

Differing impact of the COVID-19 pandemic on youth mental health: combined population and clinical study

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Background

Identifying youths most at risk to COVID-19-related mental illness is essential for the development of effective targeted interventions.

Aims

To compare trajectories of mental health throughout the pandemic in youth with and without prior mental illness and identify those most at risk of COVID-19-related mental illness.

Method

Data were collected from individuals aged 18–26 years (N = 669) from two existing cohorts: IMAGEN, a population-based cohort; and ESTRA/STRATIFY, clinical cohorts of individuals with preexisting diagnoses of mental disorders. Repeated COVID-19 surveys and standardised mental health assessments were used to compare trajectories of mental health symptoms from before the pandemic through to the second lockdown.

Results

Mental health trajectories differed significantly between cohorts. In the population cohort, depression and eating disorder symptoms increased by 33.9% (95% CI 31.78–36.57) and 15.6% (95% CI 15.39–15.68) during the pandemic, respectively. By contrast, these remained high over time in the clinical cohort. Conversely, trajectories of alcohol misuse were similar in both cohorts, decreasing continuously (a 15.2% decrease) during the

COVID-19 had detrimental effects on mental health, with worldwide rates of major depressive disorders (MDD) and anxiety disorders rising to 27.5 and 25.6%, respectively.¹ Fear of the virus itself and lockdowns implemented by governments around the globe have caused greater mental distress and lower quality of life in the general population.^{2–5} In particularly, young people, who are known to experience major social role transitions,⁶ experienced higher levels of depressive and anxiety symptoms than people in older age groups.^{1,4,5} The pandemic has also been reported to worsen symptoms of patients with pre-existing mental illness,^{7,8} although contradictory findings have been reported.^{9–12} These contradictions and the limitations of studies to date highlight the need for further research that is both longitudinal and focuses on youth.¹³

The psychosocial stress caused by this pandemic has been detrimental to youth around the world, who have experienced adverse lifestyle changes.^{14,15} Confinement measures during lockdowns and the associated personal, educational and economic disruptions created pervasive social isolation, increased stress and decreased peer interactions, which may have triggered psychological pandemic. Pre-pandemic symptom severity predicted the observed mental health trajectories in the population cohort. Surprisingly, being relatively healthy predicted increases in depression and eating disorder symptoms and in body mass index. By contrast, those initially at higher risk for depression or eating disorders reported a lasting decrease.

Conclusions

Healthier young people may be at greater risk of developing depressive or eating disorder symptoms during the COVID-19 pandemic. Targeted mental health interventions considering prior diagnostic risk may be warranted to help young people cope with the challenges of psychosocial stress and reduce the associated healthcare burden.

Keywords

COVID-19; adolescent; depression; eating disorders; alcohol use disorder.

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distress and mental health difficulties in this age group. Indeed, meta-analyses of studies of children and adolescents indicate an increased prevalence of clinically elevated depression and anxiety symptoms compared with pre-pandemic estimates, especially in adolescent females.^{16,17} However, most studies investigated the effects of the pandemic on mental health changes only at the beginning of the pandemic. Although enormously instructive, these studies do not address the longer-term effects of the pandemic. Other limitations include the considerable heterogeneity of studies, which is largely due to differences in assessments and diagnostic criteria.¹⁸ The focus of most studies on anxiety and depression has also led to a call for more research to consider the effects of the pandemic on other youth mental health conditions that may have been negatively affected by the COVID-19 pandemic, in particular, eating disorders and addiction.¹⁹ More limited evidence available suggests that pre-pandemic disordered eating is a risk factor for poorer mental health during the pandemic.8,20 However, interpretations of these findings are limited as, again, assessment of mental health was restricted to the period of eased

restrictions following the first lockdown. As for addiction, a decline in substance use has been reported, especially among adolescents initially at higher risk for substance use disorder.^{21,22} It is clear from these limitations that longitudinal trajectory research with comprehensive mental health assessments, spanning the pre-pandemic period and across multiple lockdown and release phases, is needed to understand the long-term impact of the COVID-19 pandemic on youth mental health.¹³ Research comparing data from the general population and from patient groups is also needed. Crucially, identifying the most vulnerable and resilient groups will be important for the design and delivery of the most appropriate targeted interventions.

Aims

Our study addresses these needs by using data collected before and throughout the COVID-19 pandemic in two pre-existing youth cohorts: IMAGEN, a longitudinal population-based adolescent cohort; and ESTRA/STRATIFY, a clinical cohort with diagnoses of MDD, alcohol use disorders (AUD) and eating disorders. Our repeated assessments, based on the CoRonavIruS Health Impact Survey (CRISIS)²³ and standardised mental health questionnaires, aimed to (i) establish trajectories of behaviours and mental health symptoms throughout stages of the pandemic in these cohorts; (ii) compare these trajectories to identify the most vulnerable groups; and (iii) identify pre-pandemic predictors of these mental health trajectories.

Method

Study design

Participants were drawn from three existing cohorts located in the UK, France and Germany: IMAGEN, STRATIFY and ESTRA. IMAGEN was a longitudinal population cohort, whereas STRATIFY and ESTRA were case-control cohorts. To be eligible for inclusion, participants needed to respond to our invitation and provide informed consent through an online form sent via email. Data collection was conducted through online question-naires, with the initial round taking place during the first national lockdown in the UK and Europe (April–May 2020). Subsequent follow-up surveys were administered when the first lockdown was released (July 2020) and when the second lockdown was imposed (November 2020). The design and reporting of our study were in accordance with the STROBE (Strengthening the Reporting of Observational Studies in Epidemiology) guidelines.

Participants

Population cohort

These participants, with no known history of mental illness, were drawn from the IMAGEN study, a longitudinal cohort of over 2000 adolescents recruited at age 14 years from eight study sites in Europe, with follow-up assessments at ages 16, 19 and 23 years. For detailed study protocols, please refer to Schumann et al.²⁴ Our survey was sent to those who had completed the follow-up assessment at age 23 (N=1350). A total of 458 IMAGEN participants recruited from the UK, France and Germany (London, Nottingham, Paris, Mannheim and Berlin) who completed the COVID-19 survey at baseline were included in our analyses.

Clinical cohort

This cohort was derived from two studies, STRATIFY and ESTRA, of participants aged 18–30 years (N = 628). STRATIFY participants included in this study comprised participants recruited in the UK

and Germany (London, Southampton and Berlin) who met diagnostic criteria for MDD and AUD, as assessed by self-report via online computerised screening. Participants were included if they had scores \geq 15 (moderate to severe) on the Patient Health Questionnaire (PHQ-9)²⁵ and Alcohol Use Disorders Identification Test (AUDIT)²⁶ for MDD and AUD, respectively. ESTRA consisted of participants recruited in London and meeting the DSM-5²⁷ diagnostic criteria for anorexia nervosa or bulimia nervosa. All were female. Their eating disorder symptoms were assessed using the Eating Disorder Diagnostic Scale (DSM-5 version) over a screening phone call by study researchers.²⁸ A total of 211 STRATIFY/ESTRA participants (80 MDD, 51 AUD, 47 anorexia nervosa and 33 bulimia nervosa) who completed the COVID-19 survey at baseline were included in our analyses.

Ethics statement

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. All procedures involving human subjects/patients were approved by King's College London Research Ethics Committee (17/LO/0552) for IMAGEN, London Westminster Research Ethics Committee (PNM/10/11-126) for STRATIFY and North West–Greater Manchester South Research Ethics Committee (20/NW/0143) for ESTRA. All adult participants provided written/online signature informed consent to participate in this study.

Survey and assessments

The COVID-19 survey

We adapted the CoRonavIruS Health Impact Survey (CRISIS v0.1 http://www.crisissurvey.org)²³ to examine changes to individuals' mental health and behaviours induced by the pandemic. The survey encompassed data collection at various time points, specifically: pre-pandemic (3 months prior, pre-LD1), during the first lock-down (LD1), after the first lockdown (after-LD1) and during the second lockdown (LD2). This questionnaire assessed a range of data domains including COVID-19-related health status and life changes, daily behaviours and emotions, and worries due to the COVID-19 crisis (see Supplementary Information available at https://doi.org/10.1192/bjo.2023.601 for details).

Mental health assessments

The severity of mental disorder symptoms was assessed with validated questionnaires, including the PHQ-9 for depressive symptoms, the Eating Disorder Examination Questionnaire (EDE-Q)²⁹ for eating disorder symptoms and the AUDIT Consumption²⁶ for alcohol misuse (see Supplementary Methods for details). Questionnaires were administered at three time points: (a) at the previous recruitment wave, ~3 years prior to the pandemic (pre-PD), (b) during the first lockdown (LD1) and (c) during the second lockdown (LD2). The exception was the EDE-Q, which was administered at only two time points (i.e. LD1 and LD2) in the clinical cohort.

Pre-pandemic mental health

Pre-PD symptom severity scores were used to classify participants from the population cohort, based on the following criteria. For depression, PHQ-9 scores of 0–4, 5–9 and 10+ were used to indicate minimal, mild and moderate to severe depression, respectively.²⁵ For alcohol misuse, AUDIT scores of 0–7 and 8+ were used to indicate low and high risk, respectively.³⁰ For eating disorders, EDE-Q global scores <2.8 (for females) or 1.68 (for males) were used to

indicate low risk; higher scores were considered to indicate probable eating disorders.^{31,32} For body mass index (BMI), we used the following categories: underweight or normal weight, BMI < 25; overweight or obese: BMI > 25.

Statistical analyses

Data were analysed in SPSS version 27 using mixed-effects analysis of variance (ANOVA), with within-subject effect (time) adjusted by country and between-subjects effects (cohort and sex) adjusted by country and age. For each analysis, participants were included if they had no missing data for any variable needed. Separate analyses were conducted, as detailed in the Supplementary Methods, on the whole sample or on each cohort separately. Statistical significance was set at P < 0.05.

Trajectories of lifestyle changes, worries and mental health symptoms during the pandemic

Scores from the COVID survey and mental health questionnaires were analysed across time points. In addition to time effects, we investigated cohort and sex effects, along with interaction effects (i.e. time \times cohort, time \times sex) in the whole sample. Given the strong cohort effects, we also investigated these trajectories in the population and clinical cohorts separately.

Trajectories of mental health symptoms based on pre-pandemic symptom severity

These analyses were performed in the population cohort only. Subgroups based on the severity of pre-PD symptoms (see above) were included in mixed-effects ANOVAs. Three analyses were run to investigate interactions between time and pre-PD severity of mental health symptoms (i.e. depression, alcohol misuse or eating disorder) during the pandemic.

Results

Sample description and participants' characteristics

A flowchart outlining the recruitment and follow-up of participants for this study is provided in Fig. 1. In total, 669 individuals (31.5% clinical cohort; 69.4% females) completed the COVID survey at pre-LD1, 471 (29.9% clinical cohort) at LD1 and 429 (27.0% clinical cohort) at LD2 (Supplementary Table 1). As expected, immediately prior to the pandemic, symptoms of depression (F(1,615) = 156.26, P < 0.001, $\eta_p^2 = 0.203$) and alcohol misuse (F(1,630) = 30.11, P < 0.001, $\eta_p^2 = 0.046$) were higher in the clinical cohort. BMI was higher in the population sample (F(1,543) = 17.76, P < 0.001, $\eta_p^2 = 0.032$). Females reported higher levels of depressive (F(1,615) = 13.95, P < 0.001, $\eta_p^2 = 0.022$) and eating disorder symptoms (F(1,411) = 45.92, P < 0.001, $\eta_p^2 = 0.100$), whereas males reported higher levels of alcohol misuse (F(1,630) = 23.38, P < 0.001, $\eta_p^2 = 0.036$).

Behavioural, emotional and mental health trajectories during the pandemic

We compared behavioural, emotional and mental health trajectories during the pandemic in our two cohorts using mixed-effects ANOVA. Significant main effects of time on behaviours (i.e. positive lifestyle changes, frequency of media use, average daily food consumption and frequency of substance use; all P < 0.001; Fig. 2(a–e) and Supplementary Table 2) were observed when analysing both samples together, but there were no significant time × cohort interactions. Similarly, there were significant main effects of time on emotional health, as assessed by the 'emotions and worries' and 'worries about COVID' sections of the survey (all P < 0.001; Fig. 3(a–e) and Supplementary Table 2) but no significant time × cohort interactions (all detailed in the Supplementary Material).

Comparisons of mental health symptoms (i.e. depression, alcohol drinking and eating disorders) and BMI just prior to the pandemic and during both lockdowns revealed the differential impact of the COVID-19 crisis on the cohorts (Fig. 4(a-d) and Supplementary Table 3).

There was no significant main effect of time on depressive symptoms (F(2,836) = 0.43, P = 0.65, $\eta_p^2 = 0.001$) in the whole sample. As expected, there were sex (F(1,417) = 11.84, P < 0.001, $\eta_p^2 = 0.028$) and large cohort effects (F(1, 1417) = 213.05, P < 0.001, $\eta_p^2 = 0.338$), depressive symptoms being higher in females and in the clinical cohort. A significant time × cohort interaction was also found (F(2,836) = 7.41, P < 0.001, $\eta_p^2 = 0.017$), indicating that the trajectories of depressive symptoms significantly differed between the population and the clinical samples (Fig. 4(a)). Analyses of these trajectories in the two cohorts separately revealed a significant main effect of time only in the population cohort $(F(2,606) = 16.98, P < 0.001, \eta_p^2 = 0.053)$. Depressive symptoms increased by 33.9% (95% CI, 31.78-36.57) during the lockdowns, with severity increasing to mild depression compared with minimal depression prior to the pandemic. In the clinical cohort, depressive symptoms remained high and constant across time $(F(2,220) = 0.10, P = 0.91, \eta_p^2 = 0.001).$

A significant main effect of time in harmful alcohol drinking was found (F(2,838) = 14.06, P < 0.001, $\eta_p^2 = 0.032$), with symptoms decreasing during the pandemic to reach their lowest levels (i.e. a 15.2% decrease) during the second lockdown (pre-PD > LD1 > LD2; P < 0.05). Males drank more than females (F(1,418) = 41.90, P < 0.001, $\eta_p^2 = 0.091$). There were no significant cohort or time × cohort interactions (Fig. 4(b)). Nonetheless, a time × diagnosis interaction in the clinical cohort (F(6,220) = 4.25, P < 0.001, $\eta_p^2 = 0.104$) revealed that the significant the decline in harmful drinking in the clinical cohort (i.e. a 23.04% decrease) was driven by participants with AUD.

For eating disorder symptoms, as the EDE-Q was only administered at two time points in the clinical cohort, we analysed the two cohorts separately. In the population cohort, there was a significant main effect of time on eating disorder behaviours and attitudes, as assessed by the EDE-Q global score (F(2,550) = 4.31, P = 0.01, $\eta_p^2 = 0.015$). Eating disorder symptoms increased by 15.6% (95%) CI, 15.39-15.68) during the first lockdown, returning to prepandemic levels during the second lockdown (Fig. 4(b)). As expected, eating disorder symptoms were significantly higher in females than males (F(1, 274) = 33.26, $P = \langle 0.001, \eta_p^2 = 0.108 \rangle$, but there were no significant time \times sex interactions (F(2, 550) = 1.14, P = 0.32, $\eta_p^2 = 0.004$). In contrast to our findings in the population cohort, eating disorder symptoms did not significantly differ between the two lockdowns in the clinical cohort (F(1,110) = 0.09), P = 0.77, $\eta_p^2 = 0.001$). Limiting analyses to the eating disorder subgroups also revealed no significant time effects (F(1,30) = 2.35,P = 0.14, $\eta_p^2 = 0.073$ and F(1,18) = 0.03, P = 0.87, $\eta_p^2 = 0.002$), for anorexia nervosa and bulimia nervosa, respectively.

Analyses of BMI trajectories revealed a significant main effect of time in the whole sample (F(2, 638) = 6.85, P < 0.001, $\eta_p^2 = 0.032$), with higher BMIs during the pandemic (pre-PD < LD1 and LD2, P < 0.01) (Fig. 4(d)). A significant time × sex interaction (F(2,638) = 3.81, P < 0.05, $\eta_p^2 = 0.012$) indicated that BMI significantly increased in females but not in males. No time × cohort interaction was found, but analyses of the two cohorts separately indicated that these findings were driven by the population cohort (F(2,428) = 9.61, P < 0.01, $\eta_p^2 = 0.043$). Although no main effect of time was found in the clinical cohort, analyses within each diagnostic group



symptom severity on mental health trajectories during the pandemic. For each analysis, participants were excluded if they had missing data for any required variable. ADHD, attention-deficit hyperactivity disorder; LD1, first lockdown; LD2, second lockdown; RecAN, recovered from anorexia nervosa; RecBN, recovered from bulimia nervosa.

revealed a time × sex interaction (F(2,50) = 5.48, P < 0.01, $\eta_p^2 = 0.180$) in the AUD group, with significant BMI increases observed only in females (pre-PD < LD2, P < 0.05).

Re-running the analyses described above while controlling for other potential confounders generated largely similar results (Supplementary Table 3).

Effects of pre-pandemic symptom severity on mental health trajectories during the pandemic

The following analyses were performed to identify participants from the population cohort most vulnerable to COVID-induced mental illness. We categorised participants from this cohort based on their pre-pandemic symptom severity with respect to depression, alcohol misuse, eating disorders and BMI and re-ran analyses with these categories as predictors (Fig. 5(a-e) and Supplementary Table 4).

Effects of pre-pandemic depression symptom severity on depressive symptom trajectories

There were significant interactions between time and pre-pandemic symptom severity for pandemic-related depressive symptoms (F (4,602) = 21.35, P < 0.001, $\eta_p^2 = 0.124$). *Post hoc* analyses revealed

notable group differences in trajectories (Fig. 5(a) and Supplementary Table 4). Participants with minimal pre-pandemic depression symptoms reported significant changes over time (F (2,300) = 41.73, P = <0.001, $\eta_p^2 = 0.218$), with symptoms increasing during the first lockdown and remaining higher afterwards (pre-PD < LD1 or LD2, P < 0.001). By contrast, participants with moderate to severe depression reported the opposite trend (F(2,300) = 17.54, P = <0.001, $\eta_p^2 = 0.105$), with symptoms being lower during the first and second lockdowns (pre-PD > LD1 or LD2, P < 0.001). Participants with mild depression did not report significant symptom changes with time (F(2,300) = 1.59, P = 0.21, $\eta_p^2 = 0.010$).

Effects of pre-pandemic risk for alcohol misuse on trajectories of alcohol misuse

There were significant group differences in trajectories of harmful drinking during the pandemic (F(2,606) = 16.26; P = <0.001, $\eta_P^2 = 0.051$; Fig. 5(b) and Supplementary Table 4). Participants initially more at risk of harmful drinking (i.e. prior to the pandemic) reported a significant decrease in alcohol misuse at all time points during the pandemic (pre-PD > LD1 > LD2, all, P < 0.001). A decrease was also observed for participants at low risk, and this became significant during the second lockdown (pre-PD > LD2, P < 0.001; LD1 > LD2, P < 0.05).



Fig. 2 Behavioural trajectories during the pandemic, including (a) positive life changes; (b) frequency of exercising; (c) frequency of media use; (d) daily food consumption; and (e) frequency of substance use, in the whole sample and stratified by cohort. Data are expressed as mean and standard error. Time effects from mixed-effects ANOVA in the whole sample were estimated by comparing data collected before the first lockdown with data collected at other time points (*P < 0.05, **P < 0.01, ***P < 0.001) and by comparing data collected during the first lockdown with data collected afterwards ($^{+}P < 0.05$, $^{++}P < 0.01$, $^{+++}P < 0.001$).

Effects of pre-pandemic risk for eating disorder on eating disorder symptoms and BMI trajectories

symptoms during the pandemic were observed (F(2, 548) = 18.07; P =

disorder symptoms specifically during the first lockdown, with symptoms decreasing during the second lockdown (pre-PD < LD1, P =<0.001; LD1 > LD2, P < 0.01). Conversely, for participants initially Similarly, significant group differences in trajectories of eating disorder scoring higher for eating disorder symptoms (i.e. those with probable eating disorder), symptoms significantly decreased during the first <0.001, $\eta_p^2 = 0.062$; Fig. 5(c) and Supplementary Table 4). Participants lockdown (P < 0.001), remaining lower during the second lockdown. initially at low risk for eating disorders reported a increase in eating



Fig. 3 Emotional trajectories during the pandemic, including (a) emotions and worries; (b) worries about oneself being infected; (c) worries about friends or family being infected; (d) worries about own physical health; and (e) worries about own mental health, in the whole sample and stratified by cohort. Data are expressed as mean and standard error. Time effects from mixed-effects ANOVA in the whole sample were estimated by comparing data collected before the first lockdown with data collected at other time points (*P < 0.05, **P < 0.01, ***P < 0.001) and by comparing data collected during the first lockdown with data collected afterwards ($^{+}P < 0.05$, $^{++}P < 0.01$, $^{+++}P < 0.001$).



Fig. 4 Mental health trajectories during the pandemic. Trajectories of (a) depressive symptoms; (b) harmful alcohol drinking; (c) eating disorder symptoms (d) and body mass index are indicated for the whole sample and for each cohort separately. Data are expressed as mean and standard error. Time effects from mixed-effects ANOVA in the whole sample were estimated by comparing data collected before the pandemic with data collected at other time points (*P < 0.05, **P < 0.01, ***P < 0.001), and by comparing data collected during the first lockdown with data collected afterwards (*P < 0.05, **P < 0.001).

Unsurprisingly, there were significant group differences in BMI (F(1,195) = 16.03; P < 0.001, $\eta_p^2 = 0.076$), with participants with probable eating disorder having BMIs in the overweight range and those at low risk having BMIs in the normal range (Fig. 5(d)). No significant group differences in BMI trajectories during the pandemic were observed (F(2,392) = 0.43; P = 0.65, $\eta_p^2 = 0.002$); a nominally significant increase in BMI was observed in the participants at low risk for eating disorder (pre-PD < LD2, P < 0.05) but not in those with higher eating disorder risk (Fig. 5(d)).

Effects of pre-pandemic BMI on BMI and eating disorder symptoms trajectories

Although no significant group differences on BMI trajectories were observed when comparing participants who were initially underweight/normal weight (BMI < 25) and overweight/obese (BMI > 25) (F(2,426) = 2.09; P = 0.13, $\eta_p^2 = 0.010$), significant increases in BMI were observed in the underweight/normal weight group (pre-PD < LD1, P = 0.005; pre-PD < LD2, P < 0.001) but not in the overweight/obese group, for which BMI remained constant during the pandemic (Fig. 5(e) and Supplementary Table 4). Consistent with the analyses above, the increase in BMI in the underweight/normal weight group was paralleled by a significant increase in eating disorder symptoms, specifically during the first lockdown (BMI < 25; F(2,270) = 5.74, P = 0.004, $\eta_p^2 = 0.041$; pre-PD < LD1, P = 0.003; LD1 > LD2, P = 0.045).

When re-running analyses controlling for other potential confounders, minor differences emerged in *post hoc* tests, probably owing to increased degrees of freedom and reduced sample size after adding numerous covariates. However, the overall pattern remained – participants with high levels of pre-pandemic symptoms showed improvement during lockdowns, whereas those with minimal pre-pandemic depression symptoms reported significant increases over time.

Discussion

This comparative study following population and clinical cohorts during the pandemic revealed the differing impact of the pandemic in youth with and without pre-existing mental illness. Whereas symptoms of depression and eating disorders increased during the pandemic in young people from the population, these symptoms remained high and stable in the clinical cohort. Pre-pandemic symptom severity predicted mental health trajectories in the population cohort. Participants initially at higher risk for depression, alcohol misuse or eating disorders reported a lasting decrease in their symptoms over the course of the pandemic. By contrast, being relatively healthy (i.e. having the lowest scores for depression or eating disorder) was a significant risk for deterioration in mental health during the pandemic; this was associated with relative increases in depressive symptoms throughout the pandemic and in eating disorder symptoms during the first lockdown. Being



depression (minimal, mild and moderate to severe) on trajectories of depressive symptoms; (b) effects of pre-pandemic risk for alcohol misuse (low or high risk) on trajectories of harmful alcohol drinking; (c) effects of pre-pandemic risk for eating disorders (low risk or probable eating disorder) on trajectories of eating disorder symptoms; (d) effects of pre-pandemic risk for eating disorders on body mass index (BMI) trajectories, and (e) effects of pre-pandemic BMI (low or normal and overweight or obese) on BMI trajectories. Data are expressed as mean and standard error. Mixed-effects ANOVA revealed significant time × group (i.e. pre-pandemic risk levels) interactions in all comparisons. Time effects in each group were estimated by comparing data collected before the pandemic with data collected at other time points (P < 0.05, *P < 0.01, **P < 0.001) and by comparing data collected during the first lockdown with data collected afterwards (P < 0.05, *P < 0.01, **P < 0.001).

non-overweight or non-obese predicted the observed rise in eating disorder symptoms and was associated with weight gain (i.e. BMI increase).

Our findings corroborate previous research showing an increase in depression symptoms in all age groups^{5,15,16,33} from the population during the pandemic, but particularly in young and more physically active individuals.¹⁵ This observation may reflect greater changes in lifestyle habits in this group or a reduced tolerance of uncertainty. Our findings also highlight the contrasting effects of the pandemic on other mental health outcomes in young people: a long-term negative impact on depressive symptoms lasting until the second lockdown, in contrast to the transient increase in eating disorder symptoms and continuous decrease in alcohol misuse.

Our findings also shed light on the contradictory debate concerning pre-existing mental illnesses.^{7-12,34} Contrary to previous reports of worsening symptoms during the pandemic in patients with a history of mental illness⁷ or pre-existing disordered eating,⁸ our findings indicated that although symptoms remained higher in the clinical sample, they did not worsen because of the pandemic. These discrepancies may be due to a lack of diagnostic measurement of mental illness and lack of repeated assessments to measure symptom changes during the pandemic in the relevant studies. Our observations of differences in mental health trajectories between young people from the general population and those with a clinical diagnosis suggest that pre-pandemic symptoms may have been a protective factor, and that the general population was more likely to be affected by the lockdowns than patients, which our analyses confirmed. The clinical cohort seemed to be resilient in the face of the pandemic, confirming previous reports for depression from the early stages of the pandemic^{10,11,35} and further indicating that this effect persisted as the pandemic progressed. By contrast, and in agreement with previous assessments of depression in adults9 and adolescents,12 young people without depressive or

eating disorder symptoms showed an increase in these symptoms during the pandemic, whereas those with the highest pre-pandemic risk experienced a decrease. However, it should be noted that symptoms in the higher-risk groups remained much higher than those of individuals without prior symptoms, and that patients are more vulnerable to some stressful situations due to the pandemic.³⁴

In contrast to our findings for depression and eating disorders, we observed a decline in alcohol and substance misuse during the pandemic, consistent with previous evidence.^{22,36} This decline during lockdown periods was similar in participants with and without mental health diagnoses. Among the general population, this decline could be attributed to both those at high risk and those at low risk for alcohol misuse and may reflect closures of shops, bars and pubs during lockdown.

Participants from the general population and patients differed in the intensity of their behavioural or emotional responses to the pandemic but not in their trajectories. That young people are not equally at risk from the psychosocial stress brought about by COVID-19 was to be expected; however, counterintuitively, our findings indicate that healthier individuals tended to be the most vulnerable to the negative effects of the pandemic on mental health, not those with a higher burden. Possible explanations for this are that heightened fears and worries during periods of confinement, as highlighted in this study, and increased social isolation may have contributed to deterioration of mental health in healthier individuals. By contrast, those with depression and eating disorders might have felt relief owing to reduced exposure to psychosocial stressors (e.g. social interactions). They may also have felt less isolated given the global increase in fears and worries. As for alcohol and substance use, as noted above, the general reduction may reflect restriction policies such as closures of shops, bars and pubs, which would have limited access to those substances, as evidenced by a return to pre-pandemic levels after confinement measures were lifted. In addition, the more time young people spent

at home with their families, the less likely they were to gain access to these substances.

Strengths and limitations

Strengths of our study include the use of longitudinal data collected over a period of up to 3 years prior to the pandemic and further assessments covering the two lockdowns, which allowed for a more comprehensive understanding of the impact of the pandemic. A clear strength is also the combination of data from the youth population as well as from patients with pre-existing mental illness, with both groups assessed under the same study protocol. This enabled the investigation of vulnerability and resilience and improved our understanding of how distinct groups of people may respond to challenging circumstances. However, some limitations should be acknowledged. First, our study had a relatively low response rate and high attrition during the data collection phase. It did not include underrepresented groups, such as participants from ethnic minorities that may have been disproportionately affected by the pandemic. Moreover, our clinical sample was relatively small, with the majority of participants being females. All of this may limit the generalisability of our findings. In addition, although our study used validated instruments (PHQ-9, AUDIT and EDE-Q) to measure psychiatric symptoms, these are not diagnostic tools but only measure a greater risk of the presence of clinical illness. Finally, psychiatric assessments were only conducted during periods of confinement, which precluded investigation of mental health changes once restrictions were lifted.

Clinical implications

In summary, our study revealed opposite effects of the pandemic on mental health in youth with and without mental illness. Improvements in depression, alcohol misuse or eating disorder symptoms were observed over the course of the pandemic for participants with a higher pre-pandemic risk for these disorders, suggesting that the pandemic and lockdown measures decreased the mental health burden specifically in this population group. By contrast, the increases in depressive and eating disorder symptoms in those with low prior risk suggest the detrimental effects of such measures on healthier youth. If confirmed by future studies in a more representative sample, our findings could support personalised mental health interventions to help young people to cope better with the challenges of psychosocial stress and reduce the associated healthcare burden.

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Supplementary material

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Data availability

The data that support the findings of this study are available from the corresponding author (S.D.) on reasonable request.

Author contributions

S.D. conceived and designed the study. L.R., M.B., C.G., J.W., R.A., K.A., Y.Z., S.K., E.A., T.B., A.L.W.B., M.J.B., R.B., H.F., J.H.F., H.G., A.G., A.H., S.H., M.-L.P.M., S.M., F.N., B.M.N., L.P., J.S., M.N.S., R.W., A.S., H.W., J.-L.M., G.S., U.S. and S.D. collected the data. D.P.O. managed the data. L.Q. analysed the data and drafted the initial output. L.Q., Z.Z. and S.D. contributed to the interpretation of findings. S.D. will serve as a guarantor for the contents of the paper. All authors have read and approved the final version of the manuscript.

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References

- 1 COVID-19 Mental Disorders Collaborators. Global prevalence and burden of depressive and anxiety disorders in 204 countries and territories in 2020 due to the COVID-19 pandemic. *Lancet* 2021; **398**(10312): 1700–12.
- 2 Aknin LB, Andretti B, Goldszmidt R, Helliwell JF, Petherick A, De Neve JE, et al. Policy stringency and mental health during the COVID-19 pandemic: a longitudinal analysis of data from 15 countries. *Lancet Public Health* 2022; 7(5): e417–26.
- 3 Solomon-Moore E, Lambert J, Grey E, Gillison F, Townsend N, Busam B, et al. Life in lockdown: a longitudinal study investigating the impact of the UK COVID-19 lockdown measures on lifestyle behaviours and mental health. BMC Public Health 2022; 22(1): 1495.
- 4 Glowacz F, Schmits E. Psychological distress during the COVID-19 lockdown: the young adults most at risk. *Psychiatry Res* 2020; **293**: 113486.
- 5 Fancourt D, Steptoe A, Bu F. Trajectories of anxiety and depressive symptoms during enforced isolation due to COVID-19 in England: a longitudinal observational study. *Lancet Psychiatry* 2021; 8(2): 141–9.
- 6 Sawyer SM, Azzopardi PS, Wickremarathne D, Patton GC. The age of adolescence. Lancet Child Adolesc Health 2018; 2(3): 223–8.
- 7 Lewis KJS, Lewis C, Roberts A, Richards NA, Evison C, Pearce HA, et al. The effect of the COVID-19 pandemic on mental health in individuals with pre-existing mental illness. *BJPsych Open* 2022; 8(2): e59.
- 8 Warne N, Heron J, Mars B, Kwong ASF, Solmi F, Pearson R, et al. Disordered eating and self-harm as risk factors for poorer mental health during the COVID-19 pandemic: a UK-based birth cohort study. J Eat Disord 2021; 9(1): 155.
- **9** Pan KY, Kok AAL, Eikelenboom M, Horsfall M, Jorg F, Luteijn RA, et al. The mental health impact of the COVID-19 pandemic on people with and without depressive, anxiety, or obsessive-compulsive disorders: a longitudinal study of three Dutch case-control cohorts. *Lancet Psychiatry* 2021; **8**(2): 121–9.
- 10 Pinkham AE, Ackerman RA, Depp CA, Harvey PD, Moore RC. A longitudinal investigation of the effects of the COVID-19 pandemic on the mental health of individuals with pre-existing severe mental illnesses. *Psychiatry Res* 2020; 294: 113493.

- 11 Hamm ME, Brown PJ, Karp JF, Lenard E, Cameron F, Dawdani A, et al. Experiences of American older adults with pre-existing depression during the beginnings of the COVID-19 pandemic: a multicity, mixed-methods study. Am J Geriatr Psychiatry 2020; 28(9): 924–32.
- 12 Sadeghi N, Fors PQ, Eisner L, Taigman J, Qi K, Gorham LS, et al. Mood and behaviors of adolescents with depression in a longitudinal study before and during the COVID-19 pandemic. J Am Acad Child Adolesc Psychiatry 2022; 61(11): 1341–50.
- 13 Wade M, Prime H, Browne DT. Why we need longitudinal mental health research with children and youth during (and after) the COVID-19 pandemic. *Psychiatry Res* 2020; 290: 113143.
- 14 Lee J. Mental health effects of school closures during COVID-19. Lancet Child Adolesc Health 2020; 4(6): 421.
- 15 Amerio A, Lugo A, Stival C, Fanucchi T, Gorini G, Pacifici R, et al. COVID-19 lockdown impact on mental health in a large representative sample of Italian adults. J Affect Disord 2021; 292: 398–404.
- 16 Racine N, McArthur BA, Cooke JE, Eirich R, Zhu J, Madigan S. Global prevalence of depressive and anxiety symptoms in children and adolescents during COVID-19: a meta-analysis. JAMA Pediatr 2021; 175(11): 1142–50.
- 17 Ma L, Mazidi M, Li K, Li Y, Chen S, Kirwan R, et al. Prevalence of mental health problems among children and adolescents during the COVID-19 pandemic: a systematic review and meta-analysis. J Affect Disord 2021; 293: 78–89.
- 18 Diaz Gonzalez-Colmenero F, Millan-Alanis JM, Barrera FJ, Saucedo-Uribe E. Letter to the editor: 'Prevalence of mental health problems among children and adolescents during the COVID-19 pandemic: a systematic review and meta-analysis'. J Affect Disord 2021; 294: 479–80.
- 19 Dey T, Mansell ZJ, Ranu J. Effect of the COVID-19 pandemic on adolescents with eating disorders. JAMA Pediatr 2022; 176(2): 205–6.
- 20 Hyam L, Richards KL, Allen KL, Schmidt U. The impact of the COVID-19 pandemic on referral numbers, diagnostic mix, and symptom severity in eating disorder early intervention services in England. Int J Eat Disord 2023; 56(1): 269–75.
- 21 Sheikhan NY, Hawke LD, Ma C, Courtney D, Szatmari P, Cleverley K, et al. A longitudinal cohort study of youth mental health and substance use before and during the COVID-19 pandemic in Ontario, Canada: an exploratory analysis. *Can J Psychiatry* 2022; 67(11): 841–53.
- 22 Deeken F, Reichert M, Zech H, Wenzel J, Wedemeyer F, Aguilera A, et al. Patterns of alcohol consumption among individuals with alcohol use disorder during the COVID-19 pandemic and lockdowns in Germany. JAMA Netw Open 2022; 5(8): e2224641.
- 23 Nikolaidis A, Paksarian D, Alexander L, Derosa J, Dunn J, Nielson DM, et al. The Coronavirus Health and Impact Survey (CRISIS) reveals reproducible correlates of pandemic-related mood states across the Atlantic. *Sci Rep* 2021; **11**(1): 8139.
- 24 Schumann G, Loth E, Banaschewski T, Barbot A, Barker G, Büchel C, et al. The IMAGEN study: reinforcement-related behaviour in normal brain function and psychopathology. *Mol Psychiatry* 2010; **15**(12): 1128–39.
- 25 Kroenke K, Spitzer RL, Williams JB. The PHQ-9: validity of a brief depression severity measure. J Gen Intern Med 2001; 16(9): 606–13.
- 26 Bush K, Kivlahan DR, McDonell MB, Fihn SD, Bradley KA. The AUDIT alcohol consumption questions (AUDIT-C): an effective brief screening test for problem drinking. Ambulatory Care Quality Improvement Project (ACQUIP). Alcohol Use Disorders Identification Test. Arch Intern Med 1998; 158(16): 1789–95.
- 27 Asken MJ, Grossman D, Christensen LW, American Psychiatric Association. *Diagnostic and Statistical Manual of Mental Disorders*. American Psychiatric Publishing, 2013.
- 28 Stice E, Telch CF, Rizvi SL. Development and validation of the Eating Disorder Diagnostic Scale: a brief self-report measure of anorexia, bulimia, and bingeeating disorder. *Psychol Assess* 2000; 12(2): 123–31.
- 29 Fairburn CG, Beglin SJ. Assessment of eating disorders: interview or self-report questionnaire? Int J Eat Disord 2006; 16(4): 363–70.
- 30 Piccinelli M, Tessari E, Bortolomasi M, Piasere O, Semenzin M, Garzotto N, et al. Efficacy of the Alcohol Use Disorders Identification Test as a screening tool for hazardous alcohol intake and related disorders in primary care: a validity study. Br Med J 1997; 314(7078): 420.
- **31** Schaefer LM, Smith KE, Leonard R, Wetterneck C, Smith B, Farrell N, et al. Identifying a male clinical cutoff on the Eating Disorder Examination-Questionnaire (EDE-Q). *Int J Eat Disord* 2018; **51**(12): 1357–60.
- 32 Mond JM, Myers TC, Crosby RD, Hay PJ, Rodgers B, Morgan JF, et al. Screening for eating disorders in primary care: eDE-Q versus SCOFF. *Behav Res Ther* 2008; 46(5): 612–22.
- 33 Devoe JD, Han A, Anderson A, Katzman DK, Patten SB, Soumbasis A, et al. The impact of the COVID-19 pandemic on eating disorders: a systematic review. Int J Eat Disord 2023; 56(1): 5–25.

- 34 Ambrosetti J, Macheret L, Folliet A, Wullschleger A, Amerio A, Aguglia A, et al. Impact of the COVID-19 pandemic on psychiatric admissions to a large Swiss emergency department: an observational study. Int J Environ Res Public Health 2021; 18(3):1174.
- 35 Steff A, Godinot ML, Gourlan C, Robinson L, Vidya N, Winterer J, et al. P.0509 Emotion de-regulation in a cohort of young Europeans during the spring 2020 lockdown. Eur Neuropsychopharmacol 2021; 53: S374–5.
- 36 Layman HM, Thorisdottir IE, Halldorsdottir T, Sigfusdottir ID, Allegrante JP, Kristjansson AL. Substance use among youth during the COVID-19 pandemic: a systematic review. *Curr Psychiatry Rep* 2022; 24(6): 307–24.

