



Original article

Predicting the incidence of non-suicidal self-injury in college students

G. Kiekens^{a,b,*}, P. Hasking^b, L. Claes^{c,d}, M. Boyes^b, P. Mortier^e, R.P. Auerbach^f,
P. Cuijpers^g, K. Demyttenaere^a, J.G. Green^h, R.C. Kesslerⁱ, I. Myin-Germeys^j, M.K. Nock^k,
R. Bruffaerts^{a,l}

^a Center for Public Health Psychiatry, KU Leuven, Leuven, Belgium

^b School of Psychology, Curtin University, Perth, Australia

^c Faculty of Psychology and Educational Sciences, KU Leuven, Leuven, Belgium

^d Faculty of Medicine and Health Sciences (CAPRI), University of Antwerp, Antwerp, Belgium

^e Health Services Research Unit, IMIM (Hospital del Mar Medical Research Institute), Barcelona, Spain

^f Department of Psychiatry, Vagelos College of Physicians and Surgeons, Columbia University, New York, NY, USA

^g Department of Clinical, Neuro and Developmental Psychology, Amsterdam Public Health Research Institute, Vrije Universiteit Amsterdam, Amsterdam, the Netherlands

^h Wheelock College of Education & Human Development, Boston University, Boston, MA, USA

ⁱ Harvard Medical School, Department of Health Care Policy, Harvard University, Boston, MA, USA

^j Department of Neurosciences, Centre for Contextual Psychiatry, KU Leuven, Leuven, Belgium

^k Department of Psychology, Harvard University, Cambridge, MA, USA

^l Institute for Social Research, Population Studies Center, University of Michigan, Ann Arbor, MI, USA



ARTICLE INFO

Article history:

Received 18 December 2018

Received in revised form 11 March 2019

Accepted 4 April 2019

Available online xxx

Keywords:

NSSI

Self-injury

Incidence

Prediction

Emerging adults

Prevention

ABSTRACT

Background: Despite increased awareness that non-suicidal self-injury (NSSI) poses a significant public health concern on college campuses worldwide, few studies have prospectively investigated the incidence of NSSI in college and considered targeting college entrants at high risk for onset of NSSI.

Methods: Using data from the Leuven College Surveys ($n = 4,565$; 56.8%female, $M_{age} = 18.3$, $SD = 1.1$), students provided data on NSSI, sociodemographics, traumatic experiences, stressful events, perceived social support, and mental disorders. A total of 2,163 baseline responders provided data at a two-year annual follow-up assessment (63.2% conditional response rate).

Results: One-year incidence of first onset NSSI was 10.3% in year 1 and 6.0% in year 2, with a total of 8.6% reporting sporadic NSSI (1–4 times per year) and 7.0% reporting repetitive NSSI (≥ 5 times per year) during the first two years of college. Many hypothesized proximal and distal risk factors were associated with the subsequent onset of NSSI (ORs = 1.5–18.2). Dating violence prior to age 17 and severe role impairment in daily life were the strongest predictors. Multivariate prediction suggests that an intervention focused on the 10% at highest risk would reach 23.9% of students who report sporadic, and 36.1% of students who report repetitive NSSI during college (cross-validated AUCs = .70–.75).

Discussion: The college period carries high risk for the onset of NSSI. Individualized web-based screening may be a promising approach for detecting young adults at high risk for self-injury and offering timely intervention.

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1. Introduction

The college years mark the transition from adolescence to emerging adulthood; a unique developmental period characterized by increasing opportunities in academic, personal, and social areas of life [1]. Yet, this period also is one of heightened risk for

mental disorders and risky behaviors [2–5]. Non-suicidal self-injury (NSSI), the intentional damage to one's body tissue (e.g., scraping the skin; self-battery) without suicidal intent [6], is becoming increasingly recognized as a public health concern on college campuses. While NSSI has historically been conceptualized as a symptom of mental disorders, there is now increased awareness that NSSI is not symptomatic of any particular disorder and should be conceptualized as a behavior that warrants research and intervention in its own right [7–12]. International pooled lifetime prevalence estimates of NSSI are around 20% in college students [13], with 12-month estimates in the 2–14% range [14,15].

* Corresponding author at: Kapucijnenvoer 33 building I, box 7001, 3000, Leuven, Belgium.

E-mail address: Glenn.Kiekens@kuleuven.be (G. Kiekens).

Although not always associated with suicide risk, NSSI (especially repetitive and severe self-injury) is one of the strongest independent predictors of future suicide attempts [16–19], and is associated with severe role impairment in daily life [9], stigma and feelings of shame [20,21], low levels of help-seeking [22,23], and poorer academic performance [24]. These findings underscore the necessity to address and respond to NSSI among college students [25].

Although many colleges have begun to implement risk assessment and prevention programs for mental health problems [26,27], NSSI is rarely included in these efforts. Importantly, however, although NSSI onset peaks in mid-adolescence [28], recent evidence suggests a second peak around the age of 20 [29]. Hence, there is potential to reach out to students *before* NSSI and associated negative outcomes occur. Franklin and colleagues recently demonstrated a reduction in frequency of NSSI through use of a mobile app that focuses on the barriers to NSSI (e.g., by increasing self-worth) [30]. It is possible that similar initiatives may be effective in preventing onset among at risk students. However, there is not yet any evidence-based method of identifying students at risk for onset of NSSI. This is in part due to two important limitations in the literature. First, most of what is known about potential risk factors stems from cross-sectional approaches investigating correlates among convenience samples. Whereas these studies can provide clues about potential predictors of interest, the nature of the designs and samples limit the generality of the findings [31]. Second, the prospective studies available focused primarily on risk factors for NSSI persistence (i.e., ongoing vs. ceased NSSI) [15,32–35]. While these studies provide valuable information to aid clinical decision making for persons who already engage in NSSI, it does not provide means to detect young adults who are likely to start engaging in NSSI. Two earlier studies reported 9–12-month onset rates in the 2–4% range in college students [33,34]. However, these studies may have failed to include a representative college sample (e.g., only college women, convenience sampling) and incorporated a narrow set of predictors, precluding the development of integrative prediction models to detect students at highest risk for NSSI onset. We aimed to address these gaps in the literature making use of a large longitudinal sample of college students from the Leuven College Surveys (LCS) [36], part of the WHO World Mental Health International College Student (WMH-ICS) initiative [37].

Our objectives were to: 1) estimate the incidence of NSSI during the first two years of college, 2) examine a broad range of proximal and distal risk factors, and 3) evaluate the accuracy of a multivariate risk prediction model for the onset of NSSI. In line with the proposition that NSSI is a complex behavior that is determined by a multitude of factors [38], we did not anticipate a clear set of risk factors for onset NSSI. Potential predictors we assessed are well-established correlates of NSSI in college students, including sociodemographic and college-related characteristics [23,24], childhood-adolescent trauma [39–41], recent stressful experiences and perceived social support [15,35,40], and mental disorders [9,12,42]. Consistent with the proposed DSM-5 frequency criterion (i.e., self-injury on 5 or more days in the past year) [43], we determined the predictive value of risk factors separately for the onset of both sporadic and repetitive NSSI (i.e., ≥ 5 times per year), occurring during the first two years of college.

2. Method

2.1. Procedures and sample description

Detailed recruitment strategies of the LCS have previously been described [4,24]. Recruitment involved different strategies to increase the response rate. In the first phase, all incoming students

were sent a standard invitation letter to a routine psycho-medical checkup organized by the university student health center, which included the survey. In the second phase, secured electronic links were sent to non-respondents using customized e-mails. The third phase was identical to the second, but included an additional incentive (i.e., a raffle for store coupons). Each phase included reminders, with eight as default maximum amount of contacts. In the academic years 2014–2015 and 2015–2016, all 8,530 first year students were invited to participate, and a total of 4,565 students completed the baseline survey (56.8% female, $M_{\text{age}} = 18.3$, $SD = 1.1$; Response Rate = 53.5%). Baseline responders were invited via email, to participate in the follow-up surveys 12 and 24 months after the baseline assessment. A total of 2,163 of the baseline respondents participated in at least one follow-up survey (63.2% conditional response rate after adjusting for non-participation due to college attrition). Informed consent was obtained from all participants at each wave and the study's protocol was approved by the University's Ethical Review Board.

2.2. Measures

Sociodemographic and college-related variables included gender, age, nationality, perceived parental financial situation, parental educational level, family composition, subject area enrollment (e.g., biomedical sciences, science and technology), and type of secondary school attended (i.e., vocational vs. academic track).

Non-suicidal self-injury was assessed with the self-report version of the well-validated Self-Injurious Thoughts and Behaviors Interview [44]. The self-report version showed excellent test-retest reliability and external validity in a comparison study of self-report questionnaires [45]. To assess lifetime NSSI thoughts, we asked respondents whether they ever had “thoughts of purposely hurting themselves, without wanting to die”. Incident NSSI was assessed via a checklist of 13 NSSI methods (e.g., cutting, burning, hitting) and an ‘other’ category that asked respondents whether they used that NSSI method to “hurt themselves on purpose, without wanting to die”. Using follow-up questions, we assessed whether students engaged in 12-month sporadic (i.e., 1–4 times) or repetitive NSSI (i.e., ≥ 5 times).

Traumatic experiences prior to the age of 17 were assessed using 19 items from the Composite International Diagnostic Interview-3.0 [CIDI; 46], the Adverse Childhood Experience Scale [47], and the Bully Survey [48]. Items assessed parental psychopathology (i.e., any serious mental health problems, criminal activities, or interpersonal violence), physical abuse (e.g., family member hit you so hard that it left bruises), emotional abuse (e.g., family member repeatedly said hurtful or insulting things), sexual abuse (e.g., family member touched or made you touch them in a sexual way against your will), neglect (i.e., nobody took care of you, or protected you, or made sure you had the things you needed), dating violence (i.e., you were in a romantic relationship where your partner repeatedly hit you or hurt you) and bully victimization (including verbal, physical, and cyberbullying). Items were rated on a five-point Likert scale (“never,” “rarely,” “sometimes,” “often,” and “very often”). Previous research using confirmatory factor analysis showed an excellent fit of the factor structure of the used items [4]. In order to obtain dichotomously coded variables for the calculation of population attributable risk proportions, “rarely” was used as the cutoff for experiencing each traumatic event, except bully victimization where “sometimes” was used as cutoff [49].

Stressful experiences and perceived social support. Making use of well-validated screeners [50–52], we assessed a range of 12-month stressful experiences (e.g., life-threatening illness or injury of family member or close friend). Using the Social Network

section of the CIDI-3.0 [46], participants indicated whether they felt they could rely on family, friends, and partner (when present) if they had a serious problem. Items were rated on a four-point scale (“a lot”, “some”, “a little”, and “not at all”). To allow the calculation of population attributable risk proportions, networks that were perceived as unavailable for support “a lot” were coded as unsatisfactory.

Risk for mental disorders and associated impairment. The CIDI Screening Scales [46,53], developed by the World Health Organization to deliver prevalence estimates of DSM-IV mental disorders, were used to assess 12-month Major Depressive Disorder, Broad Mania (mania/hypomania), Generalized Anxiety Disorder, and Panic Disorder. The Alcohol Use Disorders Identification Test (AUDIT) was used to identify students engaging in “risky or hazardous drinking” and students with “risk for alcohol dependence” [54]. Using additional items from the CIDI-3.0, we also screened students for intermittent explosive disorder (i.e., history of repeated attacks of anger when suddenly you lost control and either broke or smashed something, hit or tried to hurt someone, or threatened someone), eating disorders (i.e., binges at least twice a week or history of vomiting or taking laxatives or other things to avoid gaining weight), psychotic disorder (i.e., history of seeing things other people couldn't see or hear, or having thoughts like believing your mind was being controlled by outside forces) and post-traumatic stress disorder (i.e., times lasting 1 month or longer after an extremely stressful experience when you had repetitive upsetting memories or dreams, felt emotionally distant or depressed, and had trouble sleeping or concentrating). Finally, using the Sheehan Disability Scale, 12-month severe role impairment in daily life was assessed [46,55].

Twelve-month suicidal thoughts and behaviors were assessed with a modified version of the Columbia Suicidal Severity Rating Scale and included suicide ideation (i.e., having thoughts of killing yourself), suicide plan (i.e., thinking about how you might kill yourself or working out a plan of how to kill yourself), and a suicide attempt (i.e., purposefully hurt yourself with at least some intent to die) [56].

2.3. Statistical analyses

Appropriate missing data strategies were used to ensure that findings are representative for the entire student population. Specifically, non-response propensity weights were calculated based on sociodemographic and college-related variables available for the entire first year cohort [24], and multivariate imputation by chained equations was used to adjust for survey attrition and within-survey item nonresponse [57]. Using the package `mice()` in R [58], the final data consisted of 200 imputed datasets obtained after 100 iterations. For the purpose of this study, analyses were restricted to students reporting no prior history of NSSI at baseline ($n = 3,761$). Descriptive statistics and incidence estimates are reported as weighted numbers (n), and weighted proportions (%) with associated standard errors. One-year NSSI incidence proportions were calculated by using first onset NSSI follow-up cases as the numerator, and cases without NSSI at the previous wave as the denominator. Logistic regression analysis was used to test the strength of associations between risk indicators recorded at baseline and incident NSSI. Measures of association were reported as odds ratios and associated 95% confidence intervals. Each risk factor was evaluated in bivariate and multivariate models within risk domains. Population-level effect sizes were estimated using population attributable risk proportions [PARPs; 59]. PARPs provide an estimate of the proportion of cases that would potentially be prevented if it were possible to fully eliminate causal risk factor(s) under examination.

Based on multivariate equations including all risk factors in the study (>50 beta coefficients), we then calculated individual

cumulative risk probabilities. Resulting Area Under the Curve (AUC) values close to .56, .64, and .71 are considered, respectively, as small, moderate, and large effects [60]. Predicted probabilities were discretized into deciles (10 groups of equal size ordered by percentiles) and cross-classified with observed cases to visualize the concentration of risk associated with high composite predicted probabilities. Sensitivity was defined as the proportion of cases found among pre-defined proportions of respondents with the highest predicted probabilities. Positive predictive value (PPV) was defined as the probability of students developing NSSI when estimated among predefined proportions of respondents with the highest predicted probabilities. We used the method of leave-one-out cross-validation to correct for the over-estimation of prediction accuracy when both estimating and evaluating model fit in a single sample [61]. Firth's penalized likelihood estimation was applied to avoid overfitting and inconsistent estimators due to data sparseness [62]. All analyses were performed with SAS (version 9.4) and R (version 3.5.1).

3. Results

3.1. Incidence of NSSI during college

The 12-month incidence of NSSI was estimated at 10.3% (SE = 0.8) in year 1, and 6.0% (SE = 0.7) in year 2. Aggregated rates of onset of NSSI were estimated at 15.6% (SE = 0.9) during the first two college years, with 8.6% (SE = 0.8) reporting sporadic NSSI and 7.0% (SE = 0.6) reporting repetitive NSSI. The three most commonly reported methods were smashing hand or foot against the wall or other objects (52.0%, SE = 3.5), scraping the skin (37.3%, SE = 3.3), and hitting oneself (35.1%, SE = 3.1).

3.2. Bivariate and multivariate risk factors for onset of NSSI

The investigation of different risk factor domains revealed the following key findings at the individual-level. First, the most important sociodemographic and college-related variable that predicted NSSI onset was vocational secondary school track (Table 1). Second, while a variety of traumatic experiences prior to the age of 17 predicted both sporadic and repetitive NSSI (Table 2), in multivariate models only dating violence, emotional abuse, and bully victimization were significant predictors of both forms of NSSI.

Third, in bivariate models, an examination of the temporal associations between 12-month stressful experiences and incident NSSI revealed that several proximal interpersonal stressors (e.g., serious betrayal by someone other than partner) were predictive of sporadic and/or repetitive NSSI (Table 3). However, in multivariate models only unsatisfactory peer support predicted both forms of NSSI. Repetitive NSSI was also significantly associated with unsatisfactory family support, serious ongoing arguments or break-ups, and other stressful events. Fourth, all mental disorders and symptoms of psychopathology were consistently associated with increased risk for sporadic and/or repetitive onset of NSSI, with the only exception being alcohol use disorder (Table 4). In multivariate models, however, only 12-month suicidal ideation and severe role impairment in daily life were uniquely predictive of both forms of NSSI.

Next, we determined the potential population-level impact of the examined risk domains for the onset of NSSI. Assuming a causal association, we estimated that one third of sporadic NSSI (PARP = 32.9%), and nearly one half of repetitive NSSI (PARP = 46.0%), occurring for the first time in college, might have been preventable if it were possible to prevent any childhood-adolescent traumatic experiences. Somewhat smaller PARPs were observed for 12-month stressful experiences for onset of sporadic (PARP = 21.5%)

Table 1
Sociodemographic and college-related variables as baseline predictors for onset of non-suicidal self-injury.

I. Sociodemographic and college-related variables	Prevalence ^a			Sporadic NSSI			Repetitive NSSI				
	w(n)	w(%)	SE	Bivariate model ^b		Multivariate model ^c		Bivariate model ^b		Multivariate model ^c	
				OR (95% CI)	PARP (%)	OR (95% CI)	PARP (%)	OR (95% CI)	PARP (%)		
Sex (male)	1,658	44.1	0.8	1.0 (0.7-1.4)	1.0 (0.7-1.4)	0.5	0.9 (0.6-1.3)	0.9 (0.7-1.3)	–3.3		
Age > 18 years	855	22.8	0.7	1.4 (1.0-2.0)	1.2 (0.8-1.7)	3.4	1.5 (1.0-2.2)	1.2 (0.9-1.8)	3.9		
Non-Belgian nationality	172	4.6	0.3	2.3 (1.2-4.2)	1.9 (1.0-3.7)	3.5	2.5 (1.4-4.6)	1.9 (1.0-3.6)	3.8		
Parents' financial situation difficult	609	16.2	0.6	1.4 (0.9-2.1)	1.2 (0.7-1.9)	2.0	1.8 (1.2-2.7)	1.4 (0.9-2.2)	5.8		
Parental educational level ^d											
Both parents high education	2,358	62.8	0.8	ref	ref	ref	ref	ref	Ref		
One parent high education	817	21.7	0.7	1.1 (0.8-1.6)	1.0 (0.7-1.5)	0.6	1.1 (0.7-1.7)	0.9 (0.6-1.4)	–1.2		
Neither parents high education	582	15.5	0.6	1.2 (0.8-1.9)	1.0 (0.6-1.6)	–0.1	1.5 (1.0-2.3)	1.1 (0.7-1.8)	1.7		
Non-intact family composition ^e	816	21.7	0.7	1.3 (0.9-2.0)	1.2 (0.8-1.8)	2.6	1.6 (1.1-2.3)	1.2 (0.8-1.9)	4.5		
Area of enrolment											
Human Sciences	1,989	52.9	0.8	ref	ref	ref	ref	ref	ref		
Science and Technology	997	26.5	0.7	0.8 (0.6-1.2)	0.9 (0.6-1.3)	–3.0	0.6 (0.4-1.0)	0.7 (0.5-1.1)	–6.8		
Biomedical Sciences	771	20.5	0.7	0.8 (0.5-1.2)	0.8 (0.5-1.2)	–3.8	0.7 (0.5-1.1)	0.7 (0.5-1.2)	–5.2		
Vocational secondary school track	180	4.8	0.4	2.3 (1.1-4.5)	2.0 (1.0-4.0)	3.1	3.1 (1.7-5.6)	2.5 (1.4-4.7)	5.5		

Note: ^a Prevalence estimate of potential risk factors among those without a history of NSSI at baseline, ^b Bivariate associations are based on separate models for each row, with the variable in the row as predictor, ^c Multivariate associations are based on all factors shown in the table, ^d high education level was defined as holding at least a bachelor's degree, ^e defined as parents divorced or separated. w(n) = weighted number of cases, w(%) = weighted percentage of sample, OR = Odds Ratio; PARP = Population Attributable Risk Proportion; Significant odds ratios and PARPs are shown in bold ($\alpha = .05$).

Table 2
Childhood-adolescent traumatic experiences (< 17 years) as baseline predictors for onset of non-suicidal self-injury.

II. Traumatic experiences	Prevalence ^a			Sporadic NSSI			Repetitive NSSI				
	w(n)	w(%)	SE	Bivariate model ^b		Multivariate model ^c		Bivariate model ^b		Multivariate model ^c	
				OR (95% CI)	PARP (%)	OR (95% CI)	PARP (%)	OR (95% CI)	PARP (%)		
Parental psychopathology	1,156	30.8	0.8	1.7 (1.3-2.4)	1.4 (0.9-2.1)	7.5	2.2 (1.6-3.1)	1.6 (1.0-2.7)	13.9		
Physical abuse	162	4.3	0.4	3.1 (1.7-5.7)	1.9 (0.9-4.0)	3.0	4.1 (2.3-7.5)	1.8 (0.9-3.8)	3.1		
Emotional abuse	547	14.6	0.6	2.4 (1.6-3.4)	1.8 (1.0-3.2)	8.2	3.1 (2.1-4.6)	1.9 (1.0-3.6)	11.1		
Sexual abuse	32	0.8	0.2	3.5 (0.8-16.0)	1.7 (0.3-9.7)	0.2	8.1 (2.7-24.5)	2.8 (0.7-11.3)	1.6		
Neglect	222	5.9	0.4	2.4 (1.4-4.0)	1.5 (0.8-3.0)	2.4	3.3 (1.9-5.5)	1.6 (0.8-3.3)	3.3		
Dating violence	151	4.0	0.4	3.3 (1.7-6.3)	2.9 (1.4-6.1)	3.8	6.4 (3.8-11.0)	5.3 (2.7-10.3)	10.1		
Bully victimization	976	26.0	0.8	1.9 (1.3-2.6)	1.6 (1.1-2.5)	11.3	2.1 (1.5-3.0)	1.6 (1.0-2.6)	11.9		
Number of traumatic experiences											
None or exactly one	2,982	79.4	0.7	ref	ref	ref	ref	ref	ref		
Exactly 2	440	11.7	0.6	2.2 (1.5-3.2)	1.0 (0.6-1.9)	1.7	2.5 (1.6-3.9)	0.9 (0.5-1.9)	–0.2		
3 or more	335	8.9	0.5	3.3 (2.1-5.1)	0.7 (0.2-1.9)	–4.6	5.3 (3.5-8.3)	0.8 (0.3-2.3)	–3.3		

Note: ^a Prevalence estimate of potential risk factors among those without a history of NSSI at baseline, ^b Bivariate associations are based on separate models for each row, with the variable in the row as predictor, ^c Multivariate associations are based on all factors shown in the table. w(n) = weighted number of cases, w(%) = weighted percentage of sample, OR = Odds Ratio; PARP = Population Attributable Risk Proportion; Significant odds ratios and PARPs are shown in bold ($\alpha = .05$).

and repetitive (PARP = 34.9%) NSSI. The highest PARPs (sporadic = 37.8%; repetitive = 51.1%), however, were found for mental disorders and symptoms of psychopathology. The single most important risk factors at the population-level were bully victimization for sporadic NSSI (PARP = 11.3%) and unsatisfactory peer support for repetitive NSSI (PARP = 14.0%).

3.3. Evaluation of the accuracy of an integrative risk prediction model for onset of NSSI

Finally, we constructed multivariate models that included all factors in the study to predict NSSI onset. Most risk factors became non-significant in these integrative prediction models (see supplementary Table 1), with the exception of dating violence prior to age 17 (for repetitive NSSI; OR = 3.1) and severe role impairment in daily life (ORs in the 1.8–1.9 range). The generated cumulative risk probabilities showed reasonable-to-good performance for detecting onset of both sporadic and repetitive NSSI (Table 5). Cross-validated sensitivity estimates for different proportions of students at highest predicted risk show that an intervention that, for instance, targets the 10% at highest risk would effectively reach 23.9% (SE = 3.3) of students who report

sporadic, and 36.1% (SE = 3.9) of students who report repetitive NSSI, for the first-time during college. The incidence of NSSI in these subgroups would be 26.8% and 31.7%, respectively.

4. Discussion

This study presents a comprehensive examination of incident NSSI in college students. Three main findings stand out. First, the incidence of NSSI was estimated at 10.3% in year 1 and 6.0% in year 2, with 7.0% reporting onset of repetitive NSSI during the first two years of college. Second, as expected, there was no single stand-out risk factor for NSSI onset. Rather, a broad range of distal and proximal risk factors were prospectively associated with both sporadic and repetitive NSSI onset. Third, our findings show that it is possible to develop risk assessment algorithms, focused on a broad, yet feasible, range of clinically meaningful risk factors, to identify and potentially provide targeted interventions to students at high risk for onset of NSSI during college.

This is the first European study to investigate the incidence of NSSI in emerging adults. Despite evidence that emerging adulthood is a sensitive period for the onset of mental disorders and risky behaviors [2–5], the incidence of NSSI has rarely been

Table 3

Twelve-month stressful experiences and perceived social support as baseline predictors for onset of non-suicidal self-injury.

III. 12-month stressful experiences and social support	Prevalence ^a			Sporadic NSSI			Repetitive NSSI				
	w(n)	w (%)	SE	Bivariate model ^b		Multivariate model ^c		Bivariate model ^b		Multivariate model ^c	
				OR (95% CI)	PARP (%)	OR (95% CI)	PARP (%)	OR (95% CI)	OR (95% CI)	PARP (%)	PARP (%)
Life-threatening illness of a friend or family member	745	19.8	0.8	1.3 (0.9-1.9)	1.2 (0.7-2.0)	3.4	1.3 (0.9-2.0)	1.1 (0.6-1.9)	1.0		
Death of a friend or family member	697	18.5	0.8	1.2 (0.8-1.8)	1.1 (0.7-2.0)	1.2	1.4 (0.9-2.1)	1.4 (0.8-2.4)	5.4		
Breakup with a romantic partner	617	16.4	0.7	1.4 (0.9-2.1)	1.2 (0.7-2.0)	2.0	1.8 (1.2-2.7)	1.5 (0.9-2.4)	6.3		
Romantic partner cheated	123	3.3	0.3	1.9 (0.9-4.0)	1.3 (0.5-3.3)	0.7	2.7 (1.3-6.0)	1.5 (0.6-4.0)	1.8		
Serious betrayal by someone other than partner	338	9.0	0.5	1.8 (1.1-3.0)	1.3 (0.7-2.2)	2.1	2.5 (1.6-4.0)	1.4 (0.8-2.5)	4.1		
Serious ongoing arguments or breakup with friend or family member	449	12.0	0.6	2.1 (1.4-3.2)	1.7 (1.0-2.8)	5.8	2.7 (1.8-4.1)	1.8 (1.0-3.2)	8.3		
Life-threatening accident	45	1.2	0.2	2.4 (0.6-9.5)	1.8 (0.4-9.5)	0.5	5.2 (1.7-15.7)	3.5 (0.8-14.7)	2.4		
Seriously physically assaulted	75	2.0	0.3	3.2 (1.2-8.4)	2.8 (1.0-8.0)	2.3	3.5 (1.2-10.3)	2.8 (0.8-9.4)	2.4		
Sexually assaulted or raped	7	0.2	0.1	6.9 (0.5-93.3)	5.8 (0.4-93.3)	0.4	7.6 (0.5-121.7)	4.9 (0.1-366.2)	0.4		
Serious legal problem	104	2.8	0.3	2.2 (0.9-5.4)	1.8 (0.7-4.8)	1.6	2.7 (1.0-7.1)	2.0 (0.7-5.8)	2.2		
Another stressful event	291	7.7	0.5	1.7 (1.0-2.9)	1.4 (0.7-2.7)	2.2	2.5 (1.5-4.1)	1.9 (1.0-3.7)	6.0		
Number of 12-month stressful experiences											
None or exactly one	2,916	77.6	0.8	ref	ref	ref	ref	ref	ref		
Exactly 2	553	14.7	0.7	1.5 (1.0-2.3)	0.9 (0.5-1.9)	-0.2	1.9 (1.2-3.0)	1.0 (0.5-2.0)	0.2		
3 or more	289	7.7	0.5	2.6 (1.6-4.3)	0.9 (0.3-2.8)	-0.3	4.0 (2.5-6.4)	0.9 (0.3-2.9)	-1.6		
Unsatisfactory family support	638	17.0	0.8	1.7 (1.1-2.5)	1.3 (0.8-2.0)	3.8	2.3 (1.5-3.4)	1.6 (1.1-2.5)	9.2		
Unsatisfactory peer support	1,012	26.9	0.9	1.7 (1.2-2.3)	1.5 (1.1-2.2)	10.0	2.0 (1.4-2.9)	1.7 (1.2-2.6)	14.0		
Unsatisfactory or absent partner support	2,414	64.2	0.9	1.1 (0.8-1.5)	1.0 (0.7-1.4)	0.3	1.2 (0.8-1.8)	1.1 (0.7-1.6)	3.3		

Note: ^a Prevalence estimate of potential risk factors among those without a history of NSSI at baseline, ^b Bivariate associations are based on separate models for each row, with the variable in the row as predictor, ^c Multivariate associations are based on all factors shown in the table, w(n) = weighted number of cases, w(%) = weighted percentage of sample, OR = Odds Ratio; PARP = Population Attributable Risk Proportion; Significant odds ratios and PARPs are shown in bold ($\alpha = .05$).

Table 4

Risk for 12-month mental disorders, 12-month suicidal thoughts and behaviors, and associated impairment as baseline predictors for onset of non-suicidal self-injury.

IV. Mental disorders and other serious mental health symptoms	Prevalence ^a			Sporadic NSSI			Repetitive NSSI				
	w(n)	w(%)	SE	Bivariate model ^b		Multivariate model ^c		Bivariate model ^b		Multivariate model ^c	
				OR (95% CI)	PARP (%)	OR (95% CI)	PARP (%)	OR (95% CI)	OR (95% CI)	PARP (%)	PARP (%)
Major depressive disorder	265	7.1	0.4	3.3 (2.1-5.1)	1.8 (0.9-3.4)	4.0	4.8 (3.0-7.6)	1.6 (0.8-3.0)	4.5		
Generalized anxiety disorder	189	5.0	0.4	2.7 (1.5-4.7)	1.3 (0.6-2.8)	0.5	4.5 (2.7-7.5)	1.7 (0.9-3.5)	4.6		
Panic disorder	44	1.2	0.2	4.1 (1.4-12.0)	2.0 (0.6-7.2)	0.9	6.2 (2.4-15.8)	1.8 (0.6-5.8)	1.2		
Broad Mania	21	0.6	0.1	6.9 (1.3-36.3)	2.3 (0.3-16.9)	0.0	18.2 (5.2-63.9)	4.0 (0.9-18.5)	1.6		
Alcohol use disorder											
Low risk for alcohol use disorder	2,777	73.9	0.7	ref	ref	ref	ref	ref	ref		
Risky or hazardous drinking	889	23.7	0.7	1.2 (0.8-1.7)	1.2 (0.7-1.8)	2.0	1.1 (0.7-1.6)	1.0 (0.6-1.6)	-0.2		
Risk for alcohol dependence	92	2.4	0.3	1.9 (0.8-4.6)	1.4 (0.5-3.9)	0.8	2.2 (0.9-5.1)	1.2 (0.4-3.6)	0.6		
Intermittent explosive disorder item positive	608	16.2	0.6	1.5 (1.0-2.3)	1.2 (0.7-2.0)	1.3	2.1 (1.4-3.1)	1.4 (0.8-2.3)	5.2		
Any eating disorder item positive	335	8.9	0.5	2.0 (1.3-3.2)	1.4 (0.8-2.6)	2.0	3.3 (2.2-5.1)	1.7 (1.0-3.1)	6.9		
Any psychotic item positive	280	7.5	0.8	2.6 (1.4-4.7)	2.0 (1.0-4.0)	5.0	3.1 (1.6-5.9)	2.0 (0.9-4.1)	5.8		
Post-traumatic stress disorder item positive	1,044	27.8	0.7	1.8 (1.3-2.5)	1.2 (0.7-1.9)	2.8	2.7 (1.9-3.8)	1.3 (0.8-2.1)	8.5		
No suicidal thoughts and behaviors	3,605	95.9	0.3	ref	ref	ref	ref	ref	ref		
Suicidal ideation	121	3.2	0.3	3.5 (1.9-6.5)	2.3 (1.1-4.9)	2.7	4.7 (2.6-8.5)	2.6 (1.2-5.5)	4.1		
Suicide plans and/or attempts	31	0.8	0.2	3.6 (1.0-12.8)	1.4 (0.3-5.9)	0.1	7.5 (2.6-21.0)	2.3 (0.6-7.9)	1.3		
Non-suicidal self-injury thoughts	57	1.5	0.2	2.9 (1.2-7.0)	2.1 (0.7-5.7)	1.3	3.1 (1.2-8.1)	1.5 (0.4-5.3)	0.8		
Number of positive screens											
None or exactly one	2,706	72.0	0.8	ref	ref	ref	ref	ref	ref		
Exactly 2	615	16.4	0.7	1.9 (1.3-2.8)	1.1 (0.6-2.1)	4.0	2.5 (1.6-3.9)	1.4 (0.7-2.6)	5.5		
3 or more	436	11.6	0.6	3.6 (2.4-5.4)	1.0 (0.3-2.8)	3.2	6.5 (4.3-9.8)	1.3 (0.4-3.8)	5.1		
Severe role impairment in daily life	277	7.4	0.4	3.3 (2.1-5.3)	2.0 (1.2-3.5)	5.7	5.0 (3.2-7.7)	2.3 (1.3-4.1)	8.6		

Note: ^a Prevalence estimate of potential risk factors among those without a history of NSSI at baseline, ^b Bivariate associations are based on separate models for each row, with the variable in the row as predictor, ^c Multivariate associations are based on all factors shown in the table, w(n) = weighted number of cases, w(%) = weighted percentage of sample, OR = Odds Ratio; PARP = Population Attributable Risk Proportion; Significant odds ratios and PARPs are shown in bold ($\alpha = .05$).

studied outside of adolescence. The reported incidence rates are higher than two earlier American-Canadian estimates [2–4% range; 33,34]. Possible explanations may include geographical or methodological differences (i.e., we used a representative sample and made use of an exhaustive NSSI checklist [63]), cohort effects (i.e., increasing rate of NSSI [64,65]), or a combination of these. On balance, our findings confirm recent work in finding a second NSSI onset peak in emerging adulthood [29], and suggest that - although the majority of students who report onset of NSSI will not meet DSM-5 disorder criteria [9] - a large number of young adults will self-injure for the first time in college. Consistent with

studies that show that the transition to college can be a particularly stressful event [3,66,67], our findings suggest that especially incoming college students are at high risk for onset of NSSI. Interestingly, although our rates of cutting were similar to other studies [34], self-cutting was not among the most frequently reported methods. We speculate that because most individuals in our onset sample report sporadic NSSI, more severe NSSI methods such as self-cutting might be less frequently reported as early methods of NSSI. We found some evidence for this as self-cutting was more prevalent among those who reported repetitive NSSI (sporadic NSSI = 11.8% vs. repetitive NSSI = 29.0%).

Table 5

Concentration of risk for onset of NSSI in different proportions of first year students at highest predicted risk based on the final multivariate risk model.

	Sporadic NSSI		Repetitive NSSI	
	Sensitivity % (SE)	PPV % (SE)	Sensitivity % (SE)	PPV % (SE)
100	100.0 (0.0)	8.6 (0.8)	100.0 (0.0)	7.0 (0.6)
90	94.7 (1.8)	9.8 (0.9)	96.4 (1.6)	8.3 (0.7)
80	89.8 (2.4)	10.5 (1.0)	92.9 (2.2)	9.0 (0.8)
70	84.2 (3.0)	11.3 (1.1)	88.8 (2.8)	9.9 (0.9)
60	78.0 (3.4)	12.3 (1.2)	83.9 (3.1)	11.0 (1.0)
50	70.7 (3.6)	13.6 (1.4)	78.1 (3.5)	12.4 (1.2)
40	62.3 (3.8)	15.2 (1.6)	71.2 (3.9)	14.3 (1.4)
30	52.2 (3.9)	17.4 (2.0)	62.6 (4.0)	17.1 (1.8)
20	40.1 (3.9)	20.8 (2.6)	51.7 (4.2)	21.7 (2.4)
10	23.9 (3.3)	26.8 (4.0)	36.1 (3.9)	31.7 (4.0)

Note: see the multivariate models including all predictors across risk domains in supplementary materials. Model-based AUC values were 0.73 (0.02) for sporadic onset NSSI and 0.79 (0.02) for repetitive onset of NSSI. Cross-validated AUC values were 0.70 (0.03) for sporadic onset and 0.75 (0.02) for repetitive onset of NSSI. Sensitivity = proportion of onset cases found among row% of responders at highest predicted risk, based on cross-validated predicted probabilities. Positive Predictive Value = probability of effectively developing onset when being among row% of responders at highest predicted risk, based on cross-validated predicted probabilities.

With respect to prospective risk factors for first onset NSSI, there are three findings that require brief comment. First, we found evidence that the pathogenic effect of early trauma extends vulnerability for NSSI into emerging adulthood [39]. Previous research has shown that early trauma is associated with neurobiological and psychological changes that impede intrapersonal (e.g., self-critical attribution style) and interpersonal functioning (e.g., relational schemas of mistrust) [68–70]. It is worth mentioning within this context that the effect of neglect and all subtypes of abusive family relationships were attenuated when bullying and dating violence were taken into account, suggesting that the former may increase risk through re-victimization in peer and partner relationships [68,71,72]. Second, findings from this study also highlight the significance of proximal negative interactions for incident NSSI. Consistent with recent work showing the importance of positive peer relationships in mitigating risk for NSSI in emerging adults [35,73], we found that limited peer support was associated with the onset of sporadic and repetitive NSSI for approximately one in ten students who self-injured. Third, supporting the conceptualization of NSSI as a transdiagnostic behavior [12], most mental health problems were prospectively associated with sporadic and repetitive NSSI. Multivariate models suggest that the associated role impairment might partially account for these associations.

A novel and perhaps the most important contribution of our study was the development of an integrative multivariate prediction model that yields reasonable prediction accuracy for detecting students at high risk of beginning NSSI during their academic career. Consistent with recent advances in depression and suicide prevention research [4,74], risk screening at college entrance may provide a unique approach to identify those at risk for *future* NSSI and offer timely intervention. Specifically, by offering evidence-based intervention to the top 10% at greatest risk of NSSI onset, our data suggest that we could theoretically prevent nearly one in four sporadic and two in five repetitive onset cases. This figure would increase to more than half of students who report repetitive NSSI if we target the top 20% at risk, although this would also increase the risk of identifying false positives (i.e., students who would never have self-injured). However, it could also be argued that these students may still benefit from a general mental health promotion intervention because of their constellation of clinically significant risk factors. While these findings are promising, further research will almost certainly be able to

improve these models by including protective factors (e.g., emotion regulatory capability), NSSI-related cognitions (e.g., self-efficacy to resist NSS), and allowing for interactions in larger samples. Building upon these findings, the next logical step would then be to determine how preventive interventions could best be delivered (e.g., making use of the high scalability of internet- and mobile-based applications) and which type of interventions (i.e., trans-diagnostic vs. NSSI-specific) work best for students at varying levels of risk. Addressing these questions in future research will be extremely important to help guide and fully exploit the potential of individualized screening and preventive approaches for NSSI in college. Taken together, the current findings show that effectively dividing college entrants into low and high-risk groups by means of an empirically-derived prediction model has the potential to help optimize the deployment of preventive interventions aimed at reducing the incidence of NSSI and its potentially negative consequences (e.g., increased capability for suicide [75]).

Several limitations deserve attention in interpreting the results of this study. First, response rates in the 54–63% range are sub-optimal. We used state-of-the-art missing data handling techniques to tackle potential residual non-response bias, however, this remains a concern. Second, we used validated clinical screening scales instead of full diagnostic interviews to assess risk for mental disorders; hence, these prevalence rates should be interpreted cautiously. Third, the extent to which the identified risk factors are also causally predictive of NSSI onset cannot be resolved with our current approach. The best way to resolve this uncertainty is to carry out randomized trials that evaluate the effectiveness of targeting the identified risk factors. Finally, because our results are based on data from one college, replicating the findings represents an important goal for future research.

5. Conclusion

The current study makes significant advances to both science and practice by estimating the incidence of NSSI in college students and examining clinically useful prediction models that can identify students at risk for *future* NSSI. Results show that the college years are a sensitive period for the onset of NSSI. While our findings shed light on many risk factors for sporadic and repetitive incident NSSI, effect sizes of individual prospective associations were weak to moderate. Importantly, however, combining risk factors from multiple domains into an integrative prediction model enabled us to detect college entrants at high cumulative risk for incident NSSI with a reasonable degree of accuracy. Further research in this area has the potential to deliver a powerful and cost-beneficial tool that will be valuable in planning future preventive interventions for NSSI in college populations worldwide.

Declaration of interest

In the past 3 years, Dr. Kessler received support for his epidemiological studies from Sanofi Aventis, he was a consultant for Johnson & Johnson Wellness and Prevention, Shire and Takeda, and served on an advisory board for the Johnson & Johnson Services Inc. Lake Nona Life Project. Dr. Kessler is a co-owner of DataStat, Inc., a market research firm that carries out healthcare research. The other authors have no interests to declare.

Acknowledgements

The authors wish to thank the student services of KU Leuven for their assistance in data collection. This research was supported in part by grants from the Research Foundation Flanders [11N0514N (PM), 11N0516N (PM), 1114717N (GK), 1114719N (GK)], King

Baudouin Foundation [2014-J2140150-102905 (RB)], a New Independent Researcher Infrastructure Support Award [Department of Health, Government of Western Australia (MB)] and Curtin University [CIPRS/HSFIRS (GK)]. The funding sources had no role in the design and conduct of the study; collection, management, analysis, interpretation of the data; preparation, review, or approval of the manuscript; and decision to submit the manuscript for publication.

Appendix A. Supplementary data

Supplementary material related to this article can be found, in the online version, at doi:10.1016/j.eurpsy.2019.04.002.

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