

Self-referential processing in adolescents: Stability of behavioral and ERP markers

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Abstract

The self-referential encoding task (SRET)—an implicit measure of self-schema—has been used widely to probe cognitive biases associated with depression, including among adolescents. However, research testing the stability of behavioral and electrocortical effects is sparse. Therefore, the current study sought to evaluate the stability of behavioral markers and ERPs elicited from the SRET over time in healthy, female adolescents ($n = 31$). At baseline, participants were administered a diagnostic interview and a self-report measure of depression severity. In addition, they completed the SRET while 128-channel ERP data were recorded to examine early (P1) and late (late positive potential [LPP]) ERPs. Three months later, participants were readministered the depression self-report measure and the SRET in conjunction with ERPs. Results revealed that healthy adolescents endorsed, recalled, and recognized more positive and fewer negative words at each assessment, and these effects were stable over time ($r_s = .44-.83$). Similarly, they reported a faster reaction time when endorsing self-relevant positive words, as opposed to negative words, at both the initial and follow-up assessment ($r = .82$). Second, ERP responses, specifically potentiated P1 and late LPP positivity to positive versus negative words, were consistent over time ($r_s = .56-.83$), and the internal reliability of ERPs were robust at each time point ($r_s = .52-.80$). As a whole, these medium-to-large effects suggest that the SRET is a reliable behavioral and neural probe of self-referential processing.

Descriptors: Self-referential processing, Depression, P1, LPP, Reliability

Depression is a leading cause of disability worldwide (Bromet et al., 2011; Kessler, 2012), and deeply entrenched cognitive biases—particularly depressogenic self-referential processing biases (i.e., the tendency to attribute negative information as being self-relevant)—play a prominent role in the onset and maintenance of major depressive disorder (MDD) in children and adolescents (Auerbach, Stanton, Proudfit, & Pizzagalli, 2015; Goldstein, Hayden, & Klein, 2015; Jaenicke et al., 1987). Over the past three decades, the self-referential encoding task (SRET; Kuiper & Derry,

1982) has been used to probe cognitive biases implicated in MDD. During the SRET, individuals indicate whether a series of positive and negative adjectives, typically matched across relevant conditions (i.e., arousal, word length), describe themselves. The SRET is conceptualized as an implicit test of self-schema (Goldstein et al., 2015; Kuiper & Derry, 1982), and the resulting negative biases, operationalized as greater endorsement and recall of negative versus positive self-relevant adjectives, are moderately to strongly associated with core cognitive vulnerabilities linked to depression, including self-criticism, rumination, and dysfunctional attribution styles (Alloy, Abramson, Murray, Whitehouse, & Hogan, 1997; Auerbach et al., 2015; Hayden et al., 2013; Joormann, Dkane, & Gotlib, 2006).

Among adults tested with the SRET, depressed individuals endorse and recall a greater number of negative as compared to positive adjectives relative to nondepressed individuals (Derry & Kuiper, 1981; Dobson & Shaw, 1987; Matt, Vázquez, & Campbell, 1992; Moulds, Kandris, & Williams, 2007). Findings in youth are largely consistent; namely, depressed children and adolescents endorse and recall a greater number of negative words relative to positive words (Auerbach et al., 2015; Connolly, Abramson, & Alloy, 2016; Timbremont & Braet, 2004; Zupan, Hammen, &

RPA was supported through funding from the National Institute of Mental Health (NIMH) K23MH097786, the Jewett Fund, and the Tommy Fuss Fund. CAW was funded from the NIMH (1F32MH099810-01; K23MH108752-01). DAP was partially supported by grants from NIMH (1R01MH101521, 3R01MH068376). The content is solely the responsibility of the authors and does not necessarily represent the official views of the National Institutes of Health or NIMH. Over the past 3 years, Dr. Pizzagalli has received consulting fees from Akili Interactive Labs, BlackThorn Therapeutics, Otsuka America Pharmaceutical, and Pfizer for activities unrelated to the current research. No other authors report any conflicts of interest.

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Jaenicke, 1987), and, additionally, they respond faster when endorsing negative words but exhibit a slower reaction time (RT) when endorsing positive words (Auerbach et al., 2015). These depressotypic self-referential processing biases also emerged in remitted depressed children and adolescents (Timbremont & Braet, 2004) as well as at-risk children (i.e., owing to parental history of depression; Jaenicke et al., 1987). Further, in children aged 6–9 years, SRET effects demonstrated modest stability over time (Goldstein et al., 2015). Taken together, these findings suggest that the SRET provides a useful measurement of depressogenic self-referential processing biases.

More recently, research has leveraged scalp-recorded ERPs, which provide excellent temporal resolution in the milliseconds range, to better understand putative processes associated with depressotypic self-referential processing elicited during the SRET (Auerbach et al., 2015; Shestyuk & Deldin, 2010). ERP studies with the SRET have focused on early and late ERP components. Early ERP components, including the P1 and P2, are maximal over parietal-occipital areas, are stable, reflect semantic monitoring of emotional information, and are modulated by word valence (Flor, Knost, & Birbaumer, 1997; West & Holcomb, 2000). Conversely, the late positive potential (LPP) spans several hundred milliseconds to seconds and indexes sustained engagement to both emotional words (Fischler & Bradley, 2006) and images (Foti, Hajcak, & Dien, 2009). Prior work probing the LPP using an emotion-based paradigm has demonstrated strong test-retest reliability over a 2-year period (Kujawa, Klein, & Proudfit, 2013); however, to date, the stability of the LPP within an SRET context has not been tested. During the SRET, the LPP is initially maximal over parietal sites (i.e., early LPP), and later in the temporal course, it propagates to frontocentral regions (i.e., late LPP). This frontal propagation is particularly important in light of prior neuroimaging evidence implicating prefrontal cortex abnormalities in negative self-referential processing (Auerbach et al., 2015; Lemogne et al., 2010).

ERP studies using the SRET have revealed promising findings. Shestyuk and Deldin (2010) found that depressed adults exhibited enhanced P2 and late LPP positivity to negative versus positive words, whereas healthy individuals showed the opposite effect. Our group recently published similar findings in depressed youth (Auerbach et al., 2015). This study reported that depressed youth displayed a potentiated P1, but not P2, in response to negative as compared to positive words, whereas healthy adolescents showed the opposite pattern. Interestingly, greater P1 positivity to negative words was associated with depressotypic cognitive vulnerability factors, including greater self-criticism and a more negative self-view. Results among depressed adolescents also indicated enhanced early (parietal-occipital sites) and late (frontocentral sites) LPP positivity to negative versus positive words, and, again, healthy adolescents showed the opposite effect. Building on these findings in a sample of healthy low- and high-risk youth (owing to a maternal history of depression) aged 8–14 years, Speed and colleagues demonstrated that high-risk youth exhibited a potentiated LPP response to negative words; no between-group differences emerged following positive words (Speed, Nelson, Auerbach, Klein, & Hajcak, 2016). Collectively, these findings suggest that ERPs elicited through the SRET may differentiate healthy and depressed individuals, and further LPP positivity to negative words may be a trait marker that precedes depression onset.

Through its Strategic Plan for Research, the National Institute of Mental Health (NIMH) has outlined the importance of identifying “clinically useful biomarkers and behavioral indicators that predict change across the trajectory of illness” (Strategic Objective 2.2).

Toward this goal, the current study evaluated the stability of behavioral and neural (ERP) indices of the SRET over time in healthy, female adolescents. This is a critical initial step, as any proposed biomarker or behavioral indicator must demonstrate stability over time in healthy populations, prior to being used as a predictor or indicator of mental illness. To this end, we retested healthy female participants evaluated in our recent ERP study in adolescent depression (Auerbach et al., 2015) 3 months after the initial session (and recruited seven additional participants to increase sample size). We tested the following a priori hypotheses. First, we expected to confirm the behavioral and ERP task effects reported in Auerbach et al. (2015) in this extended sample; specifically, we expected that healthy adolescents will (a) endorse, recall, and recognize more positive as opposed to negative words, (b) exhibit a faster reaction to endorse self-relevant positive as opposed to negative words, and (c) show greater P1 and LPP amplitudes in response to positive versus negative words. Second, we hypothesized that these behavioral and ERP effects will remain stable and consistent at the 3-month follow-up assessment.

Method

Procedure

A full description of the procedure was provided in a recent paper, in which female adolescents with MDD and healthy females were compared at a baseline session (Auerbach et al., 2015). Briefly, the Partners Institutional Review Board approved the study. Youth aged 13 to 17 years provided assent, while 18-year-old participants and legal guardians provided written consent. The research project included three study visits. On the first visit, adolescents completed a semistructured diagnostic interview of current and past mental illness and were administered a self-report instrument assessing depressive symptoms. During the second study visit, which occurred within 1–2 weeks of the first study visit, participants completed the SRET while ERP data were recorded. The average length between the first and second visits was 7.87 ± 6.04 days. At the third study visit (i.e., the follow-up assessment), which occurred 3 months later, participants were administered the same depression self-report measure and SRET task in conjunction with ERP. Participants were remunerated \$70 for their participation.

Participants

The sample included 37 healthy female adolescents (30 healthy females included in Auerbach et al., 2015, plus seven additional participants). Participants were aged 13–18 years and recruited from the greater Boston area through online advertisements, posted flyers, and direct mailing. To meet inclusion criteria, participants were required to be fluent in English, right-handed, and female. Exclusion criteria included lifetime diagnosis of any psychopathology, mental retardation, organic brain syndrome, head injury resulting in loss of consciousness for 5 min or seizures, and use of psychiatric medication. One participant was excluded for poor data quality during the initial EEG assessment, and five of the original participants did not complete the 3-month follow-up EEG assessment. Compared to participants lost due to poor data quality and attrition ($n = 6$), the final adolescent sample ($n = 31$) did not differ in age, $t(35) = -0.22, p = .83$, or race, $\chi^2(2) = 1.54, p = .46$; they did differ in socioeconomic status (SES), $\chi^2(3) = 16.75, p = .001$, with the adolescents lost to attrition reporting a higher SES. The final sample of 31 female adolescents ($M = 15.16, SD = 1.5$) included 87.1% White, 6.5% Asian, and 6.5% multiple races. The income distribution included the following: 74.2% = more

than \$100,000, 6.5% = \$50,000 to \$75,000, and 19.4% = not reported.

Instruments

Schedule for Affective Disorders and Schizophrenia for School-Age Children—Present and Lifetime Version (K-SADS-PL). The K-SADS-PL (Kaufman et al., 1997) is a semi-structured clinical interview used to assess current and past psychiatric disorders according to the DSM-IV-TR (American Psychiatric Association, 2000), and past research has demonstrated excellent reliability and validity (Kaufman et al., 1997). Graduate students and bachelor's-level research assistants administered the clinical interview after receiving 40 h of training, which included didactics, mock interviews, and direct supervision. The principal investigator (RPA) reviewed digital audio files of 20% of the interviews selected at random to assess interrater reliability, and the Cohen's kappa coefficients were excellent ($\kappa = 1.00$).

Beck Depression Inventory II (BDI-II). The BDI-II (Beck, Steer, & Brown, 1996) is a 21-item self-report questionnaire that assesses depressive symptom severity over the previous 2 weeks. Items range from 0 to 3, and higher scores indicate higher levels of depressive symptoms. Cutoffs for the BDI-II include 0 to 13 = no or minimal depression, 14 to 19 = mild depression, 20 to 28 = moderate depression, and 29 to 63 = severe depression. In the current study, the Cronbach's alpha for the BDI-II ranged from 0.87 to 0.91, suggesting strong internal consistency.

Experimental Task

The SRET included 80 trials consisting of 40 positive and 40 negative adjectives (see Auerbach et al., 2015). Adjectives were selected from the Affective Norms for English Words based on criteria including valence, arousal, frequency, and length (Bradley & Lang, 2010). Positive and negative adjectives were significantly different in valence, $t(79) = -55.88, p < .001$, but not arousal, $t(79) = 0.68, p = .50$, frequency, $t(79) = -1.64, p = .11$, or word length, $t(79) = -0.06, p = .95$. Stimuli were presented in a pseudorandom order, with no more than two words of the same valence presented in a row. Consistent with past research (Auerbach et al., 2015; Sheshtyuk & Deldin, 2010), in each trial, the stimulus was presented for 200 ms, followed by a fixation cross (1,800 ms) and a question prompt, "Does this word describe you?" Participants responded by pressing "yes" or "no" on a button box. Intertrial intervals were jittered between 1,500 ms and 1,700 ms. Participants completed three practice trials using affectively neutral words prior to the start of data collection. After completing the 80 trials, participants were given a distractor task, consisting of counting backward from 50. Upon completing this distractor task, participants were asked to recall as many words as they could that were presented during the task. Following the recall component, participants were given a recognition task that included 160 words—80 words that appeared in the task and 80 matched distractors (i.e., an additional 40 positive and 40 negative words). In line with prior research (e.g., Goldstein et al., 2015; Prieto, Cole, & Taveson, 1992), we also created a processing bias score for positive and negative words. The positive processing bias score was calculated by dividing the number of positive words endorsed that also were recalled by the total number positive and negative words endorsed. Similarly, the negative processing bias represented the number of negative words endorsed that were recalled divided by the sum of the total number of words endorsed.

Table 1. Behavioral Data from the Self-Referential Encoding Task

Task	Initial (n = 31)		Follow-up (n = 31)	
	Mean	SD	Mean	SD
Endorse				
Positive	30.16	5.09	30.10	6.22
Negative	2.19	2.92	2.71	3.48
Reaction time (ms)				
Positive	518.64	212.01	482.03	221.03
Negative	1,221.75	1,315.41	1,065.18	1,930.94
Recall				
Positive	9.52	4.08	11.58	3.38
Negative	7.68	3.48	8.90	3.83
Recognition				
Positive	36.00	3.73	35.77	4.08
Negative	33.45	4.38	33.19	4.48
Processing				
Positive	0.22	0.09	0.27	0.08
Negative	0.03	0.04	0.03	0.04

EEG Recording, Data Reduction, and Analysis

The EEG data were recorded using a 128-channel net from Hydro-Cel GSN (Electrical Geodesics, Inc., EGI). Continuous EEG data were sampled at 250 Hz and referenced to Cz. Electrode impedances were kept below 50–75 k Ω , and offline analyses were performed using BrainVision Analyzer 2.04 software (Brain Products, Germany). EEG data were rereferenced to the average reference, and offline filters (0.1–30 Hz) were applied. Vertical and horizontal eye movement artifacts were identified and removed using an independent component analysis transform. For each trial, EEG data were segmented 200 ms before and 1,200 ms after stimulus onset. A semiautomated procedure to reject intervals for individual channels used the following criteria: (a) a voltage step $> 50 \mu\text{V}$ between sample rates, (b) a voltage difference $> 300 \mu\text{V}$ within a trial, and (c) a maximum voltage difference of $< 0.50 \mu\text{V}$ within a 100-ms interval. All trials were visually inspected for manual artifact identification and removal.

ERPs were computed time-locked to all available positive and negative words, and the average amplitude 200 ms prestimulus (i.e., word presentation) served as the baseline. ERP amplitudes were examined at sensor locations equivalent to selected electrodes in the 10/10 system. Scalp location and time windows were consistent with previously published findings using a subset of participants from the current study (Auerbach et al., 2015). The P1 and early LPP components were calculated as the mean area across electrode sites Pz, P1, PO3, POz, PO4, and P2 for the following time windows where the component was maximal: (a) P1 = 108–172 ms, and (b) early LPP = 400–600 ms poststimulus. The late LPP was examined across the average of frontocentral midline electrode sites Fz, FCz, and Cz 600–1,200 ms poststimulus.¹

All analyses were conducted with SPSS (version 20.0). A repeated measures analysis of variance (RMANOVA) tested main

1. Whereas some research has shown that the late LPP is maximal over frontocentral regions (Auerbach et al., 2015), other studies have shown that the LPP is maximal over parietal-occipital midline electrodes (e.g., Kujawa et al., 2013). In the current study, the late LPP was maximal over frontocentral regions 600–1,200 ms poststimulus (see Figure 4A/B). Nonetheless, to better integrate with prior LPP research, we also probed the late LPP effect averaged across electrode sites Pz, P1, PO3, POz, PO4, and P2. The main effect for condition was not significant, $F(1,30) = 0.41, p = .53, \eta^2 = .53$.

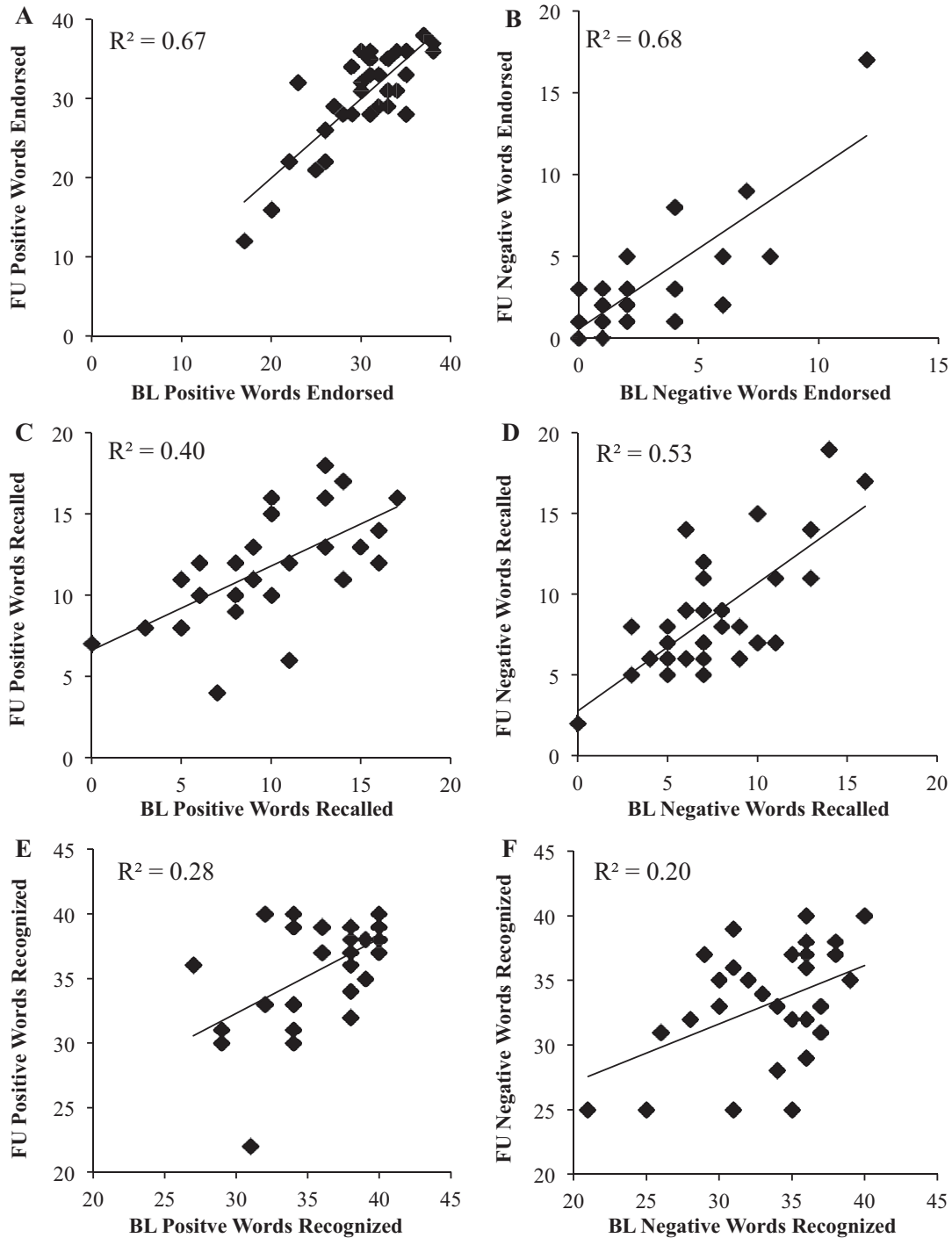


Figure 1. Test-retest for behavioral indices at the baseline and follow-up assessment. BL = baseline; FU = follow-up.

effects for time (Time 1, Time 2) and condition (positive words, negative words) as well as the Time \times Condition interaction. To demonstrate the stability of a given effect over time, we anticipated a significant main effect for condition; neither the main effect for time nor the Time \times Condition interaction was expected to be significant. All analyses included effect sizes (η^2) where .02–.12 = small, .13–.25 = medium, and $\geq .26$ = large. Test-retest reliability for behavioral and ERP indices was evaluated by performing Pearson correlations. The internal consistency of our

ERP indices was computed by examining the correlation of the odd and even trials at each time point.

Results

Descriptive Statistics

Depressive symptoms were assessed at the initial and follow-up assessments (test-retest $r = .75, p < .001$). As expected, depressive

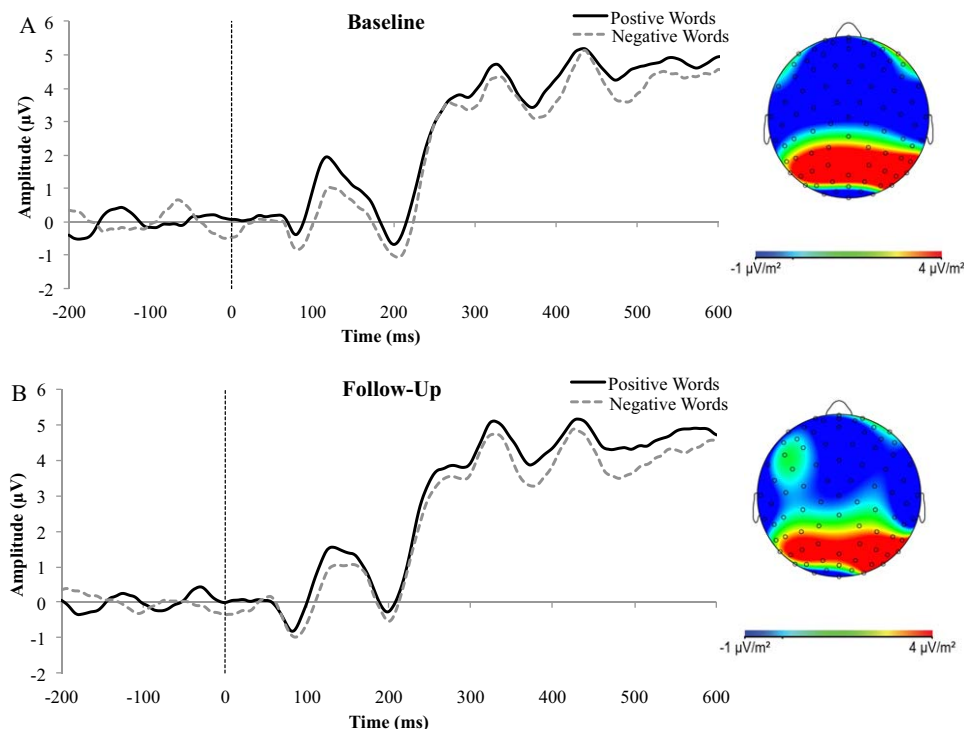


Figure 2. P1 and early LPP responses at the baseline and follow-up assessment. P1 (108–172 ms poststimulus) and early LPP (400–600 ms poststimulus) activity in response to positive and negative words averaged across P1, P2, Pz, POz, PO3, and PO4 for healthy, female adolescents ($n = 31$) during the (A) initial, and (B) follow-up assessment. Scalp topographies reflect the average topography of positive and negative words 108–172 ms poststimulus.

symptom scores were low and in the nondepressed range at baseline ($M = 1.84$, $SD = 3.72$) and follow-up ($M = 4.09$, $SD = 5.71$). While there was a significant difference in symptom scores across assessments, $t(30) = -3.26$, $p = .003$, symptom levels are indicative of healthy adolescents, and no adolescent reported clinically significant depressive symptoms at either of the assessments.

Behavioral Data

Behavioral data from the SRET are summarized in Table 1, and previously reported behavioral effects from the initial assessment (Auerbach et al., 2015) were replicated in this larger sample.

Word endorsement. As hypothesized, a main effect of condition emerged for words endorsed, $F(1,30) = 429.35$, $p < .001$, $\eta^2 = .94$, whereby participants endorsed more positive than negative words as self-relevant. Neither the main effect of time, $F(1,30) = 0.44$, $p = .51$, $\eta^2 = .02$, nor the Time \times Condition interaction, $F(1,30) = 0.55$, $p = .47$, $\eta^2 = .02$, were significant. Additionally, test-retest correlational analyses revealed associations over time for positive ($r = .82$, $p < .001$) and negative ($r = .83$, $p < .001$) words endorsed (see Figure 1A,B).

Reaction time. All participants endorsed positive words. However, 12 adolescents did not endorse any negative words as being self-relevant, and thus these individuals were excluded from the RMANOVA. A main effect of condition emerged for reaction time, $F(1,18) = 6.06$, $p = .02$, $\eta^2 = .25$, indicating that participants were overall faster to endorse positive words compared to negative words. There was no significant main effect for time, $F(1,18) = 0.15$, $p = .70$, $\eta^2 = .01$, or Time \times Condition interaction,

$F(1,18) = 0.06$, $p = .80$, $\eta^2 = .003$. There was a significant test-retest correlation for RT to positive words ($r = .82$, $p < .001$), which included all participants. The test-retest correlation for negative words was not significant ($r = .18$, $p = .45$) and included only individuals who endorsed negative words as self-relevant ($n = 19$).

Free recall. There was a main effect of condition for words recalled, $F(1,30) = 23.48$, $p < .001$, $\eta^2 = .44$, whereby participants recalled more positive than negative words across assessments. Perhaps not surprisingly, there was a significant main effect of time, $F(1,30) = 24.42$, $p < .001$, $\eta^2 = .45$. During the first administration, the recall task was unanticipated. However, it was likely expected during the follow-up assessment. Thus, participants recalled more positive and negative words at the follow-up assessment compared to the initial assessment. Nevertheless, the Time \times Condition interaction was not significant, $F(1,30) = 0.98$, $p = .33$, $\eta^2 = .03$. Analyses indicated significant test-retest correlations for positive ($r = .63$, $p < .001$) and negative ($r = .73$, $p < .001$) words recalled over time (see Figure 1B,C).

Recognition. In the recognition portion of the task, there was a main effect of condition, $F(1,30) = 30.22$, $p < .001$, $\eta^2 = .50$, as more positive words than negative words were recognized. No main effect of time emerged, $F(1,30) = 0.15$, $p = .70$, $\eta^2 = .01$, and the Time \times Condition interaction was not significant, $F(1,30) = 0.001$, $p = .97$, $\eta^2 < .001$. Test-retest correlational analyses showed associations for positive ($r = .53$, $p = .002$) and negative ($r = .44$, $p = .01$) words recognized over time (see Figure 1D,E).

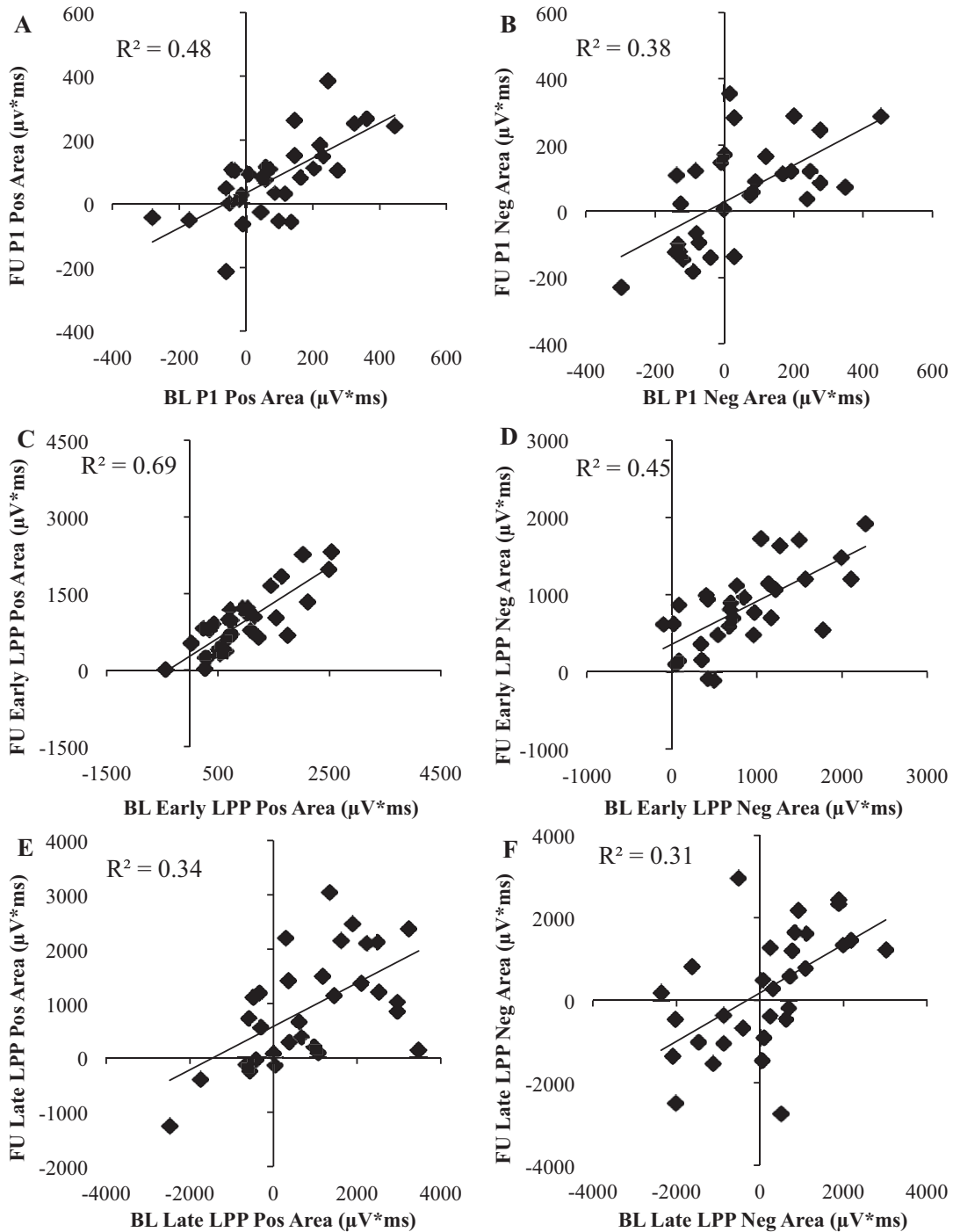


Figure 3. Test-retest for ERPs at the baseline and follow-up assessment. BL = baseline; FU = follow-up; Pos = positive words; Neg = negative words.

Processing bias. The main effect of condition was significant, $F(1,30) = 179.17, p < .001, \eta^2 = .86$, with adolescents showing a greater positive than negative processing bias. Similar to the free recall effects described earlier, there also was a main effect of time, $F(1,30) = 12.68, p = .001, \eta^2 = .30$. Interestingly, the Time \times Condition interaction was significant, $F(1,30) = 5.06, p = .03, \eta^2 = .14$, likely reflecting within-condition effects, as there was an increase in the positive ($p = .003, \eta^2 = .26$) but not the negative ($p = .35, \eta^2 = .03$) processing bias. Test-retest analyses revealed

significant associations for the positive ($r = .62, p < .001$) and negative ($r = .57, p = .001$) processing bias over time.

ERPs

The previously reported ERP results from the initial assessment (Auerbach et al., 2015) were replicated in this larger sample. When examining the P1, the Time \times Condition RMANOVA revealed a main effect of condition, $F(1,30) = 7.10, p = .01, \eta^2 = .19$, which

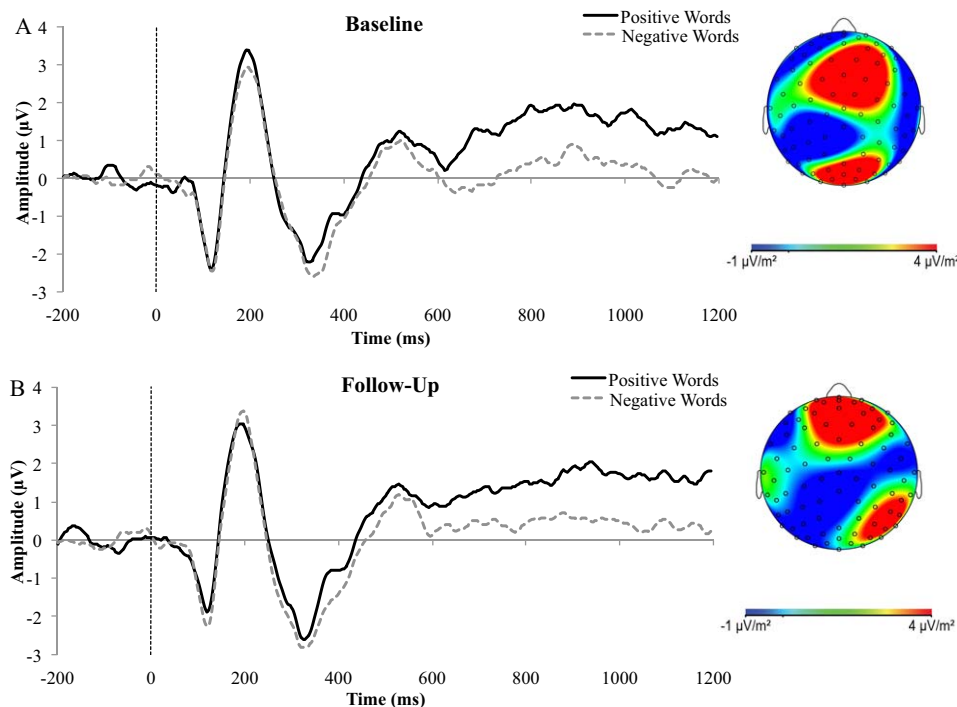


Figure 4. Late LPP responses at the baseline and follow-up assessment. Late LPP (600–1,200 ms poststimulus) activity in response to positive and negative words averaged across Fz, FCz, and Cz for healthy, female adolescents ($n = 31$) during the (A) initial, and (B) follow-up assessment. Scalp topographies reflect the difference between positive and negative words between 600–1,200 ms poststimulus.

indicated greater overall P1 positivity following positive words compared to negative words (Figure 2). The main effect of time, $F(1,30) = 0.003, p = .96, \eta^2 < .001$, and the Time \times Condition interaction, $F(1,30) = 0.42, p = .52, \eta^2 = .01$, were not significant. In line with our hypothesis, test-retest analyses demonstrated significant associations over time for P1 mean activity following positive ($r = .70, p < .001$) and negative ($r = .62, p < .001$) words (see Figure 3A,B).

For the early LPP, the Time \times Condition RMANOVA did not yield a main effect of condition; however, there was a trend in the expected direction, $F(1,30) = 2.92, p = .10, \eta^2 = .09$. The main effect for time, $F(1,30) = 0.06, p = .81, \eta^2 = .002$, and the Time \times Condition interaction, $F(1,30) = 0.02, p = .89, \eta^2 = .001$, were not significant. Test-retest analyses revealed associations over time for early LPP activity following positive ($r = .83, p < .001$) and negative ($r = .67, p < .001$) words (see Figure 3C,D).

In line with our hypothesis, analysis of the late LPP revealed a main effect of condition, $F(1,30) = 30.68, p < .001, \eta^2 = .51$, whereby participants exhibited an enhanced late LPP positivity following positive words compared to negative words (Figure 4). Fur-

ther, the main effect of time, $F(1,30) = 0.34, p = .56, \eta^2 = .01$, and the Time \times Condition interaction, $F(1,30) = 0.03, p = .86, \eta^2 = .001$, were not significant. The test-retest correlation for positive ($r = .59, p = .001$) and negative ($r = .56, p = .001$) words was significant (see Figure 3E,F).

Internal Reliability and Correlational Analyses

To test the internal reliability of our ERP indices, we computed the odd-even trial correlations for each component at the baseline and follow-up assessment. For the P1, the odd-even trial correlations were strong (Time 1: $r = .68, p < .001$; Time 2: $r = .52, p = .003$). Similarly, the internal consistency for the early LPP (Time 1: $r = .80, p < .001$; Time 2: $r = .81, p < .001$) and late LPP (Time 1: $r = .55, p = .001$; Time 2: $r = .75, p < .001$) ERPs indicated large effects.

Correlations for baseline and follow-up SRET behavioral and ERP indices are summarized in Tables 2A and 2B. Interestingly, at both the initial and follow-up ERP assessment, greater late LPP positivity following positive words was associated with greater free

Table 2A. Correlation Among Behavioral Indices and ERPs at the Initial (Below Diagonal) and Follow-Up (Above Diagonal) Assessment: Positive Words

	1	2	3	4	5	6	7	8
1. Endorse	–	–0.07	–0.10	0.04	0.13	0.11	–0.06	0.10
2. RT	0.22	–	–0.16	–0.45	–0.03	0.21	0.14	–0.19
3. Recall	–0.12	–0.47*	–	0.45*	0.77**	–0.17	–0.30	0.42*
4. Recognition	–0.29	–0.17	0.47**	–	0.33	–0.06	–0.33	0.10
5. Processing	0.12	–0.28	0.87**	0.30	–	–0.02	–0.29	0.46*
6. P1	–0.03	0.27	–0.12	–0.01	–0.19	–	0.09	0.27
7. Early LPP	0.11	–0.10	–0.26	–0.05	–0.12	0.43	–	–0.33
8. Late LPP	–0.08	–0.29	0.44*	0.26	0.42*	–0.15	–0.37*	–

Table 2B. Correlation Among Behavioral Indices and ERPs at the Initial (Below Diagonal) and Follow-Up (Above Diagonal) Assessment: Negative Words

	1	2	3	4	5	6	7	8
1. Endorse	–	–0.26	0.37*	0.30	0.83**	0.10	–0.21	0.20
2. RT	–0.31	–	0.26	0.07	–0.15	0.19	0.47*	–0.07
3. Recall	0.45**	–0.23	–	0.51**	0.55**	0.23	0.13	0.36*
4. Recognition	0.35	–0.10	0.51**	–	0.43*	–0.12	–0.07	0.33
5. Processing	0.95**	–0.36	0.55**	0.39*	–	–0.01	–0.13	0.17
6. P1	0.12	0.15	0.26	0.01	0.06	–	0.17	0.19
7. Early LPP	–0.27	0.39	–0.05	–0.05	–0.24	0.30	–	–0.25
8. Late LPP	0.04	–0.03	0.38*	0.15	0.02	–0.01	–0.10	–

Note. Participants who did not endorse any negative words as self-relevant were excluded from RT correlations (baseline excluded $n = 12$; follow-up excluded $n = 12$).

* $p < .05$. ** $p < .01$.

recall of positive words and a higher positive processing bias score. Conversely, potentiated late LPP positivity in response to negative words was associated with greater recall of negative words across assessments. No other associations emerged between behavioral and ERP indices.

Discussion

Toward the goal of identifying clinically useful biobehavioral markers of depressotypic self-referential processing, the current study sought to test the behavioral and ERP stability of the SRET among healthy, female adolescents over a 3-month period. Extending our findings from our prior study (Auerbach et al., 2015) in a larger sample, healthy adolescents endorsed, recalled, and recognized more positive and fewer negative words, and this effect was stable over time. Similarly, healthy youth reported a faster RT when endorsing self-relevant positive words, as opposed to negative words, at both the initial and follow-up assessment. Second, ERP activity to positive and negative words was consistent over time; namely, healthy youth exhibited potentiated P1 and late LPP positivity to positive versus negative words across assessments. Several findings warrant additional attention.

Similar to past research studying the stability of behavioral markers in children (Goldstein et al., 2015), behavioral indices (i.e., endorsement, RT, recall, and recognition processing bias scores) among adolescents demonstrated stability over time with medium-to-large effect sizes. Additionally, although past research has probed whether ERP responses during the SRET differ among healthy and depressed individuals (Auerbach et al., 2015; Shestyk & Deldin, 2010) and at-risk youth (Speed et al., 2016), no research has tested the stability of the early and late ERP components elicited by the SRET. The current study showed medium-to-large effect sizes when examining the test-retest correlations for the P1, early LPP, and late LPP; namely, there was greater positivity following positive versus negative words across assessments. In addition to the stability of the ERP effects, correlational analyses revealed associations between the late LPP and free recall (as well as positive processing bias). These findings were not unexpected, as the late LPP reflects sustained engagement and encoding processes (Foti et al., 2009; Naumann, Bartussek, Diedrich, & Laufer, 1992). Overall, the stability of the behavioral markers and ERPs obtained during the SRET support its use in probing biobehavioral markers of psychopathology, particularly MDD.

It is important to note several limitations in the current study, which may be addressed in future research. First, to reduce the heterogeneity of our sample, the study included only female adolescents. Although there is no reason to believe that the findings would not extend to male adolescents, future research should address this issue. Additionally, the majority of our study sample was Caucasian, and, consequently, research is warranted to test the generalizability of our findings to more diverse samples. Second, originally the SRET has been used to probe negative self-schema in depressed populations (Derry & Kuiper, 1981; Goldstein et al., 2015). In the current study, healthy adolescents endorsed few negative words as being self-relevant. These low rates of endorsement precluded us from computing ERPs only in response to endorsed adjectives. Additionally, recognition and free recall effects may be influenced by age and intelligence. Third, the current study tested stability over a 3-month period; however, future research would benefit from testing longer periods, particularly as it may relate to stability across developmental periods (e.g., adolescence to adulthood). Fourth, the study was sufficiently powered to test the stability of behavioral and ERP markers. Nonetheless, the small sample size precluded the implementation of a principal component analysis of our ERP effects. Future research also would benefit from testing the stability of these indices in a depressed sample of adolescents. Fifth, the present study provided a necessary first step to test the stability and reliability of ERP and behavioral markers in healthy youth. Moving forward, it will be essential to determine whether these indicators are stable in clinical populations (e.g., depression) both in current and remitted states. Sixth, the inclusion of the same words at the initial and follow-up assessment may reduce the novelty of the words during the second ERP assessment. Last, no pubertal information was obtained from our participants, which may have important implications for understanding the development of self-referential processing biases. This issue should be considered in future research.

In summary, prior research has demonstrated differences in behavioral (Connolly, Abramson, & Alloy, 2016; Timbremont & Braet, 2004; Zupan et al., 1987) and ERP (Auerbach et al., 2015) effects when using the SRET among healthy and depressed adolescents. The current findings also suggest that these effects remain stable over time in healthy adolescents. A question at large, however, is whether neurophysiological processes associated with depressotypic self-referential processing biases normalize in response to psychotherapeutic and pharmacological treatment. Addressing this critical issue may lead to key clinical insights with respect to designing more targeted treatment for youth with MDD.

References

- Alloy, L. B., Abramson, L. Y., Murray, L. A., Whitehouse, W. G., & Hogan, M. E. (1997). Self-referent information-processing in individuals at high and low cognitive risk for depression. *Cognition and Emotion, 11*(5–6), 539–568. doi: 10.1080/026999397379854a
- American Psychiatric Association. (2000). *Text revision (DSM-IV-TR)*. Washington, DC: American Psychiatric Association.
- Auerbach, R. P., Stanton, C. H., Proudfit, G. H., & Pizzagalli, D. A. (2015). Self-referential processing in depressed adolescents: A high-density event-related potential study. *Journal of Abnormal Psychology, 124*(2), 233–245. doi: 10.1037/abn0000023
- Beck, A. T., Steer, R. A., & Brown, G. K. (1996). *Beck Depression Inventory Manual* (2nd ed.). San Antonio, TX: Psychological Corporation.
- Bradley, M. M., & Lang, P. J. (2010). *Affective norms for English words (ANEW): Stimuli, instruction manual, and affective ratings (Technical report C-2)*. Gainesville, FL: The Center for Research in Psychophysiology, University of Florida.
- Bromet, E., Andrade, L. H., Hwang, I., Sampson, N. A., Alonso, J., de Girolamo, G., . . . Kessler, R. C. (2011). Cross-national epidemiology of DSM-IV major depressive episode. *BMC Medicine, 9*, 90. doi: 10.1186/1741-7015-9-90
- Connolly, S. L., Abramson, L. Y., & Alloy, L. B. (2016). Information processing biases concurrently and prospectively predict depressive symptoms in adolescents: Evidence from a self-referent encoding task. *Cognition and Emotion, 30*(3) 550–560. doi: 10.1080/02699931.2015.1010488
- Derry, P. A., & Kuiper, N. A. (1981). Schematic processing and self-reference in clinical depression. *Journal of Abnormal Psychology, 90*(4), 286–297. doi: 10.1037/0021-843X.90.4.286
- Dobson, K. S., & Shaw, B. F. (1987). Specificity and stability of self-referent encoding in clinical depression. *Journal of Abnormal Psychology, 96*(1), 34–40. doi: 10.1037/0021-843X.96.1.34
- Fischler, I., & Bradley, M. M. (2006). Event-related potential studies of language and emotion: Words, phrases, and task effects. *Progress in Brain Research, 156*, 185–203. doi: 10.1016/S0079-6123(06)56009-1
- Flor, H., Knost, B., & Birbaumer, N. (1997). Processing of pain- and body-related verbal material in chronic pain patients: Central and peripheral correlates. *Pain, 73*(3), 413–421. doi: 10.1016/S0304-3959(97)00137-1
- Foti, D., Hajcak, G., & Dien, J. (2009). Differentiating neural responses to emotional pictures: Evidence from temporal-spatial PCA. *Psychophysiology, 46*(3), 521–530. doi: 10.1111/j.1469-8986.2009.00796.x
- Goldstein, B. L., Hayden, E. P., & Klein, D. N. (2015). Stability of self-referent encoding task performance and associations with change in depressive symptoms from early to middle childhood. *Cognition and Emotion, 29*(8), 1445–1455. doi: 10.1080/02699931.2014.990358
- Hayden, E. P., Olino, T. M., Mackrell, S. V. M., Jordan, P. L., Desjardins, J., & Katsiroumbas, P. (2013). Cognitive vulnerability to depression during middle childhood: Stability and associations with maternal affective styles and parental depression. *Personality and Individual Differences, 55*(8), 892–897. doi: 10.1016/j.paid.2013.07.016
- Jaenicke, C., Hammen, C., Zupan, B., Hiroto, D., Gordon, D., Adrian, C., & Burge, D. (1987). Cognitive vulnerability in children at risk for depression. *Journal of Abnormal Child Psychology, 15*, 559–572. doi: 10.1007/BF00917241
- Jormann, J., Dkane, M., & Gotlib, I. H. (2006). Adaptive and maladaptive components of rumination? Diagnostic specificity and relation to depressive biases. *Behavior Therapy, 37*(3), 269–280. doi: 10.1016/j.beth.2006.01.002
- Kaufman, J., Birmaher, B., Brent, D., Rao, U., Flynn, C., Moreci, P., . . . Ryan, N. (1997). Schedule for affective disorders and schizophrenia for school-age children—Present and lifetime version (K-SADS-PL): Initial reliability and validity data. *Journal of the American Academy of Child and Adolescent Psychiatry, 36*(7), 980–988. doi: 10.1097/00004583-199707000-00021
- Kessler, R. C. (2012). The costs of depression. *Psychiatric Clinics of North America, 35*(1), 1–14. doi: 10.1016/j.psc.2011.11.005
- Kuiper, N. A., & Derry, P. A. (1982). Depressed and nondepressed content self-reference in mild depressives. *Journal of Personality, 50*(1), 67–80. doi: 10.1111/j.1467-6494.1982.tb00746.x
- Kujawa, A., Klein, D. N., & Proudfit, G. H. (2013). Two-year stability of the late positive potential across middle childhood and adolescence. *Biological Psychology, 94*(2), 290–296. doi: 10.1016/j.biopsycho.2013.07.002
- Lemogne, C., Mayberg, H., Bergouignan, L., Volle, E., Delaveau, P., Lehericy, S., . . . Fossati, P. (2010). Self-referential processing and the prefrontal cortex over the course of depression: A pilot study. *Journal of Affective Disorders, 124*(1–2), 196–201. doi: 10.1016/j.jad.2009.11.003
- Matt, G. E., Vázquez, C., & Campbell, W. K. (1992). Mood-congruent recall of affectively toned stimuli: A meta-analytic review. *Clinical Psychology Review, 12*(2), 227–255. doi: 10.1016/0272-7358(92)90116-P
- Moulds, M. L., Kandris, E., & Williams, A. D. (2007). The impact of rumination on memory for self-referent material. *Memory, 15*(8), 814–821. doi: 10.1080/09658210701725831
- Naumann, E., Bartussek, D., Diedrich, O., & Laufer, M. E. (1992). Assessing cognitive and affective information processing functions of the brain by means of the late positive complex of the event-related potential. *Journal of Psychophysiology, 6*(4), 285–298.
- Prieto, S. L., Cole, D. A., & Tageron, C. W. (1992). Depressive self-schemas in clinic and nonclinic children. *Cognitive Therapy and Research, 16*(5), 521–534.
- Shestyuk, A. Y., & Deldin, P. J. (2010). Automatic and strategic representation of the self in major depression: Trait and state abnormalities. *American Journal of Psychiatry, 167*(5), 536–544. doi: 10.1176/appi.ajp.2009.06091444
- Speed, B. C., Nelson, B. D., Auerbach, R. P., Klein, D. N., & Hajcak, G. (2016). Depression risk and electrocortical reactivity during self-referential emotional processing in 8 to 14 year-old girls. *Journal of Abnormal Psychology*. Advance online publication. doi: 10.1037/abn0000173
- Timbremont, B., & Braet, C. (2004). Cognitive vulnerability in remitted depressed children and adolescents. *Behaviour Research and Therapy, 42*(4), 423–437. doi: 10.1016/S0005-7967(03)00151-7
- West, W. C., & Holcomb, P. J. (2000). Imaginal, semantic, and surface-level processing of concrete and abstract words: An electrophysiological investigation. *Journal of Cognitive Neuroscience, 12*(6), 1024–1037. doi: 10.1162/08989290051137558
- Zupan, B. A., Hammen, C., & Jaenicke, C. (1987). The effects of current mood and prior depressive history on self-schematic processing in children. *Journal of Experimental Child Psychology, 43*(1), 149–158. doi: 10.1016/0022-0965(87)90056-7

(RECEIVED March 31, 2016; ACCEPTED May 13, 2016)