Associations of Despair With Suicidality and Substance Misuse Among Young Adults

William E. Copeland, PhD; Lauren Gaydosh, PhD; Sherika N. Hill, PhD; Jennifer Godwin, PhD; Kathleen Mullan Harris, PhD; E. Jane Costello, PhD; Lilly Shanahan, PhD

Abstract

IMPORTANCE Deaths of despair is a term that has recently been used to describe the increases in premature mortality from suicides, drug overdoses (particularly from opiates), and alcohol-related liver disease among US adults. Despite the use of the term despair, its role in these causes of premature death has not been empirically tested.

OBJECTIVE To test whether despair among young adults is associated with suicidal thoughts and behavior, alcohol misuse, and drug misuse.

DESIGN, SETTING, AND PARTICIPANTS The Great Smoky Mountains Study is a Southeastern, mixed urban-rural population-based cohort study conducted from November 10, 1992, to September 22, 2015. A total of 1420 participants originally 9, 11, and 13 years of age were followed up 11 times to 30 years of age (11,230 person-observations). A total of 1154 of 1400 living participants (82.4%) were assessed at 30 years of age. Statistical analysis was performed from May 7, 2019, to April 10, 2020.

EXPOSURES Participants were assessed with structured interviews for indicators of despair (eg, hopelessness, helplessness, low self-worth, and feeling unloved). Despair was assessed with items from structured interviews: the Child and Adolescent Psychiatric Assessment and the Young Adult Psychiatric Assessment.

MAIN OUTCOMES AND MEASURES Structured interviews were used to assess suicidal thoughts and behavior, substance use, and Diagnostic and Statistical Manual of Mental Disorders (Fifth Edition) alcohol use disorder and drug use disorder (including opioids) in young adulthood (2424 observations of 1266 individuals between 25 and 30 years of age).

RESULTS This study included 1420 individuals (790 male individuals). During young adulthood (25 and 30 years of age), the 3-month weighted prevalence of any despair was 19.5% (476 of 2424 observations) with 7.6% of participants (201 of 2424 observations) reporting 2 or more despair items. In longitudinal, lagged models, despair scores (range, 0-3) were associated with more suicidal thoughts and behaviors (odds ratio [OR], 1.5; 95% CI, 1.1-2.0), illicit drug use (OR, 1.7; 95% CI, 1.2-2.5), and opioid use (OR, 1.9; 95% CI, 1.1-3.3) but not alcohol use disorder (OR, 0.8; 95% CI, 0.6-1.2). These associations persisted after accounting for sociodemographic factors (eg, poverty and educational level), lagged outcome status, and lagged depression status. The associations between despair and study outcomes were stronger in models accounting for long-term measures of despair extending back to childhood. There was no consistent pattern of moderation by sociodemographic factors.

CONCLUSIONS AND RELEVANCE This study’s findings suggest an empirical basis for longitudinal associations between despair and several, but not all, precursors of “deaths of despair” in rural

Key Points

Question Is despair associated with drug or alcohol misuse or suicidal thoughts and behaviors among young adults?

Findings In this population-based cohort study in rural Appalachia, despair was longitudinally associated with higher rates of suicidal thoughts and behavior, illicit drug use, and opioid use, even after adjusting for sociodemographic factors, prior outcome status, and prior depressive disorder status; despair was not associated with alcohol use disorder. There was no consistent pattern of moderation by race/ethnicity, poverty status, sex, or educational level.

Meaning Despair early in life is longitudinally associated with several (but not all) putative despair-related diseases.

Open Access. This is an open access article distributed under the terms of the CC-BY License.

JAMA Network Open. 2020;3(6):e208627. doi:10.1001/jamanetworkopen.2020.8627 June 23, 2020 1/12
Abstract (continued)

Appalachia. Individual despair should be studied as a potential factor associated with morbidity and impairment in young adulthood.

Introduction

In recent decades, income stagnation and economic declines have coincided with premature deaths from suicide, alcoholism, and drug (especially opioid) overdoses, particularly among white non-Hispanic adults with low education in rural areas. Recent work reports that premature mortality from these causes occurs not only among middle-aged white adults but also among young adults and additional racial/ethnic groups. This reversal of US life-expectancy improvements was coined deaths of despair and has raised public and scientific interest. Studies on deaths of despair have not measured the central empirical claim that despair is a shared risk pathway to suicides and alcohol-and drug-related deaths or their precursors: suicidal behaviors and thoughts, alcohol misuse and dependence, and illicit drug use (ie, the “diseases of despair”). To our knowledge, to date, despair has not been studied as an independent construct in and of itself.

Despair is defined as “a state of mind in which there is an entire want of hope.” From a psychological perspective, this definition focuses on a cognitive state that includes defeat, guilt, worthlessness, learned helplessness, pessimism, and limited positive expectations for the future. Some of these cognitions overlap with specific criteria for Diagnostic and Statistical Manual of Mental Disorders Fifth Edition (DSM-5)–based depressive disorders (eg, feeling worthless and major depressive disorder), whereas others do not (eg, helplessness and loneliness). The present study tests the hypothesis that despair is a shared risk pathway to diseases of despair. We use data from a prospective, longitudinal, community-representative study of more than 25 years conducted in a rural area of the Southeastern United States. The sample was recruited from parts of the Appalachian region, which has had particularly high levels of premature mortality and deaths of despair, including from opiate overdoses. This analysis aims to test whether despair early in life might play a role in the premature mortality crisis.

We first examined the prevalence of despair from childhood to midlife. Next, we tested whether despair is associated with suicidal thoughts and behaviors, alcohol misuse, and drug misuse in young adulthood (between 25 and 30 years of age). Third, we used longitudinal lagged analyses to test directionality between despair and its hypothesized diseases over time. That is, does despair precede the diseases of despair, do the diseases precede despair, or both? We hypothesize that despair will be associated with higher levels of diseases of despair but do not have specific hypotheses for which diseases would have strongest associations. To truly illuminate the role of despair, we tested whether associations of despair with outcomes are independent of DSM-based depressive disorders with which despair is commonly associated. Finally, we tested whether associations are particularly strong in certain demographic groups, including those with low educational levels and those from a white non-Hispanic background.

Methods

Participants

The Great Smoky Mountains Study is a representative cohort study of children in 11 mostly rural counties of North Carolina. Three cohorts of children, aged 9, 11, and 13 years, were recruited from a pool of approximately 12,000 children using a 2-stage sampling design, resulting in 1420 participants (630 girls). First, potential participants were randomly selected from the population using a household equal probability design. Next, participants were screened for risk of
psychopathological conditions; participants whose screening results indicated a high risk were oversampled in addition to a random sample of the rest of the participants. In addition, Native American participants were oversampled to constitute 25% of the sample. In all statistical analyses, sampling weights are applied to adjust for the differential probability of selection and to allow results to generalize to the broader population of children from which the sample was drawn (eFigure in the Supplement). Before all interviews, parents and children signed informed consent or assent forms. The study protocol and consent forms for each assessment were approved by the Duke University Medical Center institutional review board. Participants received payment for their time ($100 for the most recent wave). This study followed the Strengthening the Reporting of Observational Studies in Epidemiology (STROBE) reporting guidelines.

Annual assessments of the 1420 children were completed during the period from 9 to 16 years of age (6674 total observations; 1993-2000) and then again at 19, 21, 25, and 30 years of age (4556 observations of 1336 participants; 1999-2015), for a total of 11 230 total assessments. Interviews were completed separately by a parent figure and the participant until 16 years of age and by the participant only thereafter.

Measures
Despair was assessed with items from structured interviews: the Child and Adolescent Psychiatric Assessment and the Young Adult Psychiatric Assessment (YAPA). These structured interviews assess the psychiatric symptoms needed to make diagnoses but also a broad range of associated constructs. To derive a scale of despair, all interview items were reviewed for consistency with how despair has been defined in recent publications on deaths of despair. Seven items of the Child and Adolescent Psychiatric Assessment and YAPA interviews capture the cognitive features of despair: loneliness, hopelessness, feeling unloved, helplessness, low self-esteem, frequent worries, and feeling sorry for oneself. eTable 1 in the Supplement includes a full list of despair-related items, their operational definitions, and their overlap with DSM constructs. Two items used in the despair scale are criteria for DSM-defined persistent depressive disorder or dysthymia (ie, a pattern of dysphoria lasting more than 2 years), 1 item is a criterion for major depressive disorder, 1 item is a criterion for generalized anxiety disorder, and 3 items are not criteria for any psychiatric disorder. The 7 dichotomous despair indicators were summed into a despair score. The internal consistency of this scale was acceptable (Cronbach α = .73). This scale was winsorized at 3 or more indicators of despair to aid in the presentation of results. Despair may also occur in the emotional (eg, sadness and anhedonia), behavioral (eg, reckless and high-risk behaviors), and biological (eg, signs of biological depletions such as high allostatic load) domains, but these domains were not the focus of the present study.

Diseases Associated With Dispair
The 3 diseases of despair assessed in this study included suicidal thoughts and behaviors, alcoholism, and drug misuse, including opioid use. All outcomes were assessed using the YAPA, which focuses on the 3 months immediately preceding the interview to minimize forgetting and recall bias. Additional sociodemographic variables were collected on sex, race/ethnicity, educational level, and falling below the federal poverty thresholds based on income and family size.

Suicidal Thoughts and Behaviors
eTable 2 in the Supplement provides definitions for passive suicidal ideation, active suicidal ideation, suicide plans, or suicide attempts. The definitions of these constructs are consistent with the Columbia Classification Algorithm of Suicide Assessment. Participants were considered positive for suicidal thoughts and behaviors if they reported any of these.
Alcohol and Drug Misuse
The substance use section of the YAPA first asks about frequency of use for specific substances, followed by a detailed section on symptoms and impairment; questions in this section were asked only if substance use was reported. The module assessed symptoms of DSM-5 substance use disorder.19 A 2-week test-retest study to determine the reliability of participant reporting for the number of substance misuse or dependence symptoms had an intraclass correlation coefficient of 0.98.15

For the present analysis, the primary outcome variable for alcohol misuse was meeting full diagnostic criteria for DSM-5 alcohol use disorder (ie, a well-known precursor of liver cirrhosis). The following illicit drugs were assessed for this study: cocaine, crack, amphetamines, methamphetamine, inhalants, nitrite inhalants, ecstasy, heroin, other opioids, oxycodone (could include lawful use), LSD (lysergic acid diethylamide), PCP (phencyclidine), psilocybin, and sedatives. We assessed any illicit drug use meeting the criteria for an illicit drug use disorder, as well as use of any opioids only (ie, oxycodone, other opioids, or heroin). Cannabis was not included in the illicit drug variable because cannabis-related overdoses are not associated with death and have not been studied in the literature on deaths of despair.

Statistical Analysis
Statistical analysis was performed from May 7, 2019, to April 10, 2020. All statistical analyses accounted for the 2-stage sampling design using sampling weights. Each participant was assigned a weight inversely proportional to their probability of selection. All models also used the generalized estimating equations option within SAS PROC GENMOD (SAS Institute Inc) to derive robust variance (sandwich-type) estimates to adjust SEs for the stratified design and multiple observations for each participant. Associations between variables were tested using weighted logistic regression models (for binary outcomes such as suicidal behavior or meeting a DSM diagnosis) and ordered multinomial regression models (for number of despair indicators).

Longitudinal analyses focused on 2424 observations of 1266 individuals between 25 and 30 years of age. In longitudinal models, the given outcome (eg, time 2 suicidality) was regressed on both the lagged values of the outcome variable (time 1 suicidality) and the independent variable (time 1 despair score). Consistent with common conventions, all percentages provided in the results are weighted, and sample sizes are unweighted. All P values were from 2-sided tests and results were deemed statistically significant at \( P < .05 \).

Missing Data
Across all assessments, 83.0% of possible interviews (11,230 of 13,529) were completed. Of the 1420 original participants, 1266 (89.2%) were followed up at least once in young adulthood at 25 or 30 years of age. Despair at earlier assessments was not associated with lower levels of participation in young adulthood. Missing individual despair items within completed young adult interviews were rare (approximately 1%). For generalized linear models, PROC GENMOD excludes any observation with a missing value for any variable involved in the model (approximately 5% of observations).

Results
Prevalence of Despair and Associations With Sociodemographic Factors
The Great Smoky Mountains Study includes 1420 participants (790 male and 630 female participants). The Figure shows the percentage of participants reporting individual indicators of despair and any despair from 9 to 30 years of age. The prevalence of despair was between 5% and 1% in childhood and adolescence (9–21 years) but increased to around 20% in young adulthood (between 25 and 30 years). Levels of any despair and total despair scores increased by age. Each of the individual indicators of despair followed this developmental pattern of increasing with age.
Next, we tested the associations between despair and sociodemographic factors in young adulthood (between 25 and 30 years of age; 2424 observations of 1266 individuals). Table 1 shows the 3-month prevalence of counts on the despair summary scale in young adulthood and associations with sociodemographic factors. The 3-month weighted prevalence of any despair was 19.5% (476 of 2424 observations) and was 7.6% (201 of 2424 observations) for reporting 2 or more items. Despair scores did not differ between male and female participants. Levels of despair differed by race/ethnicity; compared with white participants, African American participants reported much higher rates of despair and Native American participants reported lower levels of despair. Lower educational level and falling below the federal poverty line were associated with higher levels of despair. All sociodemographic factors were subsequently used as covariates in analyses testing associations with the diseases of despair.

Despair was associated with a diagnosis of a DSM depressive disorder (odds ratio [OR], 3.8; 95% CI, 2.9-4.9; \( P < .001 \)). Of the 476 young adult observations in which at least 1 symptom of despair

| Variable                  | Despair score, No. (%) | \( P \) value\(^c\) |
|---------------------------|------------------------|---------------------|
|                           | Score of 0  | Score of 1    | Score of 2    | Score of ≥3   |
| Total                     | 1937 (80.5) | 275 (12.0)    | 94 (4.1)      | 107 (3.5)     | [Reference] |
| Sex                       |            |            |              |              |
| Female                    | 893 (77.7)  | 144 (13.7)   | 54 (5.4)      | 54 (3.2)      | [Reference] |
| Male                      | 1044 (83.5) | 131 (10.2)   | 40 (2.6)      | 53 (3.8)      | .20        |
| Race/ethnicity            |            |            |              |              |
| White                     | 1288 (81.6) | 217 (11.4)   | 72 (3.8)      | 93 (3.3)      | [Reference] |
| African American          | 106 (63.1)  | 27 (21.9)    | 13 (8.2)      | 8 (6.9)       | <.001\(^d\) |
| Native American           | 543 (92.2)  | 31 (5.3)     | 9 (1.5)       | 6 (1.0)       | <.001\(^d\) |
| Educational level         |            |            |              |              |
| No high school degree     | 249 (69.8)  | 42 (17.5)    | 17 (7.5)      | 20 (5.2)      | .02\(^d\)   |
| High school only          | 418 (76.1)  | 67 (15.1)    | 14 (2.6)      | 31 (6.2)      | .04\(^d\)   |
| Some college              | 642 (80.6)  | 94 (9.2)     | 45 (5.6)      | 46 (6.6)      | .04\(^d\)   |
| 4-y Degree                | 519 (83.0)  | 72 (13.0)    | 18 (3.0)      | 10 (1.1)      | [Reference] |
| Poverty                   |            |            |              |              |
| Yes                       | 353 (64.6)  | 96 (18.9)    | 29 (6.3)      | 56 (10.2)     | [Reference] |
| No                        | 1449 (83.7) | 170 (10.6)   | 64 (3.6)      | 50 (2.0)      | <.001\(^d\) |

\(^{a}\) Based on 2424 observations of 266 individuals assessed at 25 and 30 years of age.

\(^{b}\) All percentages are weighted, and all numbers are unweighted.

\(^{c}\) Representing results from ordered multinomial regression regressing despair scores on young adult demographic variables.

\(^{d}\) Significant at \( P < .05 \).
was reported, however, 94 (19.7%) of them met criteria for a DSM depressive disorder, suggesting that despair encompasses a much broader group than those with depression.

**Associations With Suicidal Thoughts and Behaviors, Alcohol Misuse, and Illicit Drug Misuse**

Table 2 shows concurrent associations between despair scores in young adulthood and putative outcomes adjusted for sociodemographic factors. Higher despair scores were associated with higher levels of suicidal thoughts and behavior, illicit drug use, illicit drug disorders, opioid use, and any disease of despair. Despair scores were not significantly associated with alcohol use disorder between 25 and 30 years of age.

Bidirectional temporal associations between despair and outcomes were tested using a series of longitudinal lagged models (Table 3). Outcome status at 25 or 30 years of age was regressed on status of variables at the prior wave (eg, suicidal thoughts and behavior at 30 years regressed on despair score at 25 years). Models were adjusted for status of the outcome variable at the prior wave (eg, suicidal thoughts and behavior at 25 years). Because each lagged model is based on 2 time points, each participant could contribute up to 2 observations to these analyses. Models were performed using generalized estimating equations to account for repeated observations. All sociodemographic covariates were included.

The first 5 rows of results in Table 3 display logistic regression models estimating young adult outcomes (between 25 and 30 years of age; 2424 observations of 1266 individuals) from previous despair levels. Prior despair was associated with elevated levels of suicidal thoughts and behaviors (OR, 1.5; 95% CI, 1.1-2.0), illicit drug use (OR, 1.7; 95% CI, 1.2-2.5), and opioid use (OR, 1.9; 95% CI, 1.1-3.3) but not with alcohol use disorder (OR, 0.8; 95% CI, 0.6-1.2) or illicit drug disorder (OR, 1.0; 95% CI 0.6-1.7) (Table 3 and eTable 3 in the Supplement). The ORs represent the change in odds of the outcome given a 1-unit increase in the despair score (range, 0-3). In analyses assessing the difference between different despair scores, risk tended to increase with the number of despair items reported, with evidence that 2 or more items were associated with moderate to large effects.

### Table 2. Associations of Despair Score With Young Adult (at 25 and 30 Years of Age) Outcomes at the Same Observationa

| Outcome                          | Participants, No. (%) | Despair scores | P valueb | Total | 0 | 1 | 2 | 3+ | NA |
|----------------------------------|------------------------|----------------|----------|-------|---|---|---|----|----|
| Total                            | 2424 (100)             | 1937 (80.5)    | 275 (12.0) | 94 (4.1) | 107 (3.5) | NA |
| Suicide                          |                        |                |          |       |   |   |   |     |     |
| Yes                              | 160 (5.9)              | 83 (3.6)       | 29 (9.7)  | 17 (13.1) | 31 (27.9) | <.001d |
| No                               | 2253 (94.1)            | 1854 (96.4)    | 246 (90.3) | 77 (86.9) | 76 (72.1) |     |
| Alcohol use disorder             |                        |                |          |       |   |   |   |     |     |
| Yes                              | 132 (7.6)              | 99 (6.7)       | 18 (10.7) | 8 (9.8)  | 7 (3.4)  | .48 |
| No                               | 2292 (92.4)            | 1838 (93.3)    | 257 (89.3) | 86 (90.2) | 100 (96.6) |     |
| Illicit drug use                 |                        |                |          |       |   |   |   |     |     |
| Yes                              | 137 (4.7)              | 85 (5.7)       | 24 (5.6)  | 8 (6.8)  | 20 (18.5) | .04d|
| No                               | 2247 (95.3)            | 1824 (94.3)    | 250 (94.4) | 86 (93.2) | 87 (81.5) |     |
| Illicit drug use disorder        |                        |                |          |       |   |   |   |     |     |
| Yes                              | 88 (2.9)               | 46 (1.4)       | 16 (5.6)  | 7 (6.0)  | 19 (17.6) | <.001d |
| No                               | 2336 (97.1)            | 1891 (98.6)    | 259 (94.4) | 87 (94.0) | 88 (82.4) |     |
| Opioid use                       |                        |                |          |       |   |   |   |     |     |
| Yes                              | 55 (2.2)               | 31 (1.6)       | 8 (1.0)   | 6 (5.9)  | 10 (13.6) | .07d|
| No                               | 2328 (97.8)            | 1877 (98.4)    | 266 (99.0) | 88 (94.1) | 97 (86.4) |     |

Abbreviation: NA, not applicable.

a Based on 2424 observations of 1266 individuals.
b All percentages are weighted, and all numbers are unweighted. Covariates included sex, race/ethnicity, educational level, and poverty.
c Representing results from logistic models regressing young adult outcome variables on concurrent despair scores.
d Significant at P < .05.
Finally, the adjusted lagged models were performed again, adjusted for lagged depressive disorder status (in addition to other covariates). The aim was to test whether the lagged association of despair was accounted for by depression or whether despair was independently associated with young adult outcomes. In the model adjusted for depressive disorder (Table 3), lagged despair was still associated with suicidal thoughts and behaviors (OR, 1.5; 95% CI, 1.1-2.1), illicit drug use (OR, 1.7; 95% CI, 1.1-2.8), and opioid use (OR, 1.9; 95% CI, 1.0-3.5).

The final 5 rows of Table 3 show results from ordered multinomial regression models testing the reverse pattern of prior status of suicidal thoughts and behaviors and substance use variables estimating current despair scores. None of these diseases were associated with later despair scores. All models in Table 3 were performed again, with 3 variations: (1) using a nonwinsorized despair variable (ie, full range from 0 to 7), (2) using a dichotomous despair variable for the presence or absence of any despair indicators, and (3) using 7 despair sum scores that systematically left out 1 of the indicators. In all cases, the pattern of results was similar to those reported in Table 3. These results are available on request from the first author.

### Long-term Despair Exposure

Table 4 shows 2 alternative definitions of long-term despair exposure: (1) total despair scores across all prior observations (cumulative despair levels) and (2) total observations reporting any despair (cumulative despair observations). Each measure was averaged over the total number of available observations for that individual. Thus, the range for cumulative despair levels was from 0 to 4.5, and the range for cumulative despair observations was from 0 to 1. Both measures of long-term despair

| Outcome                          | Cumulative despair levels | Cumulative despair observations |
|----------------------------------|---------------------------|---------------------------------|
| Suicidal thoughts or behaviors   | 2.6 (1.5-4.7)             | 10.7 (3.5-32.9)                 |
| Alcohol use disorder             | 0.9 (0.5-1.7)             | 0.9 (0.2-3.7)                   |
| Illicit drug use                 | 2.5 (1.3-5.0)             | 6.0 (1.6-22.9)                  |
| Illicit drug use disorder        | 2.3 (1.3-4.4)             | 9.4 (2.4-36.7)                  |
| Opioid use                       | 3.3 (1.4-7.7)             | 7.5 (0.9-65.5)                  |

Abbreviation: OR, odds ratio.

* Based on 2424 observations of 1266 individuals. The top 5 rows represent results from logistic regression regressing young adult outcome variables on lagged despair scores. The bottom 5 rows represent results from ordered multinomial models regressing despair scores on lagged young suicidality and substance use variables. Sociodemographic covariates included sex, race/ethnicity, educational level, poverty, and lagged value of the outcome variable.

b Significant at \( P < .05 \).
were associated with increased risk for suicide (cumulative despair levels: OR, 2.6; 95% CI, 1.5-4.7; cumulative despair observations: OR, 10.7; 95% CI, 3.5-32.9), illicit drug use (cumulative despair levels: OR, 2.5; 95% CI, 1.3-5.0; cumulative despair observations: OR, 6.0; 95% CI, 1.6-22.9), illicit drug use disorder (cumulative despair levels: OR, 2.3; 95% CI, 1.3-4.4; cumulative despair observations: OR, 9.4; 95% CI, 2.4-36.7), and opioid use (cumulative despair levels: OR, 3.3; 95% CI, 1.4-7.7; cumulative despair observations: OR, 7.5; 95% CI, 0.9-65.5). In all cases, the associations were stronger than those observed in the lagged models, suggesting the importance of indexing long-term despair exposure.

**Moderation by Educational Level, Race/Ethnicity, Sex, and Poverty**

A series of models tested whether observed associations between despair and later outcomes were moderated by sociodemographic factors. For example, does this association differ between male and female participants or by educational level? Moderation was tested with an interaction term between despair and the sociodemographic factor. eTable 5 in the Supplement shows the result of the moderation tests for each of the variables. Although the results of 4 of the 20 interaction tests were significant, there was no consistent pattern of moderation, either in terms of outcomes or moderators. Follow-up analyses were conducted for each moderator group separately to facilitate future systematic reviews or meta-analyses (eTables 6, 7, 8, and 9 in the Supplement).

**Discussion**

Despair has been postulated as a cause of the recent increase in premature mortality associated in part with suicide and alcohol- and drug-related deaths. This study tested the longitudinal associations between despair and suicidal thoughts and behavior and substance misuse in young adulthood. Despair scores were highest among those with lower educational level and income status and among African American participants. Despair scores were longitudinally associated with increased levels of suicidal thoughts and behavior, illicit drug use, illicit drug use disorders, and opioid use over time but not alcohol use disorder. These associations were observed even after accounting for depression status. The observed despair-outcome associations were stronger in models accounting for long-term measures of despair. This study provides an empirical basis for despair as longitudinally associated with suicidality and illicit substance use.

How do our findings inform our understanding of diseases and deaths of despair? Despair was both concurrently associated with the outcomes and preceded them. These associations persisted in models accounting for a number of other sociodemographic factors strongly associated with these diseases as well as depressive disorders. Our findings suggest a 1.4 to 1.7 times prospective increase in the odds of each outcome for each additional indicator of despair reported. (In models using a dichotomized despair variable, the ORs were around 2 to 3.) The associations were stronger when accounting for long-term despair exposure extending back to childhood. In contrast, none of the diseases were associated with increased levels of despair in longitudinal models. These results suggest that the pathway from despair to diseases of despair is unidirectional.

In this analysis, which tested a limited number of outcomes, the associations of despair were specific to suicidal behavior, illicit drug use, illicit drug use disorder, and opioid use. These findings are consistent with evidence from other studies that suggest that suicidal behavior and drug use are a result of diminished valuing of oneself and one’s future. Despair was not associated with alcohol use disorder, however. Alcohol use disorder was equally common among those with the lowest levels of despair as in those with the highest levels of despair. This study does not support alcohol use disorder as a disease of despair.

As expected, despair was more common among those with low educational level or limited financial resources. At the same time, the associations of despair with outcomes were neither stronger nor weaker in specific sociodemographic groups. In other words, the relative associations of despair with outcomes did not differ systematically between groups. This finding is consistent with
recent reports that the newest premature mortality trends appear to be mostly universal across US racial/ethnic groups. Accordingly, public policy strategies should focus on population-level interventions rather than targeting specific subgroups to reduce despair.

Despair is a new construct, but the associated DSM-based depressive disorders have been studied intensely for the past 50 years or more. Is despair useful or is it simply a new way of framing an old condition? In this study, many of those who met the criteria for a depressive disorder also displayed despair. However, the association was not reciprocal: most individuals reporting despair did not meet the criteria for a depressive disorder. It might be expected that individuals with both despair and depressive disorder would be at risk for suicidal thoughts and behavior and substance misuse, but this was not the case. Depression is a complex medical disorder seen in comparatively few individuals, whereas despair is a cognitive status experienced by many individuals that may reflect the social and economic deterioration of many US communities. In this study, despair scores had an independent utility in understanding risk for these high-priority diseases.

This literature has understandably focused on the extreme outcomes of reduced life expectancy and premature mortality. Our findings suggest that despair is common in young adulthood and is associated with destructive behaviors that may or may not lead to mortality. The focus on mortality alone may miss a broader association of despair with impairment and reduced quality of life. In the language of the Global Burden of Disease study, the public health outcome of the disability-adjusted life-years associated with despair may far outstrip the association with mortality itself. Future studies should not only focus on outcomes associated with mortality but also explore how despair is associated with important areas of life functioning, such as financial and social functioning and physical health.

Limitations
This study has some limitations. The sample is not representative of the US population. Measures of despair were obtained from parent and child report to 16 years of age and only from self-report thereafter. The despair scale was created for the present study post hoc using items available from the structured interview. The focus of each observation was on despair indicators within the prior 3 months only, thus allowing it to miss despair in the intervening periods. Although participation rates were high at young adult assessments (>80%), the results may be affected by missing observations. Also, we studied common precursors of deaths of despair rather than mortality per se because the latter was not possible with our young adult sample. This study did not clarify whether opioid use was prescription or nonprescription, which may lead to some misclassification. More important, the opioid use results closely mirror the results for other illicit drug use, suggesting that such misclassification is limited. Finally, the study is a cohort study and unable to test causal associations between despair and suicidality or substance misuse.

Conclusions
In the play Caesar and Cleopatra, George Bernard Shaw observes “he who has not hoped can never despair.” Despair may be a risk factor for diseases, but it is also the end point of a process in which hope is lost. The public health successes of the 20th century have consistently allowed children to live longer than their parents. It is precisely because this recent pattern deviates from a century of progress that it has captured the attention of the public and the scientific community. This study takes a step toward understanding how a psychological state could derail the seemingly inexorable progress of modern medicine. However, this study is only 1 step, and additional work is needed to understand the origins of despair and premature mortality within individuals, families, and communities and how we can intervene to recover hope and forestall further morbidity and mortality.
ARTICLE INFORMATION

Accepted for Publication: April 16, 2020.
Published: June 23, 2020. doi:10.1001/jamanetworkopen.2020.8627

Open Access: This is an open access article distributed under the terms of the CC-BY License. © 2020 Copeland WE et al. JAMA Network Open.

Corresponding Author: William E. Copeland, PhD, Vermont Center for Children, Youth, and Families, Department of Psychiatry, University of Vermont College of Medicine, One S Prospect St, PO Box 3454, Burlington, VT 05401 (william.copeland@med.uvm.edu).

Author Affiliations: Vermont Center for Children, Youth, and Families, Department of Psychiatry, University of Vermont College of Medicine, Burlington (Copeland); Public Policy Studies, Center for Medicine, Health, and Society, Vanderbilt University, Nashville, Tennessee (Gaydosh); Department of Psychiatry, Duke University Medical Center, Durham, North Carolina (Hill); Center for Child and Family Policy, Duke University, Durham, North Carolina (Godwin); Carolina Population Center, Department of Sociology, University of North Carolina at Chapel Hill (Harris); Department of Psychiatry and Behavioral Sciences, Duke University Medical Center, Durham, North Carolina (Costello); Jacobs Center for Productive Youth Development, Department of Psychology, University of Zürich, Zürich, Switzerland (Shanahan).

Author Contributions: Dr Copeland had full access to all of the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis.

Concept and design: Copeland, Gaydosh, Hill, Harris, Shanahan.

Acquisition, analysis, or interpretation of data: Copeland, Gaydosh, Godwin, Harris, Costello.

Drafting of the manuscript: Copeland, Gaydosh, Shanahan.

Critical revision of the manuscript for important intellectual content: Copeland, Gaydosh, Hill, Godwin, Harris, Costello.

Statistical analysis: Copeland, Harris.

Obtained funding: Copeland, Harris, Costello, Shanahan.

Administrative, technical, or material support: Gaydosh, Hill, Godwin.

Supervision: Copeland, Costello.

Conflict of Interest Disclosures: Drs Copeland, Hill, and Godwin reported receiving grants from the National Institute of Mental Health during the conduct of the study. Dr Hill reported receiving grants from the Eunice Kennedy Shriver National Institute of Child Health and Human Development (training grant) and the National Institute on Drug Abuse; and personal fees from University of North Carolina at Chapel Hill Frank Porter Graham Child Development Institute outside the submitted work. No other disclosures were reported.

Funding/Support: The work presented here was supported by grants MH117559, MO80230, MH63970, MH63671, MH48085, MH075766, MH094605, MH104576 from the National Institute of Mental Health; grants DAO16977, DAO13130, DAO36523, DAO23026 from the National Institute on Drug Abuse; grant HD093651 from the National Institute of Child health and Development; Brain and Behavior Research Foundation (Early Career Award to Dr Copeland); and the William T Grant Foundation.

Role of the Funder/Sponsor: The funding sources had no role in the design and conduct of the study; collection, management, analysis, and interpretation of the data; preparation, review, or approval of the manuscript; and decision to submit the manuscript for publication.

REFERENCES

1. Case A, Deaton A. Rising morbidity and mortality in midlife among white non-Hispanic Americans in the 21st century. Proc Natl Acad Sci USA. 2015;112(49):15078-15083. doi:10.1073/pnas.1518393112

2. Woolf SH, Schoomaker H. Life expectancy and mortality rates in the United States, 1959-2017. JAMA. 2019;322(20):1996-2016. doi:10.1001/jama.2019.16932

3. Stein EM, Gennuso KP, Ugboaja DC, Remington PL. The epidemic of despair among white Americans: trends in the leading causes of premature death, 1999-2015. Am J Public Health. 2017;107(10):1541-1547. doi:10.2105/AJPH.2017.303941

4. Hu G, Wilcox HC, Wissow L, Baker SP. Mid-life suicide: an increasing problem in U.S. whites, 1999-2005. Am J Prev Med. 2008;35(6):589-593. doi:10.1016/j.amepre.2008.07.005

5. Shiels MS, Chernyavskiy P, Anderson WF, et al. Trends in premature mortality in the USA by sex, race, and ethnicity from 1999 to 2014: an analysis of death certificate data. Lancet. 2017;389(10073):1043-1054. doi:10.1016/S0140-6736(17)30187-3
6. Shanahan L, Hill SN, Gaydosh LM, et al. Does despair really kill? a roadmap for an evidence-based answer. Am J Public Health. 2019;109(6):854-858. doi:10.2105/AJPH.2019.305016

7. “Despair, n.” In: Oxford English Dictionary. Oxford University Press; 2020.

8. Erwin PC. Despair in the American heartland? a focus on rural health. Am J Public Health. 2017;107(10):1533-1534. doi:10.2105/AJPH.2017.304029

9. Meit M, Heffernan M, Tanebaum E, Hoffmann T. Appalachian diseases of despair: report for the Appalachian Regional Commission. The Walsh Center for Rural Health Analysis National Opinion Research Center at the University of Chicago. Published August 2017. Accessed May 7, 2019. https://www.arc.gov/assets/research_reports/AppalachianDiseasesofDespairAugust2017.pdf

10. Costello EJ, Mustillo S, Erkanli A, Keeler G, Angold A. Prevalence and development of psychiatric disorders in childhood and adolescence. Arch Gen Psychiatry. 2003;60(8):837-844. doi:10.1001/archpsyc.60.8.837

11. Costello EJ, Angold A, Burns B, et al. The Great Smoky Mountains Study of Youth: goals, design, methods, and the prevalence of DSM-III-R disorders. Arch Gen Psychiatry. 1996;53(12):1129-1136. doi:10.1001/archpsyc.1996.01830120067012

12. Copeland WE, Angold A, Shanahan L, Costello EJ. Longitudinal patterns of anxiety from childhood to adulthood: the Great Smoky Mountains Study. J Am Acad Child Adolesc Psychiatry. 2014;53(1):21-33. doi:10.1016/j.jaac.2013.09.017

13. von Elm E, Altman DG, Egger M, Pocock SJ, Gøtzsche PC, Vandenbroucke JP. STROBE Initiative. The Strengthening the Reporting of Observational Studies in Epidemiology (STROBE) statement: guidelines for reporting observational studies. Ann Intern Med. 2007;147(8):573-577. doi:10.7326/0003-4819-147-8-200710160-00010

14. Angold A, Costello EJ. The Child and Adolescent Psychiatric Assessment (CAPA). J Am Acad Child Adolescent Psychiatry. 2000;39(1):39-48. doi:10.1097/00004583-200001000-00015

15. Angold A, Costello EJ. A test-retest reliability study of child-reported psychiatric symptoms and diagnoses using the Child and Adolescent Psychiatric Assessment (CAPA-C). Psychol Med. 1995;25(4):755-762. doi:10.1017/S0033291700034991

16. Angold A, Cox A, Prendergast M, et al. The Young Adult Psychiatric Assessment (YAPA). Duke University Medical Center; 1999.

17. Case A, Deaton A. Mortality and morbidity in the 21st century. Brookings Pap Econ Act. 2017;2017:397-476. doi:10.1353/eca.2017.0005

18. Posner K, Oquendo MA, Gould M, Stanley B, Davies M. Columbia Classification Algorithm of Suicide Assessment (C-CASA): classification of suicidal events in the FDA's pediatric suicidal risk analysis of antidepressants. Am J Psychiatry. 2007;164(7):1035-1043. doi:10.1176/ajp.2007.164.7.1035

19. American Psychiatric Association. Diagnostic and Statistical Manual of Mental Disorders. 5th ed. American Psychiatric Association; 2013.

20. Kraemer HC, Kazdin AE, Offord DR, Kessler RC, Jensen PS, Kupfer DJ. Coming to terms with the terms of risk. Arch Gen Psychiatry. 1997;54(4):337-343. doi:10.1001/archpsyc.1997.01830160065009

21. Bodner E, Bergman YS. Loneliness and depressive symptoms among older adults: the moderating role of subjective life expectancy. Psychiatry Res. 2016;237:78-82. doi:10.1016/j.psychres.2016.01.074

22. Piquero AR. “Take my license n all that jive, I can’t see...35”: little hope for the future encourages offending over time. Justice Q. 2016;33(1):73-99. doi:10.1080/07418825.2014.896396

23. Scott-Sheldon LA, Carey MP, Vanable PA, Senn TE. Subjective life expectancy and health behaviors among STD clinic patients. Am J Health Behav. 2010;34(3):349-361. doi:10.5993/AJHB.34.3.10

24. Hyde JS, Mezzulis AH, Abramson LY. The ABCs of depression: integrating affective, biological, and cognitive models to explain the emergence of the gender difference in depression. Psychol Rev. 2008;115(2):291-313. doi:10.1037/0033-295X.115.2.291

25. Kendler KS, Gardner CO, Prescott CA. Toward a comprehensive developmental model for major depression in men. Am J Psychiatry. 2006;163(1):115-124. doi:10.1176/appi.ajp.163.1.115

26. Murray C.L., Lopez AD. The Global Burden of Disease. Vol 1. World Health Organization; 1996.

27. Shaw GB. Three Plays for Puritans: The Devil’s Disciple, Caesar and Cleopatra, & Captain Brassbound’s Conversion. G. Richards; 1901.
SUPPLEMENT.

**Table 1.** Individual Indicators of Cognitive Despair and Their Definitions

**Table 2.** Definitions of Different Types of Suicidal Thoughts and Behaviors

**Table 3.** Associations of Lagged Despair Scores With Young Adult (age 25, 30) Outcomes

**Table 4.** Models Predicting Young Adult Outcomes (ages 25 and 30) Comparing Different Levels of Lagged Despair (0 vs 1, 2, 3)

**Table 5.** Lagged Longitudinal Models Testing Interactions Between Despair and Each of, Sex, Race/Ethnicity, Poverty, and Educational Attainment in the Prediction of Each Outcome

**Table 6.** Lagged Models Predicting Young Adult Outcomes Separately by Poverty Status

**Table 7.** Lagged Models Predicting Young Adult Outcomes Separately by Educational Attainment

**Table 8.** Lagged Models Predicting Young Adult Outcomes Separately by Race/Ethnicity

**Table 9.** Lagged Models Predicting Young Adult Outcomes Separately by Sex

**Figure.** Ascertainment of the Original Great Smoky Mountains Study Sample