GUEST EDITORIAL

Toward fine-grained phenotyping of suicidal behavior: the role of suicidal subtypes

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Despite advances in the assessment and management of suicidal behavior (SB), suicide remains a leading cause of morbidity and mortality in the United States, and suicide rates have increased dramatically over the past 30 years.¹ There is ample evidence that suicidal thoughts and behaviors are transdiagnostic phenomena that can arise even in the absence of other diagnosable mental health disorders.² While attempts to elucidate a unified model of SB have identified genetic, neurobiological, and psychological factors that are associated with increased risk, these risk factors, or their combination, have only modest statistical and limited clinical utility.³ Thus, predicting who will engage in SB remains challenging. We assert that, rather than being a unified construct, SB likely represents a final common pathway of multiple separate pathological processes. Here, we hypothesize that the pattern of suicidal thinking helps distinguish at least two of these suicidal subtypes.

Three lines of evidence support the notion that SB results from multiple suicidal subtypes. First, there is evidence of associations between particular clinical contexts and features of SB and distinct neurobiological underpinnings. For example, limbic circuits have been implicated in unplanned SB in late life depression, and serotonin receptor 1A binding potential in the raphe nuclei has been linked to more lethal SB in depression.^{4,5} Second, contrary to what we would expect with a unified stress-diathesis model, there

appears to be an inconsistent relationship between stressful life events and suicidal behavior.⁶ Third, fine-grained assessments have shown that suicidal thinking varies from fluctuating to persistent, and that fluctuating and persistent thoughts of suicide are associated with different risks of SB.^{7,8}

In light of this, we posit that there are at least two patterns of suicidal thinking: stress-responsive and non-stress-responsive.⁹ We further propose that these patterns have their own risk factors and pathophysiology, and that they moderate or mediate the relationship between environmental factors, such as life stressors and access to means, and the timing and lethality of SB. Finally, we think these patterns fit with existing knowledge about SB while potentially explaining some of the observed inconsistencies and limited overall predictive value of established risk factors (Figure 1).

In the stress-responsive pattern of suicidal thoughts, individuals report sudden, potentially fleeting increases in suicidal thoughts following stressful life events. We think that individuals with this pattern of suicidal thinking are at higher risk of less planned SB and that limiting access to means might be particularly important for them. We hypothesize that the stress-responsive pattern is rooted in childhood trauma (CT). CT is a risk factor for suicidal behavior across psychiatric diagnoses, suggesting it plays an independent role in the etiology of suicidal behaviors.¹⁰ CT is also associated with impulsive aggression—the tendency to react to perceived threats with an intense, emotional, angry response—among adults.¹¹ Indeed, conceptually, impulsive aggression may serve as a model for the kind of adulthood response to stress following CT that we propose and has, in fact, also been associated with SB.¹²

Figure 1. Two proposed suicidal subtypes distinguished by the pattern of suicidal thinking.

Additionally, human and animal studies have identified persistent serotonin system and hypothalamic–pituitary axis dysregulation following CT, and these same systems have been implicated in SB.^{13,14} Moreover, polymorphisms in serotonin- and cortisol-related genes, and also epigenetic modifications to serotonin- and cortisol-related genes, have been specifically linked with SB following CT.¹⁵ However, findings regarding hypothalamic–pituitary axis dysfunction, and especially the relationship between cortisol response to stress and SB, are mixed, with some studies showing blunted cortisol response in attempters compared to non-attempters, and others showing heightened response or no difference.^{16,17} The existence of suicidal subtypes might explain these inconsistent findings: our preliminary data suggest that a potentiated cortisol response to stress is integral to the stress-responsive pattern of suicidal thoughts, but not to the non-stress-responsive pattern.¹⁸

In the non-stress-responsive pattern, individuals are more likely to report persistent thoughts of suicide. We think this pattern is linked to depressive affect and is more likely to result in more carefully planned SB. A variety of studies support these connections. In addition to being one of the criteria for diagnosing major depression, statistical modeling suggests that major depressive episodes play a role in the development of suicidal thinking.^{19,20} Biomarker studies in depression point to a neurobiological basis for more carefully planned SB, with greater intent to die, including an inverse relationship between cerebrospinal fluid serotonin metabolite levels, and SB planning and lethality.²¹ Studies distinguishing less planned and more planned attempts among depressed patients have shown that some risk factors previously linked to SB generally are associated with more planned attempts specifically. These risk factors include more severe and frequent thoughts of suicide, family history of suicide, and greater suicidal intent.²² Finally, individuals with histories of more carefully planned attempts demonstrate better cognitive control than those with prior impulsive attempts, implicating executive function and reward circuits in the differential pathogenesis of these suicidal subtypes.²³

We have described two suicidal subtypes reflecting different patterns of suicidal thinking and stress responsivity, but we suspect there are others. Furthermore, these patterns might overlap or change over time in the same way that psychological and physiological responses to stress alter over the lifespan.²⁴ Much work still needs to be done to further operationalize these terms, and to connect this work with existing knowledge. We advocate for integrated research to further delineate suicidal subtypes.

CONFLICT OF INTEREST

Dr. Bernanke reports no financial relationships with commercial interests. Dr. Stanley receives royalties for the commercial use of the Columbia Suicide Severity Rating Scale. Dr. Oquendo receives royalties for the commercial use of the Columbia Suicide Severity Rating Scale. Her family owns stock in Bristol Myers Squibb.

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