including participants between 0 and 9 years of age. Adversity was defined as having a household member who was diagnosed with cancer, psychiatric disease or social problems. We compared these patients with controls matched for gender, age and general practice. The primary outcome was any new episode defined with a psychological and psychiatric label. Secondarily, the encounter rates at the general practices after adversity were analysed.

Results
Participants in both groups were followed for an average of 12 years, whereby patients with an adversity were more likely to develop psychiatric morbidities compared with matched references (odds ratio 1.38, 95% CI 1.12–1.68, P = 0.002), also revealing higher encounter rates at general practices. We found no statistically significant association between adversities in the family and increased psychosocial symptoms.

Conclusions
The short- and long-term consequences of exposure to negative events in childhood are of great public health importance. Our data suggest screening more proactively for consequences of commonly occurring adversities in families, as they are a risk factor for subsequent psychiatric symptoms. Enhanced consultation frequency at general practitioners following adversities should be differentiated in more detail.

Keywords
Childhood experience; comorbidity; outcome studies; primary care; risk assessment.

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There are significant relationships between adverse life events, psychosocial resources and mental and somatic well-being throughout the lifespan. The earlier adverse events occur in life, the more likely they are to affect all subsequent levels of mental and physical development. Childhood adversity promotes subsequent health problems through pathways, including physiological challenge of the hypothalamic–pituitary–adrenal axis and social mechanisms, both of which affect the brain development (see Nelson et al.1 for a recent review). It has been acknowledged that mental health services for practitioners (GPs) can take to help improve outcomes.5

Early identification is thus an important prerequisite for a successful prevention of long-term consequences of childhood adversity in children. Recently, Nelson and colleagues1 have suggested to broaden assessment of interventions beyond mental health measures, and include stress-related health outcomes such as asthma, infection, inflammation and insulin resistance. Primary care in general practices could play an important role in this,4 and there are several actions that general practitioners (GPs) can take to help improve outcomes.5

Thus far, the emphasis in monitoring in GP practices seems to have been on more severe adversity,2 but translational studies have shown that more subtle indirect adversities may also do great harm and are possibly not screened for often enough. Among more

general psychosocial circumstances, cancer in household members has been identified to be an important risk factors (for review, see Walczak et al.1). For example, Huizinga and colleagues7 have shown that parental cancer is a significant inducer of psychological fragility in terms of internalising and cognitive problems in children. Likewise, the impact of severe family distress owing to parental psychiatric disorders has been investigated, often with respect to the gene×environment interaction effect on developing the same problems later in life.8 Other more general psychosocial problems include, for example, stress of parents at work. Moreover, we know that limited financial resources and workplaces with fewer family-friendly policies are risk factors for family life and child development.9

Aims
The overall goal of this study was to investigate the difference in occurrence of psychosocial and psychiatric problems in children who experienced indirect forms of frequently occurring early adversity, compared with matched controls. To realise this aim, we analysed data from a prospective database of GPs, allowing us to look at the course of these patients in their primary care setting. The form of registration in primary care allowed us not only to identify episodes of morbidities after adversity that were registered in primary care,
Table 1 Overview of the classification of the indirect adversities, according to the International Classification of Primary Care, Second Edition (ICPC-2)

| Code | Definition |
|------|------------|
| 1. Cancer in household members | **A79** Malignancy not otherwise specified  
**B72** Hodgkin’s disease/lymphoma  
**B73** Leukaemia  
**B74** Malignant neoplasm blood other  
**D74** Malignant neoplasm stomach  
**D75** Malignant neoplasm colon/rectum  
**D76** Malignant neoplasm pancreas  
**D77** Malignant neoplasm digest other/not otherwise specified  
**F74** Neoplasm of eye/adnexa  
**H75** Neoplasm of ear  
**K72** Neoplasm cardiovascular  
**L71** Malignant neoplasm musculoskeletal  
**N74** Malignant neoplasm nervous system  
**N76** Neoplasm nervous system unspecified  
**R84** Malignant neoplasm bronchus/lung  
**R85** Malignant neoplasm respiratory, other  
**S77** Malignant neoplasm of skin  
**T71** Malignant neoplasm thyroid  
**U75** Malignant neoplasm of kidney  
**U76** Malignant neoplasm of bladder  
**U77** Malignant neoplasm urinary tract  
**X75** Malignant neoplasm cervix  
**X76** Malignant neoplasm breast female  
**X77** Malignant neoplasm genit التسلسل  
**Y78** Malignant neoplasm female genital other  
**Z12** Relationship problem with partner  
**Z13** Partner’s behaviour problem  
**Z14** Partner illness problem  
**Z15** Loss/death of partner problem  
**Z16** Relationship problem with child  
**Z18** Illness problem with child  
**Z19** Loss/death of child problem  
**Z20** Relationship problem parent/family  
**Z21** Behaviour problem parent/family  
**Z22** Illness problem parent/family  
**Z23** Loss/death parent/family member  
**Z24** Relationship problem friend  
**Z25** Assault/harmful event problem  
**Z27** Fear of a social problem  
**Z29** Social problem not otherwise specified  |
| 2. Psychiatric diagnoses of household members | **P71** Organic psychosis other  
**P72** Schizophrenia  
**P73** Affective psychosis  
**P74** Anxiety disorder/anxiety state  
**P77** Suicide/suicide attempt  
**P80** Personality disorder  
**P82** Post-traumatic stress disorder  
**P86** Psychological disorders, other  |
| 3. Social problems of household members | **A94** Death  
**Z01** Poverty/financial problem  
**Z02** Food/water problem  
**Z03** Housing/neighbourhood problem  
**Z04** Social cultural problem  
**Z05** Work problem  
**Z06** Unemployment problem  
**Z07** Education problem  
**Z08** Social welfare problem  
**Z09** Legal problem  
**Z10** Healthcare system problem  
**Z11** Compliance/being ill problem  
**Z12** Relationship problem with partner  
**Z13** Partner’s behaviour problem  
**Z14** Partner illness problem  
**Z15** Loss/death of partner problem  
**Z16** Relationship problem with child  
**Z18** Illness problem with child  
**Z19** Loss/death of child problem  
**Z20** Relationship problem parent/family  
**Z21** Behaviour problem parent/family  
**Z22** Illness problem parent/family  
**Z23** Loss/death parent/family member  
**Z24** Relationship problem friend  
**Z25** Assault/harmful event problem  
**Z27** Fear of a social problem  
**Z29** Social problem not otherwise specified  |

Method

Data source

This retrospective study of prospectively collected data was conducted on the basis of the Family Medicine Network (FaMe-Net database16) in The Netherlands. The Dutch Practice-Based Research Network (PBRN) FaMe-Net, the world’s oldest PBRN, has a long history of systematically recording all morbidity presented to the family physician in episodes. FaMe-Net is a practice-based research network from the Department of Primary and Community Care at the Radboud University Medical Center, which aims to contribute to research and education and thereby improve the quality of primary care. FaMe-Net—associated physicians systematically and prospectively register data electronically on the reason for encounter, diagnostic procedures, diagnoses, interventions and referrals. All data in this database are anonymised and patients in FaMe-Net gave permission to use data for research purposes by an opting-out procedure. All doctors are trained in registration as part of belonging to the network. There is regular supervision and quality control on the registration of the data. Diagnosis in all patient encounters are coded by the GP, according to the International Classification of Primary Care (ICPC-211), extended by the ICD-10.12 In this network, which consists of seven Dutch family practices in the East Netherlands, all encounters between family practices and patients are registered since 1971. All information belonging to one health problem is ordered in one episode, whereby information from other institutions and specialised diagnoses are included to support diagnoses.

Extraction rules

In this study, patients were included who encountered an indirect adversity before the age of 9 years. To this end, data were extracted from 1 January 1995 until 31 December 2016, whereby a potential adversity occurred before 2013, such that minimum age of inclusion was 4 years. We defined an indirect adversity as the patient having a household member diagnosed with one of the following conditions: cancer, a severe psychiatric diagnosis or social problems (indexed in Table 1). Because of the given evidence in the literature, we defined a hierarchy whereby cancer was taken as the primary adversity, followed by psychiatry diagnosis and then social problems, which are more frequently reported in GP practices and can be the least specific in a database. To account for potential mistakes owing to double registrations, we selected one household member (i.e. family member) with an indirect adversity for each patient. The follow-up period was determined as a minimum of 4 years since the first experience of adversity. An event for a family member subsequently leading to indirect adversity in our patient group could therefore potentially have occurred in the recent period before the birth of that patient, so that the adversity ‘started’ at the age of 0 days.

The control group were matched to the index group for gender, age and the general practices. Matched references were not diagnosed with cancer or a social problem, and did not have a family member that was diagnosed with cancer, a psychiatric disease or a social problem. Importantly, also with respect to the time frame of the observed results, the matched reference was selected based...
on the date of indirect adversity diagnosis of the index patient, to analyse potential psychosocial and psychiatric problems in the same time period. We included index patients with a minimum of one and maximum of four matched control patients, to have more data to calculate statistics.

Extracted data for this study were age, gender, practice, diagnosis coded by ICPC codes, date of diagnosis, date of practice registration or deregistration, and the reason for deregistration.

The Radboud University Medical Center’s Technology Center Health Data supports FaMe-Net in distillation and secure storage of routine data from the affiliated general practices. It adheres to the regulations of Dutch and European laws, and has gained ethical approval from the research ethics committee of Radboud University Medical Center for this procedure (medical ethics number 2020-6871).

Analyses

The primary outcome variables were new ICPC-2 ‘episodes’ (for an overview, see https://ehelse.no/kodeverk/icpc-2e-english-version),11 We distinguished episodes with psychiatric or psychosocial symptoms only (ICPC codes P1–P29), and episodes resulting in a psychiatric diagnosis (ICPC codes P70–P99), as can be found in Table 2. We selected the first psychiatric and psychosocial complications that were reported in the follow-up period after the adversity, whereby we adhered to the hierarchy that subsequent report of psychiatric diagnoses was noted before psychosocial symptoms, as this was thought to reflect the more clinically relevant outcome.

The second outcome variable was the number of visits at the GP practice in a follow-up period of at least 4 years since the first adversity event. Consultations for all different ICPC codes were counted for this rate. We also calculated the different follow-up length in terms of years after the index event.

Data management and analysis were performed with SPSS version 22.0 (SPSS Inc., Chicago, Illinois, USA) software for Windows. Descriptive statistics were used to describe the main findings. Given that our primary outcome was associated with a dependent variable with two possible values (yes/no psychiatric diagnosis), we opted for a binary logistic regression model to compare our group of index patients with control patients. Age and gender were taken as confounding variables and were corrected for. Negative binomial regression analysis was used to test the difference in number of encounter between index patients and control patients. A P-value of <0.05 was considered to be statistically significant, based on two-sided tests.

Results

The whole database consisted of 28 659 patients from all seven practices. Based on the aforementioned selection criteria, we included 1029 patients and 3849 controls. Participants encountering adversity were followed around 12 years in the same GP practice, as shown in Table 3. There were slightly more males than females in the sample, with a median age of experienced adversity of around 4 years (Table 3). Note, however, an indirect adversity could also occur before the birth of the index patient, which was the case in 70 out of 1029 patients, in which severe social problems were the most common (see Table 3).

Results from our primary question can be found in Table 2. Patients with indirect adversity are significantly more likely to develop psychiatric symptoms compared with controls (odds ratio 1.38, 95% CI 1.12–1.68; P = 0.002). We found no statistically significant association between indirect adversities and the onset of psychosocial symptoms (odds ratio 1.04, 95% CI 0.89–1.21; P = 0.620). Moreover, patients with indirect adversities demonstrated significantly higher encounter rates (hazard ratio 1.19, 95% CI 1.10–1.28; P ≤ 0.001) compared with matched controls (mean encounter per year 16.27 (s.d. 15.53) vs. 13.94 (s.d. 13.13).

When looking at the different psychosocial and psychiatric problems (Table 4) in both groups, we found that having behavioural problems, specific intellectual disabilities and feeling anxious were the most common problems in this follow-up period. This was mirrored by the psychiatric morbidities, whereby attention-deficit hyperactivity disorder and anxiety were most common.

Discussion

This is, to the best of our knowledge, the first study in a primary care setting that has prospectively investigated the effect of commonly
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Table 4 Absolute frequencies and percentage of psychosocial symptoms and psychiatric morbidity according to the International Classification of Primary Care in patients that experienced indirect adversities and their matched controls

| Indirect adversity group (n = 1029) | Control group (n = 3849) |
|------------------------------------|-------------------------|
| Any symptom or morbidity           | 542 (52.7)              |
| Psychosocial symptoms              | 1582 (41.0)             |
| P01 Feeling anxious/tense          | 43 (4.2)                |
|                                   | 155 (4.0)               |
| P02 Acute stress reaction          | 8 (0.8)                 |
|                                   | 38 (1.0)                |
| P03 Feeling depressed              | 18 (1.7)                |
|                                   | 47 (1.2)                |
| P04 Feeling/behaving irritable/angry| 12 (1.2)               |
|                                   | 23 (0.6)                |
| P06 Sexual fulfilment reduced      | 1 (0.1)                 |
|                                   | 4 (0.1)                 |
| P09 Sexual preference concern      | 1 (0.1)                 |
|                                   | 1 (0.03)                |
| P11 Eating problem with child      | 7 (0.7)                 |
|                                   | 27 (0.7)                |
| P22 Child behaviour symptom/complaint | 108 (10.5)          |
|                                   | 292 (7.6)               |
| P23 Adolescent behaviour symptom/complaint | 5 (0.5) |
|                                   | 33 (0.9)                |
| P24 Specific learning problem      | 129 (12.5)              |
|                                   | 444 (11.5)              |
| P25 Phase of life problem adult    | 3 (0.3)                 |
|                                   | 9 (0.2)                 |
| P29 Psychological symptom other    | 23 (2.2)                |
|                                   | 44 (1.1)                |
| Psychiatric morbidity              |                         |
| P15 Chronic alcohol abuse          | 0                       |
|                                   | 3 (0.08)                |
| P18 Medication abuse               | 2 (0.2)                 |
|                                   | 2 (0.05)                |
| P19 Drug abuse                     | 7 (0.7)                 |
|                                   | 19 (0.3)                |
| P72 Schizophrenia                  | 0                       |
|                                   | 0                       |
| P73 Affective psychosis            | 0                       |
|                                   | 2 (0.05)                |
| P74 Anxiety disorder/anxiety state | 26 (2.5)                |
|                                   | 74 (1.9)                |
| P75 Somatisation disorder          | 3 (0.3)                 |
|                                   | 5 (0.1)                 |
| P76 Depressive disorder            | 19 (1.8)                |
|                                   | 56 (1.3)                |
| P77 Suicide/suicide attempt        | 0                       |
|                                   | 1 (0.03)                |
| P78 Neurasthenia/surmenage         | 5 (0.5)                 |
|                                   | 14 (0.4)                |
| P79 Phobia/composite disorder      | 5 (0.5)                 |
|                                   | 18 (0.3)                |
| P80 Personality disorder           | 3 (0.3)                 |
|                                   | 9 (0.2)                 |
| P81 Hyperkinetic disorder          | 81 (7.9)                |
|                                   | 175 (4.5)               |
| (attention-deficit hyperactivity disorder) |               |
| P82 Post-traumatic stress disorder | 2 (0.2)                 |
|                                   | 7 (0.2)                 |
| P86 Anorexia nervosa/bulimia       | 0                       |
|                                   | 9 (0.2)                 |
| P98 Psychosis not otherwise        | 3 (0.3)                 |
|                                   | 3 (0.08)                |
| specified/other                    |                         |
| P99 Psychological disorders, other | 28 (2.7)                |
|                                   | 68 (1.8)                |

a. For the control group, only the first symptom diagnosis was extracted for this study.

occurring indirect adversities before 9 years of age on the subsequent development of psychosocial problems and psychiatric morbidity. Using a large sample of index samples and matched controls, we were able to support our hypothesis that commonly occurring adversities are related to mental stability in young children. More precisely, we found that children with a parent with cancer, a psychiatric disease or a social problem were more likely to develop psychiatric morbidity, compared with controls from the similar primary care setting and thereby likely share the same socioeconomic setting. Hyperkinetic disorders (P81), followed by anxiety disorders (P74) and then depressive disorder (P76) were most commonly registered in the follow-up course for both groups, mimicking the prevalence rate of so-called internalising disorders at this time in the life course. Systematic reviews suggest that the reported range in the community prevalence of attention-deficit hyperactivity disorder (2.2%–7.2%) reflects variation in study methodology, and the prevalence rate in our control group is in line with this prevalence.13 Bias in the registration of disorders cannot be entirely ruled out, but we do not see a higher prevalence of developmental disorders in the control group. When comparing the distribution in the indirect adversity and control groups, indirect adversity was not related to another distribution of psychosocial and psychiatric complications in childhood and adolescence, but was related to a higher frequency. Finally, mean encounter rates per year at primary care centres were significant higher in participants that had experienced adversities, compared with controls. Unfortunately, we were not able to differentiate whether adversity led to greater need for care in the context of psychiatric complications or somatic problems, which are also well-documented consequences. We cannot rule out that as a result of more encounters, there was a higher likelihood of detecting problems within primary care in the indirect adversity group. However, the method of registration in the GP practices allowed us not only to identify post-adversity morbidity episodes that were registered in primary care, but also all morbidity episodes that were registered in secondary and tertiary care and reported back to the GP, so that it seems likely that these are valid diagnoses.

The onset of mental disorders in those that have experienced childhood adversity has been extensively studied with retrospective questionnaires that cover a wide array of adversities. Childhood adversity has been linked to the onset of different dimensions of stress-related psychopathology, such as depression, schizophrenia, severity of bipolar disorder and increased risk of psychosis.14–16 Focusing on more frequently occurring adversities, we can support this notion from the perspective of a GP, at least in the observed period of approximately 12 years after a reported episode. Although there were limitations in assessing all kinds of chronic stress in the families, our findings support other studies showing that children with parental cancer in the age group of 11–23 years have reported internalising and cognitive problems.7 It has been acknowledged that GPs should be alert for somatic and psychosocial problems in partners of patients with cancer, but studies prospectively looking into the effects on children are sparse. Thastum and colleagues17 investigated a cohort of children and adolescents whose parents were suffering from cancer. The authors showed a higher risk of problems, particularly when the father was ill, but it remains unclear whether this difference was because of the different diagnoses of fathers and mothers, gender or other factors. Moreover, internalising problems in children and adolescents were best predicted by parental depression, whereas family dysfunction was related to externalising problems in the offspring. Chen and Parebianco18 showed that emotional well-being of ill parents was directly associated with adolescent distress, which also led to mental health problems later in development.

In the present study, we explicitly tried to include the concept of indirect adversity because longer-lasting stressful events in the household are an important risk factor even when children are not directly affected. Our study is limited by the fact that the relationship between the affected child and household members is not well defined. Yet, we know that stress impact also occurs as any household member may induce psychological and health problems. Of course, we have to acknowledge that the concept of an indirect adversity is difficult to define in a naturalistic cohort setting. We cannot clearly identify the length and severity of the indirect adversity period. Indirect adversity, in our case, could even occur before birth, and thus longer-lasting programming effects on the developing brain may have effects ranging from stunted physical growth and cognitive delays to problems regulating attention.19 As is also evident from the demographics, most of the patients we investigated were still in their early adolescence during the time of the follow-up in this study. Based on the epidemiological evidence, it is likely that more mental health problems occur later on in life, so we can only reflect on the impact at that point. From naturalistic cohorts in adulthood, we know that the frequency of childhood adversity of any kind was positively associated with psychiatric comorbidity.20

Currently, interventions informed by specific trauma in early life have not yet been adapted from mental health settings for use in primary care. A few studies have investigated the use of additional screening lists in primary care. Notably, however, such
questionnaires mainly covered more severe forms of adversity and not the common psychosocial episodes we have investigated. For example, Glowa and colleagues used a ten-item childhood adversity questionnaire as a screening tool in a family practice setting, with patients presenting for follow-up of chronic illness or annual physicals. Based on the sum score, they divided patients with higher and lower risk scores and found that based on the outcome scores, clinicians were more likely to have discussed adversity issues for high-risk patients. If these data could be replicated in larger samples, they could lead to policy changes in staging risks after such events, promoting more preventive actions in primary care. This is of relevance because in many countries, secondary mental healthcare comes with long waiting lists and area-specific availability. As there is an ongoing concern that longer waiting times for treatment leads to poorer health outcomes, earlier interventions in primary care, if possible, could circumvent side-effects of restricted specialised care. In fact, all of the psychiatric morbidities we found in the follow-up period of the children with indirect adversity show high comorbidity with somatic pathology, such as cardiovascular disease, rheumatoid arthritis and diabetes. Such comorbidity is associated with unfavourable outcomes for the individual, such as low quality of life and mortality. Not surprisingly, it leads to high levels of healthcare utilisation, which was also supported by increased encounter rates with the GP practices in our data-set.

Strengths and limitations

A strength of this study is the large size of our study population and the longitudinal primary care data investigating the effect of childhood adversity on psychiatric morbidities and psychosocial symptoms, using ICPC-2 codes. Many studies have investigated the effect of adversity concerning only one psychiatric diagnosis, e.g. depression, schizophrenia, bipolar disorder and psychosis. Lastly, the follow-up period of this study has an average of 12 years, which is long enough to provide clear information and draw conclusions about our findings.

This study also has a number of limitations. We included residential connection members (e.g. investigating parental cancer), assuming that they are family members. However, the precise relationship of a patient with their household member is not registered in our database. Yet, we also know that stress effects can occur more broadly within a whole household/family, and in this pilot analysis, we particularly focused on more severe and disabling psychiatric diagnoses of household members. Depression and substance use disorders are also highly prevalent and can be disabling, but in the ICPC system, no distinction is possible between (frequently occurring) mild depression and the level of alcohol misuse. Future studies will need to disentangle risk for indirect adversity in more detail. To assess potential causality in more depth, these studies should prospectively collect more detailed data on family and socioeconomic circumstances, as well as the impact of indirect adversity (e.g. severity of the event). It is known that all patients lived with the person experiencing adversity. Moreover, a household member in the context of the registration in the GP system is, per definition, meaningful because they live together and the definition of a household member within the GP system excluded incidental circumstances where, for example, somebody else is temporarily living as a lodger at the same address. Another limitation is that the encounter rates for different adversity groups could be overestimated, since the registration date of an ICPC-2 code is used as the start date of an adversity. Selecting only one household member (i.e. family member) with indirect adversity for each patient did not allow us to dissociate the quantitative effects in further detail, such that other family members could have also experienced an event that may have led to indirect adversity. Further, the comparison between absolute frequencies of symptoms diagnoses between the two groups was hampered by the fact that for the control group, only the first diagnosis was extracted. However, this did not affect the primary outcome because this was based on the occurrence of any first outcome. Finally, a limitation can be the possibility of incomplete data entry, as in any large administrative data-set; however, the necessity for GPs to indicate the episode correctly with regards to qualitative checks and financial controlling helps to minimise this risk.

In sum, our data show that common indirect adversities a child can experience in their household is an important risk factor for developing psychosocial and psychiatric morbidities in later childhood and adolescence. Further research also using prediction models of these more common indirect adversity types would be of value to determine the high-risk population that would benefit from direct intervention in primary and potentially mental healthcare. In particular, our results may be of relevance in countries with a strong primary care system, such as the UK and The Netherlands. The findings support a systemic approach for both adults who present with cancer, severe psychiatric disorders and social problems, and for the children who may present to a higher extent with associated psychiatric problems. Moreover, psychiatric expertise might be helpful in caring for families who experience adversity, thus making a case for interdisciplinary care.

Data availability

The data that support the findings of this study are available from the corresponding author, I.T., upon reasonable request. The data are not publicly available as they contain information that could compromise the privacy of research participants.

Author contributions

I.T., T.P., R.A. and F.v.d.L. were responsible for study conceptualisation and design. I.T. and F.v.d.L. formulated the specific research question. T.P. conducted the statistical analyses under supervision of I.T., R.A. and F.v.d.L. H.P. and R.A. were responsible for the data extraction and management, quality control and manuscript preparation. All authors contributed to writing and/or editing of the manuscript and approved the final version.

Funding

This research received no specific grant from any funding agency, commercial or not-for-profit sectors.

Declaration of interest

None.
References

1. Nelson CA, Scott RD, Bhutta ZA, Harris NB, Danese A, Samer AM. Adversity in childhood is linked to mental and physical health throughout life. BMJ 2020; 371: m3048.

2. Danese A, Smith P, Chitsabesan P, Dubicka B. Child and adolescent mental health amidst emergencies and disasters. Br J Psychiatry 2020; 216(3): 159–62.

3. Kessler R, Berglund P, Demler O, Jin O, Merikangas KR, Walters EE. Lifetime prevalence and age-of-onset distributions of DSM-IV disorders in national comorbidity survey replication. Arch Gen Psychiatry 2005; 62(6): 593–602.

4. Mughal F, England E. The mental health of young people: the view from primary care. Br J Gen Pract 2016; 66(651): 502–3.

5. Mail EJ, Hargraves DS. The UK's global standing in child and young people's health and how GPs can help. Br J Gen Pract 2020; 70(693): 156–7.

6. Waiczak A, McDonald F, Patterson P, Dobinson K, Allison K. How does parental cancer affect adolescent and young adult offspring? A systematic review. Int J Nurs Stud 2018; 77: 54–80.

7. Hulzinga GA, Visser A, van der Graaf WT, Hoekstra HJ, Klijn EC, Pras E, et al. Stress response symptoms in adolescent and young adult children of parents diagnosed with cancer. Eur J Cancer 2005; 41(2): 288–95.

8. Shalev A, Merranko J, Goldstein M, Goldstein Bl, et al. A longitudinal study of family functioning in offspring of parents diagnosed with bipolar disorder. J Am Acad Child Adolesc Psychiatry 2019; 58(10): 961–70.

9. Perry-Jenkins M, Laws HB, Sayer A, Newkirk K. Parents’ work and children’s development: a longitudinal investigation of working-class families. J Fam Psychol 2020; 34(3): 257–68.

10. Luijks H, Van Boven K, Olde Hartman T, Uijen A, Van Weel C, Schers H. Making of children is linked to mental and physical health throughout life. BMJ 2020; 371: m3048.

11. Luijks H, Van Boven K, Olde Hartman T, Uijen A, Van Weel C, Schers H. Making of children is linked to mental and physical health throughout life. BMJ 2020; 371: m3048.

12. Nelson CA, Scott RD, Bhutta ZA, Harris NB, Danese A, Samer AM. Adversity in childhood is linked to mental and physical health throughout life. BMJ 2020; 371: m3048.

13. Sayal K, Prasad V, Daley D, Ford T, Coghill D. ADHD in children and young people: prevalence, care pathways, and service provision. Lancet Psychiatry 2018; 5(2): 175–86.

14. Matheson SL, Shepherd AM, Pinchbeck RM, Laurens KR, Carr VI. Childhood adversity in schizophrenia: a systematic meta-analysis. Psychol Med 2013; 43(2): 225–38.

15. Nanni V, Uher R, Danese A. Childhood maltreatment predicts unfavorable course of illness and treatment outcome in depression: a meta-analysis. Am J Psychiatry 2012; 169(2): 141–51.

16. Varese F, Smeets F, Drukker M, Lieverse R, Lataster T, Viechtbauer W, et al. Childhood adversities increase the risk of psychosis: a meta-analysis of patient-control, prospective and cross-sectional cohort studies. Schizophr Bull 2012; 38(4): 661–71.

17. Thastum M, Watson M, Kienbacher C, Piha J, Steck B, Zachariae R, et al. Prevalence and predictors of emotional and behavioural functioning of children where a parent has cancer: a multinational study. Cancer 2009; 115(17): 4030–9.

18. Chen CY, Panebianco A. Physical and psychological conditions of parental chronic illness, parentification and adolescent psychological adjustment. Psychol Health 2020; 35(9): 1075–94.

19. van Izendoorn MH, Bakermans-Kranenburg MJ, Duschinsky R, Fox NA, Goldman PS, Gunnar MR, et al. Institutionalisation and deinstitutionalisation of children 1: a systematic and integrative review of evidence regarding effects on development. Lancet Psychiatry 2020; 7(6): 703–20.

20. Vrijen JW, van Amer CT, Koekkoek B, van Oostrom I, Schene AH, Tendolkar I. Childhood trauma and negative memory bias as shared risk factors for psycho-pathology and comorbidity in a naturalistic psychiatric patient sample. Brain Behav 2018; 8(12): e01181.

21. Glowa PT, Olson AL, Johnson DJ. Screening of adverse childhood experiences in a family medicine setting: a feasibility study. J Am Board Fam Med 2016; 29(3): 303–7.

22. Cohen BE, Edmondson D, Kronish IM. State of the art review: depression, stress, anxiety, and cardiovascular disease. Am J Hypertension 2015; 28(11): 1295–302.

23. Marrie RA, Walla R, Bolton JM, Sareen J, Walker JR, Palten SB, et al. CIHR team in defining the burden and managing the effects of psychiatric comorbidity in chronic immunoinflammatory disease. Increased incidence of psychiatric disorders in immune-mediated inflammatory disease. J Psych Res 2017; 101: 17–23.

24. Schram MT, Baan CA, Pouwer F. Depression and quality of life in patients with diabetes: a systematic review from the European Depression in Diabetes (EDID) Research Consortium. Curr Diabetes Rev 2009; 5(2): 112–9.

25. van Dooren FE, Nefs G, Schram MT, Verhey FR, Denollet J, Pouwer F. Depression and risk of mortality in people with diabetes mellitus: a systematic review and meta-analysis. PLoS One 2013; 8(3): e57058.

26. Wagner JA, Pietrzak RH, Petry NM. Psychiatric disorders are associated with hospital care utilization in persons with hypertension. Soc Psychiatr Epi 2008; 43(11): 878–8.

27. Giannelli A, Palmos A, Hagaenaas SP, Breen G, Lewis CM, Mutz J. Examining the association between family status and depression in the UK Biobank. J Affect Disord 2021; 279: 585–98.

28. Rayner C, Coleman JR, Purves KL, Cheesman R, Hübel C, Gaspar H, et al. Genetic influences on treatment-seeking for common mental health problems in the UK Biobank. Behav Res Ther 2019; 121: 103413.