

A Neural Signature for Reappraisal as an Emotion Regulation Strategy: Relationship to Stress-Related Suicidal Ideation and Negative Affect in Major Depression

Sarah Herzog, Noam Schneck, Hanga Galfalvy, Tse Hwei-Choo, Mike Schmidt, Christina A. Michel, M. Elizabeth Sublette, Ainsley Burke, Kevin Ochsner, J. John Mann, Maria A. Oquendo, and Barbara H. Stanley

ABSTRACT

BACKGROUND: Impaired emotion regulation (ER) contributes to major depression and suicidal ideation and behavior. ER is typically studied by explicitly directing participants to regulate, but this may not capture spontaneous tendencies of individuals with depression to engage ER in daily life.

METHODS: In 82 participants with major depressive disorder, we examined the relationship of spontaneous engagement of ER to real-world responses to stress. We used a machine learning–derived neural signature reflecting neural systems that underlie cognitive reappraisal (an ER strategy) to identify reappraisal-related activity while participants recalled negative autobiographical memories under the following conditions: 1) unstructured recall; 2) distanced recall, a form of reappraisal; and 3) immersed recall (comparison condition). Participants also completed a week of ecological momentary assessment measuring daily stressors, suicidal ideation, and negative affect.

RESULTS: Higher reappraisal signature output for the unstructured period, a proxy for the spontaneous tendency to engage ER, was associated with greater increases in suicidal ideation following stressors ($b = 0.083$, $p = .041$). Higher signature output for distanced recall, a proxy for the capacity to engage ER when directed, was associated with lower negative affect following stressors ($b = -0.085$, $p = .029$). Output for the immerse period was not associated with ecological momentary assessment outcomes.

CONCLUSIONS: Findings suggest that in major depressive disorder, the spontaneous tendency to react to negative memories with attempts to reappraise may indicate greater reactivity to negative cues, while intact capacity to use reappraisal when directed may be associated with more adaptive responses to stress. These data have implications for understanding stress-related increases in suicide risk in depression.

<https://doi.org/10.1016/j.bpsc.2024.08.011>

Emotion regulation (ER), the process whereby emotional responses are modulated to meet situational demands and personal goals (1), contributes to psychological well-being. Impaired ER is implicated in mood- and anxiety-related psychiatric disorders including major depression (2,3), and a growing body of literature suggests that suicidal ideation (SI) and behavior are likewise associated with ER deficits (4,5). However, while the nature of ER deficits in depression have been relatively well characterized (6–9), research on suicide-related ER deficits lacks similar granularity. The preponderance of such studies have used self-report measures of global impairments in ER (10), and few empirical studies have characterized the specific nature of ER deficits that contribute to vulnerability toward suicide risk.

Perhaps the most well-researched ER strategy is cognitive reappraisal, which involves changing one's interpretation of the meaning of a stimulus to alter its emotional impact (11). Cognitive reappraisal is effective at downregulating negative

affect (12,13) and diminishes the relationship between mental pain and SI (14). Notably, most behavioral studies of cognitive reappraisal in depressed and suicidal cohorts have utilized structured tasks that explicitly prompt participants when and how to regulate emotions (15). Such tasks were developed for first-generation ER studies to ensure that observed changes in brain activity and behavior were related to engagement of self-regulatory mental processes (16–18). These studies usefully elucidated the behavioral consequences of various ER strategies and their underlying neural mechanisms (19). However, these directed behavioral tasks may not be informative about how ER is engaged in daily life, when it usually occurs in an undirected, spontaneous manner (20), without conscious goals to alter emotional reactions (21). Directed behavioral tasks can gauge an individual's capacity to engage ER when instructed but may not reflect their natural spontaneous tendency to engage ER under ordinary circumstances.

SEE COMMENTARY ON PAGE 5

Meaningful individual differences in the spontaneous tendency to engage ER may be of important clinical relevance to suicidal populations. Depressed participants with SI perform comparably to healthy volunteers when explicitly instructed to regulate (9) but report greater difficulty employing ER (4). This observation suggests that individuals with SI have the capacity to regulate emotions upon instruction in the laboratory but may be less likely to engage these regulatory strategies in daily life (22). Consistent with this interpretation, less frequent use of cognitive reappraisal has been linked to higher levels of past-week SI in psychiatric inpatients with mood disorders (23). Conversely, a tendency toward greater reappraisal is associated with reduced risk of suicidal behavior (24). The literature also suggests that cognitive reappraisal is important to the regulation of emergent suicide risk in the context of stress in individuals with major depressive disorder (MDD). A month-long ecological momentary assessment (EMA) study in patients with MDD showed that greater use of cognitive reappraisal attenuated the relationship between daily stress levels and same-day SI (25). While cognitive reappraisal appears to be useful for mitigating suicide risk, no study has assessed—either behaviorally or neurally—whether the emergence of SI in individuals with MDD is linked to their spontaneous tendency to use cognitive reappraisal.

The aim of the current study was to quantify the tendency to engage an ER strategy of reappraisal in adults with MDD and assess how reappraisal is associated with real-world suicide-related responses to stress. To address this aim, we leveraged a neural signature for reappraisal that we developed and validated in a previous study (26). This signature was derived using a 2-step multivoxel pattern analysis trained on task-based functional magnetic resonance imaging (fMRI) data to identify a pattern of neural activity associated with directed attempts at cognitive reappraisal (26). The neural signature allows for quantification of cognitive reappraisal without the limitations of self-report measures, which are subject to incomplete/biased recall and constrained by level of insight into emotional experience (27), without relying on indirect behavioral measures [e.g., reaction time or decision making, which are assumed to be the products of successful regulation (28)]. Another benefit of the neural signature is that it overcomes the need to explicitly instruct participants to regulate and therefore allows assessment of the natural tendency to engage in reappraisal.

In the current study, the signature was used to detect and quantify engagement of neural mechanisms that supported reappraisal while participants recalled negative autobiographical memories. Reappraisal was quantified 1) under unstructured recall conditions as a proxy for the spontaneous tendency to engage reappraisal, 2) in response to explicit instruction to engage reappraisal using a distancing strategy, and 3) in response to explicit instruction to immerse in the memory. Participants completed a 7-day EMA period in which they reported on stressors, SI, and negative affect up to 6 times daily. Based on previous work that has linked mood disorders to deficits in spontaneous ER (6,29), we hypothesized that signature output during the unstructured recall period would be associated with lower stress-induced increases in SI and negative affect.

METHODS AND MATERIALS

Sample

The sample consisted of 82 participants with major depressive disorder, 33 of whom were part of the training sample in which the neural signature was initially derived [see Schneck *et al.* (26) for a full description of the training sample]. All participants were screened to confirm English reading fluency, normal or corrected-to-normal vision, and absence of conditions contraindicated for MRI. Study procedures were approved by the Institutional Review Board at the New York State Psychiatric Institute.

Clinical Assessment. Psychiatric diagnoses were established using the Structured Clinical Interview for DSM-IV (30), conducted by doctoral- or masters-level psychologists. Depression severity was quantified with the 24-item Hamilton Depression Rating Scale.

Inclusion/Exclusion Criteria. Participants with depression met criteria for a current major depressive episode, were between ages 18 and 65 years, and had been medication free for ≥ 21 days at the time of scan. The medication washout protocol, which was performed as part of participants' participation in a positron emission tomography study of 5-HT_{1A} autoreceptor binding, involved a 1-week medication taper and 3 weeks off any medication that affects the serotonergic system. Exclusion criteria consisted of 1) lifetime psychosis, 2) substance/alcohol abuse (past 2 months) or past-year substance/alcohol dependence, 3) past-year anorexia nervosa or bulimia nervosa, 4) lifetime intravenous drug use, 5) >3 lifetime incidents of MDMA use, 6) first-degree family member with schizophrenia (for participants under age 33), 7) significant active physical illness, 8) electroconvulsive therapy in the past 6 months, and 9) previous head trauma with loss of consciousness or cognitive impairment.

Ecological Momentary Assessment

The EMA period spanned 7 consecutive days during which participants reported on SI, stressors, and negative affect 6 times daily (see Table S1 for prompts) on a personal device. Prompts were presented at random intervals within 2-hour epochs over a 12-hour wake period customized to each participant (31). This ensured that a participant's fixed schedule (e.g., sleep) did not interfere with data collection on a regular basis.

Total scores for EMA SI were computed by summing responses to the 9 SI items within that same epoch, yielding a time-varying (lagged) total SI score. Change in SI at a given time t (e.g., epochs with stressors) was computed as the difference between the SI score at time t and the SI score at the previous epoch ($t - 1$), as long as both observations occurred on the same day. Change in negative affect was calculated in a similar manner. A time-varying stress indicator was also computed to identify epochs with versus without stressors, denoted as "yes/no." SI change following epochs with stressors was regarded as stress-reactive SI.

Neural Signature Development

To identify a neural signature that reflects engagement of reappraisal-related processes, we employed a multistep procedure involving 2 separate fMRI tasks (see Figure 1 for a

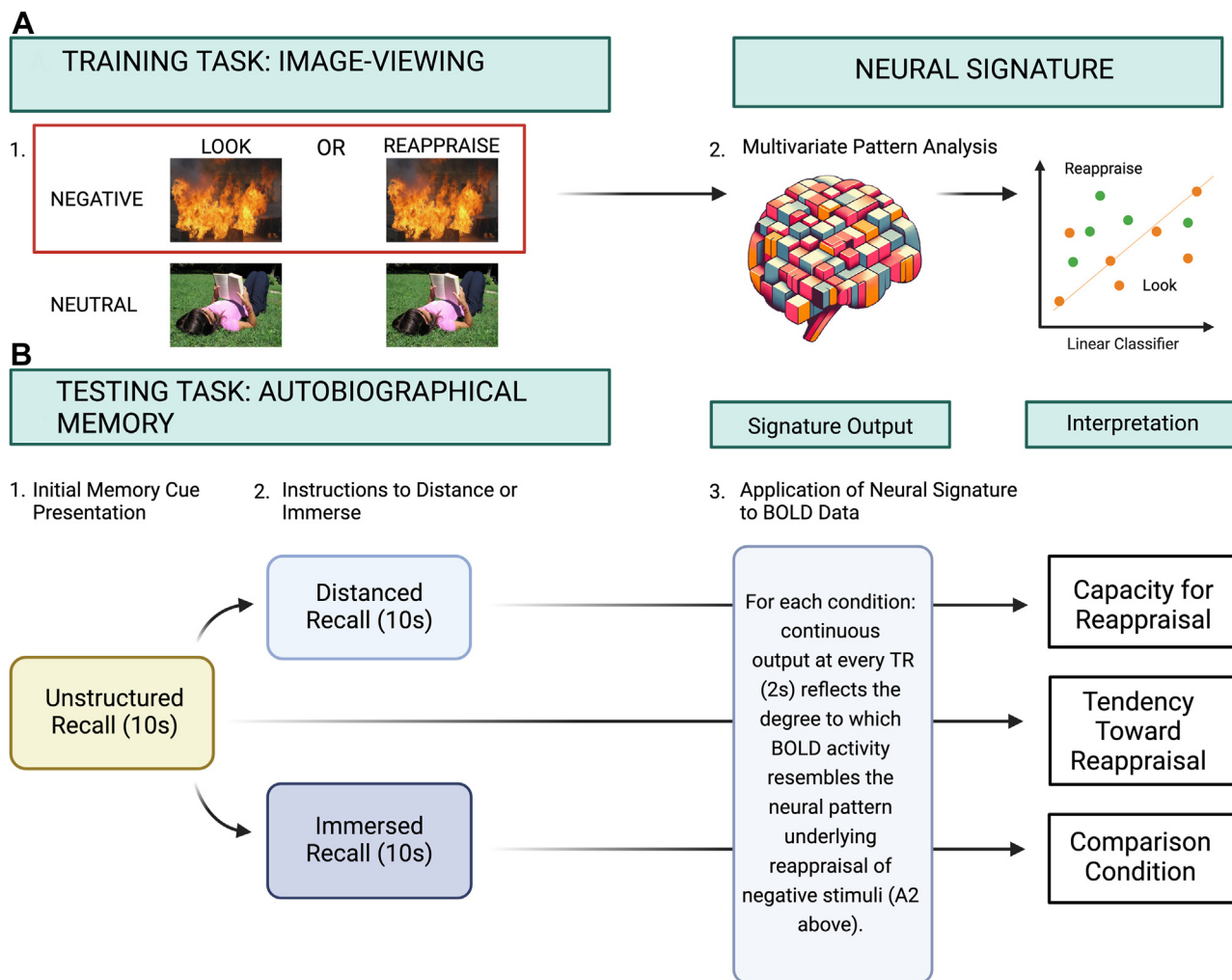


Figure 1. Schematic depiction of the development and application of the neural signature for reappraisal. **(A)** Top: The signature for reappraisal was trained on blood oxygen level–dependent (BOLD) activity from an image-viewing task **(A1)**. A linear classifier **(A2)** was trained to distinguish a pattern of neural activity within a prespecified ventromedial mask that predicts reappraisal vs. look trials for negative images. **(B)** Bottom: Participants completed a negative autobiographical memory task in which they were cued to remember personal memories with brief phrases (e.g., “broke up with boyfriend”). Following a 10-second unstructured recall period **(B1)**, participants were prompted to either distance or immerse while continuing to recall the memory **(B2)**. The neural classifier was applied to BOLD activity from the memory task within the same prespecified ventromedial mask that was used to develop the signature. This yielded continuous output at each 2-second repetition time (TR) reflecting the degree to which BOLD activity resembled the neural pattern underlying reappraisal of negative images **(B3)**. Signature output during the distanced recall period was regarded as a proxy for the capacity to reappraise, while output for the unstructured recall period was regarded as a proxy for the tendency to reappraise.

depiction of signature development). As reported in Schneck *et al.* (26), a linear classifier was trained on fMRI data collected during the course of an image-based reappraisal task (19). The classifier identifies a pattern of spatially distributed neural activity associated with trials for which participants were instructed to reappraise versus a condition where participants were instructed to look at images and respond naturally. This image-based neural signature was validated within the same training sample in a separate testing dataset comprised of fMRI data collected during recall of negative autobiographical memories (task described below). This demonstrated that the signature was associated with reappraisal activity in a separate task and that it was not overfit to the stimuli characteristics of the image-viewing task.

In the current study, the classifier, or neural signature, was applied to a larger sample of participants with MDD who completed both the autobiographical memory task and a week-long EMA of stress, SI, and negative affect.

fMRI Task

Autobiographical Memory Task. In a prescan interview, participants provided 8 personal negative memories of events that had occurred within the last 6 months and generated 2 to 4 words per event to be used as cues to elicit those memories. Before scanning, participants were tested to confirm that they could recall their memories with the cues provided. All trials began with a 10-second presentation of a memory cue, and

this comprised the unstructured recall period. After the initial 10 seconds and a brief jittered interstimulus interval, participants were directed to either distance or immerse while continuing to recall the memory for another 10 seconds. On immerse trials, participants recalled their negative memories from a first-person perspective, allowing recollected events to unfold naturally (“as if re-living the event through your own eyes”). On distance trials, participants recalled their memories from a fact-focused, third-person perspective (“as if watching events unfold from the viewpoint of a camera”). In previous work using this paradigm, distance trials were associated with lower negative affect than the immerse condition (26,32,33), supporting the use of distancing as a reappraisal strategy that can be used to downregulate negative emotion (34–36). See Figure 1 for a depiction of task structure. Participants were trained on the autobiographical memory task prior to scanning.

Each of 8 memories were recalled twice, once for distanced and once for immersed trials, in counterbalanced order across participants. After each distance or immerse trial, participants rated their negative affect and memory vividness. Trials were followed by a 26-second active baseline task in which participants indicated the direction of arrows on the screen to provide a perceptual baseline condition that was not emotion focused and did not involve recall (37). The task was completed in 4 runs of 4 trials, with each trial being approximately 8.5 minutes in total duration.

Image Acquisition. fMRI scans were conducted with small variations in scanning protocols. See the Supplement for details on image sequence parameters and MR preprocessing steps. For all participants, runs began with an 8-second fixation, and the corresponding 4 volumes were discarded. During functional scanning, task stimuli were viewed on an MR-compatible back-projection screen seen in a mirror mounted atop the head coil. Stimuli were presented using E-prime software (Psychology Software Tools, Inc.) on a personal computer. Affect ratings were collected using an MR-compatible 5-button response box.

Application of Neural Signature

The neural signature was applied to the autobiographical memory task data within the same prespecified ventral-frontal mask used to train the signature (see the Supplement for the description and rationale of the mask used). The classifier was applied to 2 distinct periods of all trials: 1) an unstructured 10-second recall period at the beginning of each trial and 2) the subsequent 10 seconds during which participants were directed to use distanced recall or immersed recall (see Figure 1). Application of the neural signature entailed voxelwise multiplication of the weighted vector from the image-based task with values for blood oxygen level-dependent (BOLD) data collected during the autobiographical memory task, followed by a linear summation across voxels. Neural signature output is continuous and reflects the degree to which fMRI activity during the memory task is similar to the voxel pattern associated with reappraisal trials on the image-based task. See the Supplement for a detailed explanation of how the signature was trained and validated.

Statistical Analyses

Auxiliary Analyses. Conditionwise differences in signature output were probed with longitudinal mixed effects linear models. Models featured repetition time (TR)-by-TR signature output as the outcome variable, condition (unstructured/distance/immerse) as a categorical predictor, and participant-specific random intercepts. Models covaried for task run number (1–4), trial number (1–4), and average BOLD signal. A similar model was used to assess the relationship between affect ratings on TR-by-TR signature output and included a main effect for affect rating (continuous) and a condition \times affect rating interaction term.

Data Preparation. Because EMA data are longitudinal (i.e., repeated), neural signature output values were reduced to participant-level averages for models examining associations between signature output and EMA outcome variables. First, to account for the potential influence of fluctuation in overall BOLD signal over time, signature output values were residualized on the average BOLD signal using a generalized least squares regression model featuring an autoregressive of order 1 within-participant correlation structure. Participant-level mean values for the residualized signature output were calculated separately for unstructured recall, distance trials, and immerse trials.

Primary Analyses. Three separate models were used to examine the respective effects of participant-level average neural signature output for unstructured, distanced, and immersed recall on EMA-assessed change in SI following epochs with versus without stressors. The first of those models (unstructured recall) was the main analysis of interest, and the latter two were conducted for purposes of comparison. Longitudinal mixed effects linear regression models featured EMA-assessed SI change as the outcome variable and participant-specific random intercepts. Predictors included neural signature output (continuous) and the time-varying EMA stress indicator (stressor vs. nonstressor epochs) as main effects and an interaction term for signature output \times stress. To model change in ideation, the model included a time-lagged SI total score as a covariate, reflecting average ideation at the previous epoch.

We examined the effect of neural signature output on EMA-assessed change in negative affect in a similar manner to SI. The abovementioned 3 models were replicated while substituting the outcome variable for negative affect change. The time-lagged total SI score was replaced with a time-lagged negative affect score.

All aforementioned models included a binary covariate representing scanning protocol to account for differences in fMRI sequence and participation in the training sample. Standardized beta coefficients are reported for all models.

RESULTS

Sample Characteristics

Participants were recruited as part of a study on biomarkers of suicide risk in depression, and the sample was enriched for suicidal behavior. The sample ($N = 82$) had a mean age of 30.2 years ($SD = 8.9$), was largely female (64.6%), and most

participants had higher education (91.5%). Participants self-identified as Asian (17.1%), Black/African American (22.0%), Hispanic (29.3%), and White (45.1%). Participants were moderately to severely depressed (mean Beck Depression Inventory score 25.6 ± 8.2), and 45.0% of the sample reported a previous suicide attempt. See Table 1 for descriptive statistics for sample demographic and clinical characteristics.

Ecological Momentary Assessment

The mean number of epochs with responses per participant was 31.8 (SD = 10.2), reflecting a completion rate of 75.7%. On average, participants reported experiencing stressors during 45.7% of EMA epochs. The occurrence of stressors (binary) was associated with increases in SI compared with ideation level during the previous epoch ($b = 0.331$, SE = 0.05, $p < .001$). A similar pattern was evident for the impact of stressors on negative affect ($b = 0.720$, SE = 0.05, $p < .001$). See Figures S3 and S4 for a graphical depiction of SI, negative affect, and stress reported across the EMA period.

Autobiographical Memory Task

Mixed linear models with random participant-level intercepts indicated that neural signature output was greater on distance than on immerse trials in the memory task ($b = 0.006$, SE = 0.002, $p < .001$). Output did not differ on distance trials compared with the unstructured recall period ($b = 0.002$, SE = 0.001, $p = .118$) (see Table S2 for descriptive statistics). Participant-level average output on distance and immerse trials were positively correlated (Pearson's $r = 0.265$, $p = .016$). Average output from the unstructured period was not correlated with distanced recall output ($r = -0.127$, $p = .256$) but was negatively correlated with immersed recall output ($r = -0.319$, $p = .003$).

Memory Task Affect Ratings. Participants reported greater negative affect on posttrial affect ratings for immerse trials than for distance trials ($b = 0.671$, SE = 0.02, $p < .001$). Higher negative affect ratings were associated with lower ER signature output ($b = -0.002$, SE = 0.001, $p = .002$) regardless of condition ($b = 0.000$, SE = 0.000, $p = .839$).

Neural Signature Output and EMA

Unstructured Recall. Results of the mixed linear model indicated no main effect for ER signature output for the unstructured recall period on change in EMA SI ($b = 0.040$; SE = 0.12; $p = .736$; 95% CI, -0.196 to 0.276) (Figure 2). There was an interaction between signature output and the EMA stress indicator, wherein greater ER signature output during the unstructured period was associated with greater increases in SI in the context of stressors ($b = 0.083$; SE = 0.04; $p = .041$; 95% CI, 0.003 to 0.162).

There was no main effect of ER signature output during unstructured recall on EMA negative affect change ($b = -0.049$; SE = 0.10; $p = .638$; 95% CI, -0.254 to 0.157) and no interaction for ER signature relationship to negative affect change by presence or absence of EMA stress ($b = 0.019$; SE = 0.04; $p = .625$; 95% CI, -0.056 to 0.093).

Distanced Recall. There was no relationship between ER signature output during distance trials and EMA SI change

Table 1. Demographic and Clinical Characteristics

Variables	Patients With MDD, N = 82
Age, Years	30.2 (8.9)
Sex	
Female	53 (64.6%)
Male	29 (35.4%)
Education—Some Higher Education	75 (91.5%)
Race	
African American/Black	18 (22.0%)
Asian	14 (17.1%)
Multiracial/Unknown	13 (15.8%)
White	37 (45.1%)
Ethnicity, Hispanic	24 (29.3%)
Duration of Current Episode, Weeks	210.8 (276.6)
Psychiatric Medication in the Past 3 Months—Yes	23 (28.0%)
Nonpsychiatric Medication in the Past 3 Months—Yes	32 (39.0%)
Comorbid Psychopathology	
Borderline personality disorder	17 (20.7%)
Other personality disorder	20 (24.4%)
Past substance disorder	25 (30.5%)
Clinical Rating Scale	
Beck Depression Inventory	25.6 (8.2)
24-Item Hamilton Depression Rating Scale	25.0 (7.7)
Suicidal Ideation and Behavior	
Scale for Suicidal Ideation—Past 2 Weeks	6.6 (8.2)
Previous suicide attempt—Yes	37 (45.0%)
Ecological Momentary Assessment	
Epochs with responses	31.8 (10.2)
Epochs with stressors	13.8 (9.8)
Proportion of epochs with stressors	45.7 (27.9)
Suicidal ideation total	7.0 (4.5)
Negative affect total	45.1 (14.1)

Values are presented as mean (SD) or *n* (%).

($b = -0.004$; SE = 0.12; $p = .976$; 95% CI, -0.240 to 0.247) and no interaction between signature output and SI change due to presence or absence of EMA stress ($b = 0.028$; SE = 0.04; $p = .491$; 95% CI, -0.053 to 0.109) (Figure 3).

There was no main effect of signature output during distance trials on EMA negative affect change ($b = -0.163$; SE = 0.11; $p = .127$; 95% CI, -0.373 to 0.047). There was an interaction between signature output and EMA stress on negative affect change such that higher ER signature output was associated with attenuated increases in negative affect in the context of stress ($b = -0.085$; SE = 0.04; $p = .029$; 95% CI, -0.009 to -0.161).

Immersed Recall. There was no main effect of signature output during immerse trials on EMA SI change ($b = -0.122$; SE = 0.13; $p = .340$; 95% CI, -0.376 to 0.131) and no interaction based on presence or absence of EMA stress regarding the relationship between signature output and SI change ($b = 0.016$; SE = 0.06; $p = .764$; 95% CI, -0.091 to 0.124).

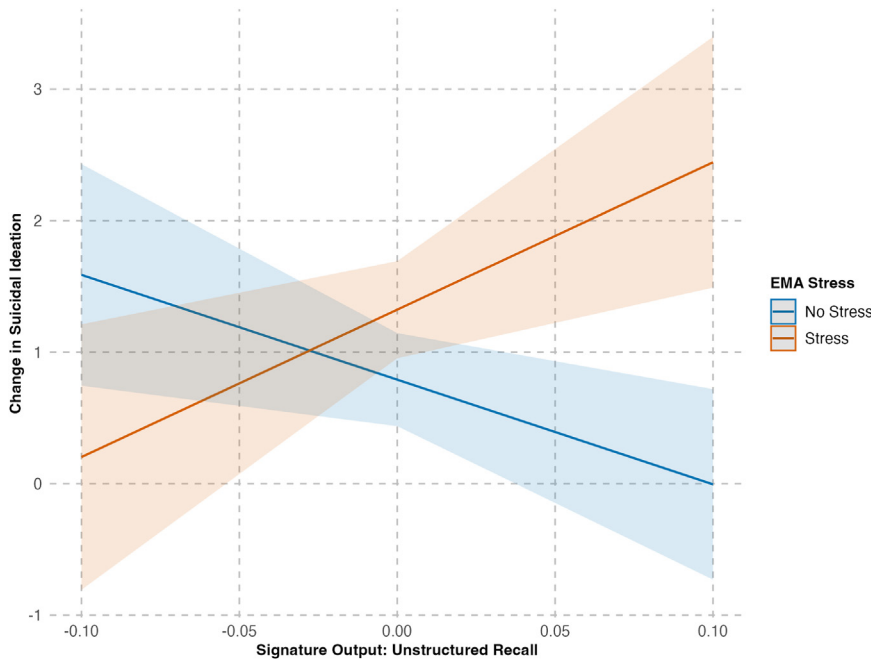


Figure 2. Interaction of signature output for unstructured recall and ecological momentary assessment (EMA) stress change in suicidal ideation. The plot depicts the estimated marginal means of the interaction model. There was a significant interaction between reappraisal signature output for the unstructured recall period and EMA stress on epoch-to-epoch change in suicidal ideation. Greater reappraisal signature output during the unstructured recall period, a proxy for the tendency to engage reappraisal, was associated with greater increases in suicidal ideation following epochs with stressors.

There was also no main effect of signature output during immerse trials on EMA negative affect change ($b = -0.094$; $SE = 0.11$; $p = .403$; 95% CI, -0.316 to 0.128) and no interaction for signature output by EMA stress on negative affect change ($b = 0.013$; $SE = 0.05$; $p = .796$; 95% CI, -0.065 to 0.137).

DISCUSSION

In the current study, we aimed to understand whether engagement of an ER strategy of cognitive reappraisal by individuals with MDD was related to the emergence of SI and negative affect in response to real-world stressors. We found

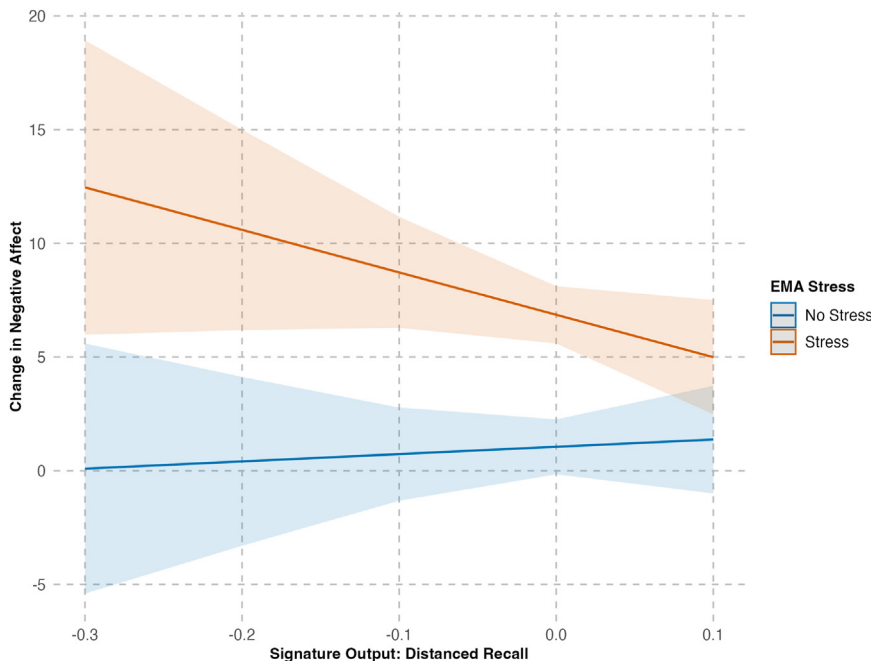


Figure 3. Interaction of signature output for distanced recall and ecological momentary assessment (EMA) stress on change in negative affect. The plot depicts the estimated marginal means of the interaction model. There was a significant interaction between reappraisal signature output for the directed recall period and EMA stress on epoch-to-epoch change in negative affect. Greater reappraisal signature output during the directed reappraisal, a proxy for the capacity to engage reappraisal, was associated with lower negative affect change following epochs with stressors.

that reappraisal neural signature output for the unstructured period was associated with greater increases in SI following stressors and was not associated with negative affect. Conversely, signature output for the distanced recall period was associated with lower negative affect following stressors and was not associated with SI. Thus, the spontaneous tendency to reappraise during exposure to negative memories was related to more pronounced SI in response to stressors, while greater use of reappraisal when directed was associated with less acute negative affective responses to stress. Because the unstructured recall period consisted of the first 10 seconds of exposure to the memory cue, the positive association between signature output during unstructured recall and more acute SI following stressors may imply that stress-sensitive individuals attempt to dampen initial emotional responses to upsetting cues. This could perhaps be due to the difficulty that individuals with MDD have managing negative reactions. Vulnerability to SI in individuals with MDD is associated with nonacceptance of negative emotions (38) and frequent use of ineffective or detrimental emotion regulation strategies (23,39,40). Reduced confidence in the ability to effectively cope with negative emotions may result in attempts to regulate emotional reactions even in instances when the stakes are relatively low, as is the case in the current study, or when regulation may not be advantageous or functional (13,29).

This study is one of the first to neurally quantify both the spontaneous tendency of adults with MDD to engage cognitive reappraisal, as well as the capacity to reappraise when directed. The tendency versus capacity to regulate is a distinction that has only entered the discourse on ER more recently (41,42). The importance of this distinction to clinical outcomes is underscored by evidence suggesting that the tendency to regulate captures individual variation in ER more closely than capacity to regulate (22). In the current study, neural signature output for unstructured and distanced recall periods were not correlated, supporting their distinction as separate constructs. Our findings suggest that the capacity of individuals with MDD to engage reappraisal when instructed may be a positive prognostic indicator of their ability to cope with stress in daily life, while the spontaneous tendency to engage reappraisal under conditions of benign risk (as in the autobiographical memory task) may reflect a general tendency toward experiencing negative cues with a greater sense of threat. That is, greater sensitivity to a relatively benign negative stimulus may provoke both increased efforts at regulation and greater increases in SI in response to daily stress. Relatedly, our finding that spontaneous engagement of ER was associated only with stress-related SI but not negative affect may reflect a general tendency toward engaging maladaptive methods of coping. SI has been understood by clinical theorists as an attempt at coping with psychological pain or problem solving in difficult circumstances (43). This understanding is central to the theoretical approach of prominent evidence-based interventions for suicide risk (44). It is possible that individuals who spontaneously regulate their emotional reactions may be more likely to cope with stress through SI but do not necessarily experience stress with heightened negative emotion.

The distinction between tendency versus capacity for reappraisal may help to bridge seemingly divergent findings in

the literature on ER in depressed populations. Individuals with depression report higher daily stress and negative emotion than individuals without depression (45,46) and exhibit difficulty with the cognitive inhibition of negative emotion (2), but they demonstrate performance comparable to that of healthy individuals on directed tasks of ER (26). There is also evidence to suggest that depression-related deficits in inhibitory control are associated with a lower tendency to use cognitive reappraisal (2). Potentially, individuals with MDD have the capacity to compensate for inhibitory control deficits when reappraising in the laboratory, but such deficits may discourage spontaneous use of reappraisal in daily life. Another explanation for divergence between self-reported difficulties in ER and lab-based measures of ER in depression is that lab-based tasks may not be sufficiently potent (compared to real-world stressors) to distinguish between individuals with MDD and healthy individuals.

We note that the neural signature for reappraisal used here was limited to ventrolateral prefrontal regions involved in general cognitive control (see the Supplement), and therefore the signature may capture processes that are not uniquely involved in reappraisal per se but are related to other forms of cognitive effort governed by this brain region. Some such forms of cognitive effort include selective attention or the effortful selection between competing mental representations. However, previous literature suggests that many cognitive forms of emotion regulation, such as reappraisal, do not depend on processes entirely distinct from those involved in the cognitive control of attention or memory but rather reflect cognitive control exerted in the context of emotional distress (19,47). In addition to reappraisal, another coping mechanism that involves applying cognitive effort in the context of distress is rumination, which is a maladaptive regulation strategy that has been implicated in the maintenance of depression (48). In the current study, signature output for the immerse period, which approximates ruminative activity, was unrelated to EMA-assessed SI and negative affect. This may lend further support to the neural signature as an indicator of reappraisal.

A potential limitation of the current study is that the neural signature was developed using a task wherein participants were explicitly prompted to reappraise, and it can therefore be argued that signature output better reflects neural systems associated with explicit regulation, i.e., consciously engaged or controlled reappraisal. However, whether participants deployed reappraisal during the unstructured memory period in an explicit or implicit manner [i.e., consciously aware of a goal to regulate or without awareness of such a goal (21)] cannot be confirmed from the current data. Spontaneous or implicit attempts at reappraisal can be applied with or without a conscious goal to regulate emotions (34,49) and can be deployed with controlled effort or relatively automatically (21). Additionally, because explicit and implicit regulation exist on a continuum, they are unlikely to engage entirely dissimilar neural systems. In fact, neural systems engaged by explicit and implicit regulation are overlapping and include areas such as the ventrolateral prefrontal cortex, dorsal anterior cingulate cortex, and medial prefrontal cortex (19,50). Similarly, both implicit and explicit regulation, when engaged without direct instruction, involve engagement of the ventromedial prefrontal cortex (51,52). Another potential limitation of this study

Neural Emotion Regulation and Responses to Stress

concerns the lack of affect ratings immediately following the unstructured period. This prevented examination of whether unprompted downregulation of negative affect is associated with real-world responses to stress.

Conclusions

We used an innovative machine learning–based method to identify neural activity underlying reappraisal. Our findings link spontaneous reappraisal to increases in SI following real-world stressors. Results suggest that the spontaneous tendency to reappraise negative memories in individuals with MDD indicates lower tolerance of negative affective cues rather than reflecting adaptive efforts at coping. Future research could apply the signature to the study of real-time temporal patterns in emotional responses to dynamic stimuli to characterize dynamic fluctuation in the process of regulation in depressed populations. The neural signature approach could also be extended to characterize a broader range of emotion regulatory strategies, which could then be applied to the monitoring of medication and psychotherapy treatment outcomes.

ACKNOWLEDGMENTS AND DISCLOSURES

This work was supported in part by the National Institute of Mental Health (Grant Nos. R01 MH109326 [to BHS and MAO], P50 MH090964 [to JM], and K23 MH114021 [to NS]) and by an Early Career Researcher Innovation Grant from the American Foundation for Suicide Prevention (to SH).

JM and MAO receive royalties for commercial use of the Columbia Suicide Severity Rating Scale from the Research Foundation for Mental Hygiene. JM received royalties from Columbia University for the Columbia Pathways App. MAO serves as an adviser to Alkermes, Mind Medicine, Sage Therapeutics, St. George's University, and Fundación Jiménez Díaz. Her family owns stock in Bristol-Myers Squibb. All other authors declare no biomedical financial interests or potential conflicts of interest.

ARTICLE INFORMATION

From the Molecular Imaging and Neuropathology Division, New York State Psychiatric Institute, New York, New York (SH, NS, MS, CAM, MES, AB, JJM, BHS); Department of Psychiatry, Columbia University Vagelos College of Physicians and Surgeons, New York, New York (SH, NS, HG, TH-C, MS, CAM, MES, AB, JJM, BHS); Mental Health Data Science Division, New York State Psychiatric Institute, New York, New York (TH-C); Department of Biostatistics, Mailman School of Public Health, Columbia University, New York, New York (HG); Department of Psychiatry, Columbia University in the City of New York, New York, New York (KO); Department of Psychiatry, Perelman School of Medicine, University of Pennsylvania, Philadelphia, Pennsylvania (MAO); and Department of Radiology, Columbia University Vagelos College of Physicians and Surgeons, New York, New York (JJM). Deceased (BHS).

MAO and BHS are joint senior authors.

Address correspondence to Sarah Herzog, Ph.D., at sarah.herzog@nyspi.columbia.edu.

Received Apr 12, 2024; revised Jul 16, 2024; accepted Aug 6, 2024.

Supplementary material cited in this article is available online at <https://doi.org/10.1016/j.bpsc.2024.08.011>.

REFERENCES

- Thompson RA (1994): Emotion regulation: A theme in search of definition. *Monogr Soc Res Child Dev* 59:25–52.
- Joormann J, Gotlib IH (2010): Emotion regulation in depression: Relation to cognitive inhibition. *Cogn Emot* 24:281–298.
- Gyurak A, Gross JJ, Etkin A (2011): Explicit and implicit emotion regulation: A dual-process framework. *Cogn Emot* 25:400–412.
- Colmenero-Navarrete L, García-Sancho E, Salguero JM (2021): Relationship between emotion regulation and suicide ideation and attempt in adults and adolescents: A systematic review. *Arch Suicide Res* 26:1702–1735.
- Franklin JC, Ribeiro JD, Fox KR, Bentley KH, Kleiman EM, Huang X, et al. (2017): Risk factors for suicidal thoughts and behaviors: A meta-analysis of 50 years of research. *Psychol Bull* 143:187–232.
- Joormann J, Stanton CH (2016): Examining emotion regulation in depression: A review and future directions. *Behav Res Ther* 86:35–49.
- Vanderlind WM, Millgram Y, Baskin-Sommers AR, Clark MS, Joormann J (2020): Understanding positive emotion deficits in depression: From emotion preferences to emotion regulation. *Clin Psychol Rev* 76:101826.
- Visted E, Vøllestad J, Nielsen MB, Schanche E (2018): Emotion regulation in current and remitted depression: A systematic review and meta-analysis. *Front Psychol* 9:756.
- Liu DY, Thompson RJ (2017): Selection and implementation of emotion regulation strategies in major depressive disorder: An integrative review. *Clin Psychol Rev* 57:183–194.
- Turton H, Berry K, Danquah A, Pratt D (2021): The relationship between emotion dysregulation and suicide ideation and behaviour: A systematic review. *J Affect Disord Rep* 5:100136.
- McRae K, Ciesielski B, Gross JJ (2012): Unpacking cognitive reappraisal: Goals, tactics, and outcomes. *Emotion* 12:250–255.
- Troy AS, Shallcross AJ, Brunner A, Friedman R, Jones MC (2018): Cognitive reappraisal and acceptance: Effects on emotion, physiology, and perceived cognitive costs. *Emotion* 18:58–74.
- Gross JJ (1998): Antecedent- and response-focused emotion regulation: Divergent consequences for experience, expression, and physiology. *J Pers Soc Psychol* 74:224–237.
- Amazue LO, Ozor T, Chukwuorji JC, Ifeagwazi CM, Onu DU, Onyedire NG (2019): Mental pain and suicidal ideation in nursing students: The moderating role of emotion regulation. *Cogn Brain Behav* 23:171–191.
- Morawetz C, Bode S, Dementl B, Heekeren HR (2017): The effect of strategies, goals and stimulus material on the neural mechanisms of emotion regulation: A meta-analysis of fMRI studies. *Neurosci Biobehav Rev* 72:111–128.
- Schaefer SM, Jackson DC, Davidson RJ, Aguirre GK, Kimberg DY, Thompson-Schill SL (2002): Modulation of amygdalar activity by the conscious regulation of negative emotion. *J Cogn Neurosci* 14:913–921.
- Ochsner KN, Bunge SA, Gross JJ, Gabrieli JD (2002): Rethinking feelings: An FMRI study of the cognitive regulation of emotion. *J Cogn Neurosci* 14:1215–1229.
- Ochsner KN (2019): From the self to the social regulation of emotion: An evolving psychological and neural model. In: Neta M, Haas IJ, editors. *Emotion in the Mind and Body*. Nebraska Symposium on Motivation, vol 66. Cham, Switzerland: Springer, 43–75.
- Ochsner KN, Silvers JA, Buhle JT (2012): Functional imaging studies of emotion regulation: A synthetic and evolving model of the cognitive control of emotion. *Ann N Y Acad Sci* 1251:E1–E24.
- Koole SL, Rothermund K (2011): “I feel better but I don’t know why”: The psychology of implicit emotion regulation. *Cogn Emot* 25:389–399.
- Braunstein LM, Gross JJ, Ochsner KN (2017): Explicit and implicit emotion regulation: A multi-level framework. *Soc Cogn Affect Neurosci* 12:1545–1557.
- Doré BP, Silvers JA, Ochsner KN (2016): Toward a personalized science of emotion regulation. *Soc Personal Psychol Compass* 10:171–187.
- Forkmann T, Scherer A, Böcker M, Pawelzik M, Guggel S, Glaesmer H (2014): The relation of cognitive reappraisal and expressive suppression to suicidal ideation and suicidal desire. *Suicide Life Threat Behav* 44:524–536.
- Ong E, Thompson C (2019): The importance of coping and emotion regulation in the occurrence of suicidal behavior. *Psychol Rep* 122:1192–1210.
- Franz PJ, Kleiman EM, Nock MK (2021): Reappraisal and suppression each moderate the association between stress and suicidal ideation: Preliminary evidence from a daily diary study. *Cognit Ther Res* 45:1120–1127.
- Schneck N, Herzog S, Lu J, Yttredahl A, Ogden RT, Galfalvy H, et al. (2023): The temporal dynamics of emotion regulation in subjects with

- major depression and healthy control subjects. *Biol Psychiatry* 93:260–267.
27. McMahon TP, Naragon-Gainey K (2020): Ecological validity of trait emotion regulation strategy measures. *Psychol Assess* 32:796–802.
 28. Etkin A, Wager TD (2007): Functional neuroimaging of anxiety: A meta-analysis of emotional processing in PTSD, social anxiety disorder, and specific phobia. *Am J Psychiatry* 164:1476–1488.
 29. Ehring T, Tuschen-Caffier B, Schnulle J, Fischer S, Gross JJ (2010): Emotion regulation and vulnerability to depression: Spontaneous versus instructed use of emotion suppression and reappraisal. *Emotion* 10:563–572.
 30. Spitzer R, Williams J, Gibbon M, First M (1990): *Instruction Manual for the Structured Clinical Interview for the DSM-IV (SCID-P)*. Washington: American Psychiatric Press.
 31. Chaudhury SR, Galfalvy H, Biggs E, Choo TH, Mann JJ, Stanley B (2017): Affect in response to stressors and coping strategies: An ecological momentary assessment study of borderline personality disorder. *Borderline Personal Disord Emot Dysregul* 4:8.
 32. Doré BP, Rodrik O, Boccagno C, Hubbard A, Weber J, Stanley B, *et al.* (2018): Negative autobiographical memory in depression reflects elevated amygdala-hippocampal reactivity and hippocampally associated emotion regulation. *Biol Psychiatry Cogn Neurosci Neuroimaging* 3:358–366.
 33. Silvers JA, Hubbard AD, Chaudhury S, Biggs E, Shu J, Grunebaum MF, *et al.* (2016): Suicide attempters with borderline personality disorder show differential orbitofrontal and parietal recruitment when reflecting on aversive memories. *J Psychiatr Res* 81:71–78.
 34. Denny BT, Ochsner KN (2014): Behavioral effects of longitudinal training in cognitive reappraisal. *Emotion* 14:425–433.
 35. Koenigsberg HW, Fan J, Ochsner KN, Liu X, Guise K, Pizzarello S, *et al.* (2010): Neural correlates of using distancing to regulate emotional responses to social situations. *Neuropsychologia* 48:1813–1822.
 36. Powers JP, LaBar KS (2019): Regulating emotion through distancing: A taxonomy, neurocognitive model, and supporting meta-analysis. *Neurosci Biobehav Rev* 96:155–173.
 37. Stark CEL, Squire LR (2001): When zero is not zero: The problem of ambiguous baseline conditions in fMRI. *Proc Natl Acad Sci U S A* 98:12760–12766.
 38. Weinberg A, Klonsky ED (2009): Measurement of emotion dysregulation in adolescents. *Psychol Assess* 21:616–621.
 39. Flores-Kanter PE, Garcia-Batista ZE, Moretti LS, Medrano LA (2019): Towards an explanatory model of suicidal ideation: The effects of cognitive emotional regulation strategies, affectivity and hopelessness. *Span J Psychol* 22:E43.
 40. Azadi S, Khosravani V, King S, Mohammadzadeh A, Baseri A (2020): Effects of neuropsychological systems on psychopathology through cognitive emotion regulation strategies in individuals with suicide attempts. *Cogn Ther Res* 44:229–239.
 41. Silvers JA, Guassi Moreira JF (2019): Capacity and tendency: A neuroscientific framework for the study of emotion regulation. *Neurosci Lett* 693:35–39.
 42. Shu J, Ochsner KN, Phelps EA (2022): Trait intolerance of uncertainty is associated with decreased reappraisal capacity and increased suppression tendency. *Affect Sci* 3:528–538.
 43. Jobes DA (2009): The CAMS approach to suicide risk: Philosophy and clinical procedures. *Suicidologi* 14.
 44. Jobes DA, Drozd JF (2004): The CAMS approach to working with suicidal patients. *J Contemp Psychotherapy* 34:73–85.
 45. Dunkley DA, Ma D, Lee IA, Preacher KJ, Zuroff DC (2014): Advancing complex explanatory conceptualizations of daily negative and positive affect: Trigger and maintenance coping action patterns [published correction appears in *J Couns Psychol* 2014;61:263]. *J Couns Psychol* 61:93–109.
 46. Bylsma LM, Taylor-Clift A, Rottenberg J (2011): Emotional reactivity to daily events in major and minor depression. *J Abnorm Psychol* 120:155–167.
 47. Ochsner KN, Gross JJ (2005): The cognitive control of emotion. *Trends Cogn Sci* 9:242–249.
 48. Nolen-Hoeksema S (2000): The role of rumination in depressive disorders and mixed anxiety/depressive symptoms. *J Abnorm Psychol* 109:504–511.
 49. Denny BT, Inhoff MC, Zerubavel N, Davachi L, Ochsner KN (2015): Getting over it: Long-lasting effects of emotion regulation on amygdala response. *Psychol Sci* 26:1377–1388.
 50. Buhle JT, Silvers JA, Wager TD, Lopez R, Onyemekwu C, Kober H, *et al.* (2014): Cognitive reappraisal of emotion: A meta-analysis of human neuroimaging studies. *Cereb Cortex* 24:2981–2990.
 51. Wager TD, Rilling JK, Smith EE, Sokolik A, Casey KL, Davidson RJ, *et al.* (2004): Placebo-induced changes in FMRI in the anticipation and experience of pain. *Science* 303:1162–1167.
 52. Motzkin JC, Philippi CL, Wolf RC, Baskaya MK, Koenigs M (2015): Ventromedial prefrontal cortex is critical for the regulation of amygdala activity in humans. *Biol Psychiatry* 77:276–284.