- A Meta-Analysis of Expressive Writing on Quality of Life, Posttraumatic Growth, and
- Posttraumatic Stress

research are discussed.

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Abstract

Emotional expression has been shown to be beneficial for promoting both positive psychological and physical health outcomes. Unfortunately, inhibiting emotions can lead to impairments in physical and psychological health. James Pennebaker showed that expressive writing is an effective form of emotional expression, and he and others have used expressive writing as an experimental manipulation to gauge its efficacy in treating a wide variety of health-related and psychological outcomes. While many studies have been conducted that examine the efficacy of expressive writing across such outcomes, a considerable amount of 10 these studies tend to neglect necessary considerations such as different levels of 11 symptomology, power, and meaningfulness of respective effect sizes. Six previous 12 meta-analyses have been conducted that examine expressive writing's effect on psychological 13 outcomes. However, these studies focus on the experimental versus control group effect size. 14 Thus, our meta-analysis sought to examine the efficacy of an expressive writing task on only 15 the experimental conditions in studies measuring posttraumatic growth, posttraumatic stress, and quality of life using random effects models. Results indicated a small overall effect size for posttraumatic stress and negligible to small effect sizes for posttraumatic growth and 18 quality of life. However, those studies requiring a diagnosis of PTSD exhibited a medium to 19 large effect size. Implications for future research design and interpretation of published

Keywords: meta-analysis, posttraumatic stress, expressive writing

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$_{26}$ Emotional Expression

Emotional expression relating to negative emotions or trauma has been shown to 27 enhance both mental and physical health outcomes (Esterling, Antoni, Kumar, & 28 Schneiderman, 1990; Fawzy et al., 1993; Lieberman & Goldstein, 2006; Rachman, 1980; Scheff, 1979). For example, the disclosure of traumatic or stressful events has been shown to reduce stress and lead to positive health outcomes in those with diabetes (Bodor, 2002) and breast cancer (Stanton et al., 2002), among others. Inhibiting thoughts or emotions, rather, may be detrimental to both physical and psychological health (Goldstein, Edelberg, Meier, & Davis, 1988; Gross & Levenson, 1997; Larson & Chastain, 1990). While some studies suggest that emotional expression in the form of "truth telling" may cause psychological harm to 35 individuals (Brounéus, 2010), the literature presents evidence confirming the negative effects of a lack of emotional expression, such as social concerns, overall psychological dysfunction, 37 and lack of value-congruent behaviors (Frankl, 1959; Pennebaker, 1989; Pennebaker & Beall, 38 1986; Schulenberg, Hutzell, Nassif, & Rogina, 2008; Wilson & DuFrene, 2009). These resulting negative outcomes may lead to detrimental effects on health (Pennebaker & Beall, 1986). Individuals having experienced a traumatic or stressful life event are significantly more likely to repress thoughts and feelings about their experience compared to individuals who have not experienced such events, thereby subjecting them to potential negative outcomes caused by a lack of emotional expression (Bodor, 2002).

Expressive Writing as Effective Emotional Expression

Pennebaker and Beall (1986) first showed that emotional expression can be both
experimentally manipulated and have positive benefits to participants. In their seminal
study, they randomly assigned participants to several writing groups, including writing about
a "stressful or traumatic" life event or a neutral event. As such, the content of the writing

likely varies widely based on contextual factors (e.g., topic, setting, sample, health concern).

The group that disclosed both regarding their trauma and the emotions surrounding said trauma later showed a reduction in health visits. Pennebaker has replicated the use of expressive writing across a number of studies ranging from improved health (Pennebaker, Colder, & Sharp, 1990; Pennebaker, Kiecolt-Glaser, & Glaser, 1988) to improvements in school (Pennebaker & Francis, 1996) and work (Francis & Pennebaker, 1992). Others have expanded this work to show positive effects on mood (Schoutrop, Lange, Hanewald, Davidovich, & Salomon, 2002) and asthma (Smyth, Stone, Hurewitz, & Kaell, 1999); however, several controlled studies have shown to not replicate (Harris, Thoresen, Humphreys, & Faul, 2005) or null effects (Gidron, Peri, Connolly, & Shalev, 1996; Walker, Nail, & Croyle, 1999). This protocol, more generally, has been termed written emotional disclosure (WED).

The idea that a brief, controlled writing task can have numerous positive health and 62 psychological benefits can certainly be controversial, given the existing literature. For 63 example, Henry, Schlegel, Talley, Molix, and Bettencourt (2010) found that expressive writing only benefited a rural population for those individuals surviving breast cancer on physical and psychological health outcomes, while Lancaster, Klein, and Heifner (2015) found no significant evidence that expressive writing can be considered an effective approach in measuring posttraumatic growth. Regardless, the concept remains interesting due to the nature and inexpensive implementation of expressive writing. Many individuals who have experienced traumatic events do not wish to disclose their feelings regarding the events with others. Additionally, those who do not meet diagnostic criteria (e.g., subclinical symptoms) are sometimes neglected despite probable suffering (Wilson & DuFrene, 2009). However, by utilizing expressive writing as a personal method of treatment, individuals are able to effectively express their emotions while avoiding talking to another individual or clinician about the traumatic event (Smyth, 1998). Pennebaker (1993) found that participants in 75 experimental conditions assigned to participate in an expressive writing task generally report more positive changes than those in control conditions. Some controversy has been observed
over whether or not writing about a formerly disclosed event is more effective than writing
about an undisclosed event. Greenberg and Stone (1992) conducted an experiment where
they separated participants into three groups: writing about a formerly disclosed trauma,
writing about an undisclosed trauma, and a control group. They found no difference between
groups in efficacy. However, they did find that those who disclosed more severe traumas
reportedly experienced fewer physical symptoms at follow up, which suggests that the type
of trauma revealed can play a significant impact on symptom reduction and physical health.
A review of current meta-analyses relative to expressive writing is presented in a subsequent
section.

87 Possible Mechanisms Underlying WED Efficacy

In order to understand why expressive writing is considered to be efficacious, one must 88 examine the cognitive, social, and behavioral processes by which it promotes information processing. Pennebaker et al. (1990) discovered that individuals who benefited from 90 expressive writing attributed their success from the writing task to a renewed sense of 91 understanding. Further, Pennebaker (1993) conducted a textual analysis on expressive 92 writing content and found that those who were more successful during the task used causation words. Pennebaker thus concluded that expressive writing was a way for individuals to effectively process the event cognitively, which may explain the aforementioned renewed sense of understanding and excess of causation-oriented words. Aside from theories related to cognitive-processing and inhibition, there are a number of other theories related to emotional disclosure that warrant mentioning. The first is the social integration model (Pennebaker & Graybeal, 2001). This model suggests emotional disclosure can have a positive impact on how people interact in their environment. This increased 100 environmental interaction has been shown to have positive benefits on health (Frattaroli, 101 2006). Second, expressive writing parallels exposure therapy for phobias and Posttraumatic 102

Stress Disorder (PTSD), which suggests that repeatedly exposing oneself to the fear or 103 trauma can reduce the negative emotions or physical sensations associated with that fear or 104 trauma (Meshberg-Cohen, Svikis, & McMahon, 2014). Given that exposure therapy has 105 been shown to be effective for reducing symptoms of posttraumatic stress (PTS; Sloan, 106 Marx, & Epstein, 2005), one would expect individuals in these studies to experience a 107 reduction in PTS symptoms after taking part in an expressive writing task. Third, Wilson 108 and DuFrene (2009) discussed how the nonjudgmental acceptance of emotions leads to 109 positive health benefits by promoting value-congruent behavior, one of the main facets of 110 Functional Contextualism theory and the field of positive psychology (Frankl, 1959; 111 Schulenberg et al., 2008). Indeed, emotional expression in the form of expressive writing 112 could be considered a form of nonjudgmental acceptance, although it may not necessarily 113 lead to behavior change in all cases. Finally, a recently proposed theory that may help explain positive outcomes is referred to as a distance perspective (Kross & Ayduk, 2011). 115 This theory posits that, when individuals adopt a psychologically distanced perspective, they are better able to better understand their life situation. In sum, it seems likely that there are 117 multiple underlying mechanisms that account for the beneficial outcomes associated with 118 expressive writing described below. Indeed, the wide range of theoretical perspectives 119 provide further evidence which suggests that expressive writing is applicable in a variety of 120 contexts. Previously conducted meta-analyses, however, present varying results. 121

22 Meta-Analytic Techniques

Meta-analyses allow researchers the opportunity to collectively examine the efficacy of different psychological interventions/tasks on outcome variables (Borenstein, Hedges, & Rothstein, 2007; Glass, 1976; Hedges, 1982). Although many studies produced positive outcomes associated with expressive writing, some of these studies tend to neglect important questions, the most important of which is whether or not the effect sizes are meaningful (Smyth, 1998). Meta-analyses are a technique that allows researchers to pool studies to

examine an overall, weighted, population effect (Borenstein et al., 2007). Several
meta-analyses of expressive writing and emotional expression have been explored that
warrant explanation: Smyth (1998); Frisina, Borod, and Lepore (2004); Frattaroli (2006);
Mogk, Otte, Reinhold-Hurley, and Kröner-Herwig (2006); van Emmerik, Reijntjes, and
Kamphuis (2013); and Reinhold, Bürkner, and Holling (2018). These meta-analyses have
laid a foundation for exploring the effects of writing on psychological outcomes.

Smyth (1998) conducted the seminal meta-analysis regarding the efficacy of expressive 135 writing. They included studies utilizing an expressive writing group and control group 136 (neutral topic). This particular analysis examined the efficacy of expressive writing on 137 psychological well-being, general health, and physical functioning. In sum, thirteen 138 studies/effect sizes were included, and the authors found an overall medium effect size, d =130 0.47, for the experimental group compared to the control group. A later meta-analysis 140 conducted by Frisina et al. (2004) expanded these analyses. They included studies utilizing 141 clinical samples and employing the paradigm adapted by Pennebaker and Beall (1986). This 142 meta-analysis included nine studies in total and found an effect size of d=0.19 for 143 health-related outcomes and d = 0.07 for psychological outcomes. The next expressive 144 writing meta-analysis was conducted by Mogk et al. (2006) and aimed to update the state of the literature on expressive writing. Similar to previously-conducted analysis, they included studies employing Pennebaker's paradigm on experimental and control groups. Additionally, they only included studies with a four-week follow up that included at least ten participants. In sum, thirty studies met their criteria. They found nonsignificant effects on somatic and 149 psychological health outcomes and concluded that expressive writing does not promote 150 health-related outcomes. These findings corroborate those from Frisina et al. (2004). 151

Frattaroli (2006) conducted perhaps the most notable meta-analysis to date examining
the efficacy of emotional disclosure on the following constructs using only randomized and
control conditions: psychological health, physiological functioning, reported health, health
behaviors, and general functioning/life outcomes. Additionally, their meta-analysis was the

first to employ random effects models, which estimate the mean of a proposed distribution of 156 population effect sizes. Prior meta-analyses employed fixed effects models, which assume 157 that all studies assess the same "true" population effect size, which may be an untenable 158 assumption across different populations (Borenstein et al., 2007). They included a wide 159 range of studies N=146. Individual studies were again collapsed into one publication effect 160 size, although these effects were also examined separately by health outcome. Overall, 161 Frattaroli (2006) found a weighted r effect size of .08 for all outcomes combined, which 162 would be considered small. Additionally, they examined potential moderators and found 163 larger effect sizes for the following samples: those with physical health problems, those with 164 a history of having experienced traumatic or stressful events, samples not including college 165 students, samples where expressive writing tasks were conducted at home and in private 166 settings, paid participants, more male participants, and fewer participants (see Frattaroli, 2006 for a complete list of moderators). A recent analysis conducted by van Emmerik et al. (2013) employing Pennebaker's paradigm included six eligible studies that compared 169 treatment to control groups. In regards to inclusion criteria, they included studies where 170 participants had a diagnosis of Acute Stress Disorder (ASD) or PTSD. They found that 171 those who participated in the expressive writing group experienced short-term reductions in 172 PTS and comorbid depressive symptoms, combined Hedges' g = 0.81. The most recently 173 published meta-analysis was conducted by Reinhold et al. (2018) and examined the effects of 174 expressive writing on depression by randomizing participants to conditions (expressive 175 writing vs. control). They included thirty-nine randomized controlled trials and excluded 176 individuals with diagnoses of PTSD. This study did not support utilizing expressive writing 177 for depression outcome measures for the specified sample, $g_{post} = -0.09$. Further, they found 178 that expressive writing did not yield any type of long-term effect on depression outcomes. 179

Posttraumatic Stress

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or events after a trauma. This generates a context where individuals are prone to 182 affect-related deficiencies and maladaptive behaviors (American Psychiatric Association, 183 2013). DSM-5 criteria are based on twenty symptoms structured into four different subsets 184 in those having experienced a traumatic event. These subsets are as follows: re-experiencing, 185 avoidance, negative alterations in cognition and mood, and increased arousal (Crespo & 186 Gomez, 2016). While the renewed DSM-5 criteria are now increasingly employed, the current 187 meta-analysis considers studies using DSM-IV criteria. DSM-IV criteria are similar and 188 include the following: exposure to a traumatic event, re-experiencing (intrusion), avoidance, 189 and increased arousal (American Psychiatric Association, 2013). Further, the studies employed in the current meta-analysis are divided according to these subsets (arousal, intrusion, and avoidance). PTSD affects a wide variety of groups, a few of which are sexual 192 assault survivors (Klump, 2008), Iraq and Afghanistan war veterans (Gentes et al., 2014), 193 and those exposed to natural disasters (Wang et al., 2000). 194 Research conducted on the efficacy of expressive writing on PTSD symptoms presents 195 intriguing results. Sloan, Marx, Epstein, and Lexington (2007) examined individuals with at 196 least moderate PTSD symptom severity and found that individuals assigned to an emotional 197 expression writing condition reported fewer PTSD and depression symptoms during follow 198 up. Sloan, Marx, and Greenberg (2011) found that PTSD symptoms decreased after a 199 written emotional disclosure task, although this decrease was not significantly different than 200 a control group change. Di Blasio et al. (2015) recruited women who had just given birth and assessed them a few days after experiencing childbirth along with a three-month follow-up. Results showed that women who had participated in the expressive writing task had lower depression and posttraumatic stress symptoms than the group assigned to a neutral writing condition. Additionally, regression models showed that expressive writing 205 was significantly linked to a reduction of PTSD symptoms across different dimensional levels 206

Posttraumatic Stress Disorder (PTSD) is a disorder involving re-experiencing thoughts

of symptom severity. Only 20 of the 113 women recruited for this study qualified for a 207 diagnosis of PTSD, but those who reported mild symptomology responded better to the task 208 than those meeting criteria for PTSD. This limitation suggests that those with moderate 209 distress could perhaps benefit more from an expressive writing task than those diagnosed 210 with or meeting the qualifications for PTSD. It may also explain the differences in results in 211 comparing to Sloan et al. (2011), as they found that those with a clinical diagnosis of PTSD 212 did not respond to an emotional disclosure writing task. Perhaps it may be more 213 advantageous to examine effect sizes separately for diagnoses of PTSD and subclinical 214 symptoms. Further Sloan, Marx, Bovin, Feinstein, and Gallagher (2012) adapted a writing 215 protocol to focus primarily on the emotions, meaning, and "hot spots" associated with the 216 trauma. They referred to this procedure as the written exposure therapy (WET) protocol, 217 distinguishable from the paradigm adapted by Pennebaker and Beall (1986). In their seminal study examining the efficacy of WET for motor-vehicle accident related PTSD, they found 219 that those in the WET condition experienced significant reductions in PTSD symptoms throughout the course of the study. Since then, a small number of other studies employing 221 the WET procedure have been employed in those with PTSD. While these will be included 222 in the current review, the newness of this protocol does not allow exclusive examination using meta-analytic techniques.

25 Posttraumatic Growth

While the literature mostly discusses potentially harmful outcomes to traumatic events such as emotional distress, traumatic events also provide opportunities for personal growth (Aslam & Kamal, 2013). Traumatic events, either natural or human-inflicted, can lead to positive outcomes by allowing the individual to take a different perspective (Cobb, Tedeschi, Calhoun, & Cann, 2006; Taku, Calhoun, Cann, & Tedeschi, 2008). The relationship between positive growth after a traumatic event and symptom reduction is unclear, as it is a complex process. Thus, it is necessary to examine how expressive writing might influence each

variable separately, which is one of the key goals of this meta-analysis (Slavin-Spenny, Cohen, 233 Oberleitner, & Lumley, 2011). Models receiving empirical support within the last decade 234 suggest that traumatic events offer opportunities for both negative and positive experiences 235 (Tedeschi & Calhoun, 1995; Weiss, 2002). Posttraumatic Growth (PTG) is a positive 236 experience after a traumatic event (Aslam & Kamal, 2013; Yilmaz & Zara, 2016). 237 Specifically, PTG is classified as broad cognitive benefits that are seen after a traumatic 238 experience. These benefits can be categorized into building closer relationships, examining 230 new possibilities, appreciating life, recognizing personal strengths, and undergoing spiritual changes (Dursun, Steger, Bentele, & Schulenberg, 2016; Tedeschi & Calhoun, 2004). 241 PTG is associated with a variety of desired outcomes (Dursun et al., 2016). PTG has 242 been studied in those experiencing natural disasters, war, and other harms such as sexual 243 assault. Finally, PTG has been studied in those experiencing medical diagnoses such as 244 different types of cancer and diseases. Although the relationship between PTG and symptom 245 reduction is not yet fully understood, perhaps expressive writing allows the individual to 246 fully comprehend the event. Pennebaker and Graybeal (2001) speculated that expressive 247 writing allows an individual to feel more connected with his or her surroundings. Although 248 this speculation does not directly explain positive outcomes after an expressive writing task, perhaps individuals gain a better appreciation for life after gaining a better sense of 250 connectedness with that individual's surroundings. One might expect effect sizes to be larger 251 for those studies requiring a diagnosis of PTSD, as such growth may not be possible in those 252 with subclinical symptomology.

254 Quality of Life

Quality of Life (QOL), according to Theofilou (2013) is an evaluation of the "goodness" that an individual experiences, separated into domains of reactions to life events, disposition, life fulfillment, and satisfaction with life experiences. More generally, QOL refers to an individual's attitude towards the target life situation (Costanza et al., 2007), delineated into

objective and subjective components. Objectively, QOL refers to components outside of an 259 individual and measurable by others, while subjective QOL is an individual's assessment of 260 his or her own experiences (Costanza et al., 2007). The current meta-analysis will focus 261 solely on the subjective components of QOL, as it is obtainable through questionnaires. 262 Pennebaker and Graybeal (2001) suggested that expressive writing allows one to feel more 263 connected with their surroundings. Further, they explain that expressive writing allows 264 people to see things in a different way and better understand themselves. By understanding 265 a traumatic or stressful event, one is said to see things differently and perhaps look at the 266 situation with a more positive mindset. The changes that occur after expressive writing may 267 also allow one to find meaning in the traumatic event, thereby increasing the QOL of that 268 individual (Frankl, 1959). Higher QOL may be considered a type of PTG, which is why the 269 current meta-analysis sought to examine the efficacy of studies utilizing expressive writing to 270 improve QOL and PTG in the same study. 271

272 Current Meta-Analysis

The purpose of the current meta-analysis is to examine studies employing expressive 273 writing procedures using Pennebaker's paradigm (WED) and the more recent WET protocol 274 on variables relevant to the field of positive psychology (PTG and QOL) and PTS, with effect 275 sizes separated by the paper's indication of PTSD diagnosis when sample sizes are large 276 enough. Based on recently published literature regarding efficacy of expressive writing for 277 different levels of PTSD symptoms, this marker is an important facet to consider (Di Blasio 278 et al., 2015; Reinhold et al., 2018; Sloan et al., 2011). No review has examined the effects of expressive writing on PTS separated by diagnosis. Additionally, no meta-analysis has been 280 conducted that examines the efficacy of expressive writing on positive outcome variables such 281 as PTG and QOL, in line with the field of positive psychology. The meta-analyses described 282 sequentially above also focused on experimental versus control group effect sizes or p-values, 283 rather than emphasizing change for the expressive writing group. This focus is likely because

of the analyses provided in these publications, especially when using randomized controlled 285 trial research designs. While this design is the gold standard for medicine, the current 286 meta-analysis sought to examine the magnitude of change for participants who experienced 287 an expressive writing task. For example, a comparison group may increase their quality of 288 life scores by two points in a controlled study, while the experimental group increases their 280 quality of life scores by four points; thus, creating a significant difference in change between 290 the two groups. This information is valuable, but it does not tell the reader the magnitude of 291 the change for the writing group, wherein four points might only be a small effect when 292 examined within the group who received the writing task. 293

This analysis will also focus on changes across time for groups who received the 294 expressive writing task to determine what size of effects one might expect given a specific 295 measurement schedule (i.e., one to three months, three months to six months, etc.). This 296 analysis should present researchers with a renewed examination of the efficacy of expressive 297 writing on the aforementioned variables using newer meta-analytic techniques. Newer 298 methods of meta-analysis, including p-curve (Simonsohn, Nelson, & Simmons, 2014; 290 Simonsohn, Simmons, & Nelson, 2015), p-uniform (van Aert, Wicherts, & van Assen, 2016), 300 PET-PEESE (Stanley & Doucouliagos, 2014), selection models (Vevea & Hedges, 1995), and 301 trim and fill methods (Carter & McCullough, 2014) allow for better estimation of 302 meta-analytic effect sizes. These analyses would be best performed by examining each 303 potential effect separately, rather than averaging effects of each publication into one study effect size (a common trend in the previously mentioned meta-analysis). In addition to an estimate of overall effect sizes using updated techniques, the current meta-analysis estimates power for effects on writing groups, as research has shown a consistent under powering of 307 psychological studies, combined with a misunderstanding of the sample size needed for 308 adequately powering one's work (Bakker, Hartgerink, Wicherts, & van der Maas, 2016). 309

310 Method

Data Collection

Studies were collected through online databases, such as PsycINFO and Google 312 Scholar, using the following search terms and their combinations: Posttraumatic Growth, 313 PTG, Quality of Life, QOL, Posttraumatic Stress, PTS, Expressive Writing, Emotional Disclosure, Written Emotional Disclosure (WED), Written Exposure Therapy (WET). 315 Within these articles, the change in outcome variables (PTS, PTG, QOL) from pre- to 316 post-test was the dependent variable of interest. Generally, groups were separated into an 317 experimental and control group and then examined at different time points. For purposes of 318 this meta-analysis, only participants assigned to the experimental condition were examined 319 due to having received the expressive writing task. If a study included multiple assessment 320 time points, then these measurements were examined sequentially (i.e., time 1 to time 2, 321 time 2 to time 3) to determine change across time for the dependent variable. 322 264 citations focusing on PTS, PTG, and QOL were identified through the literature 323 search and previous meta-analyses. After screening these studies, 53 articles were retained for 324 containing the appropriate information for this meta-analysis. This manuscript was written 325 with papaja in R (Aust & Barth, 2017) with the analyses inline with the text. The complete 326 set of data, excluded article list with reasoning, and other relevant information can be found 327 at: https://osf.io/4mjqt. Generally, studies were included if they utilized WED or WET, 328 included relevant numbers to compute an effect size, and included the relevant outcome 329 variables. After having two reviewers independently code articles, 223 effect sizes were 330 calculated. On average, each study represented M = 4.21, SD = 3.31 effects, ranging from 1 to 16 effects. 163 effects were calculated for PTS, 21 for PTG, and 37 for QOL. Studies were coded for PTSD diagnosis as no (not mentioned or not included), mixed (mentioned number 333 of participants but all included), and yes (included as criteria). After examining the number 334 of effects in each of these categories for each variable, only the PTS results will be split by 335 PTSD diagnosis with 16 no mention, 16 in the mixed category, and 86 yeses. 336

Calculations for Effect Size, Variance, and Confidence Intervals

For our purposes, we used Cohen's (1988) standards for nomenclature for small (0.20), medium (0.50), and large (0.80) d values, although it is important to note that Cohen himself suggested that these values should be based on the area of study. Generally, however, these effect size criteria are used within the social sciences. Each study implemented a pre-test to post-test style repeated measures design, usually with paired t-tests, ANOVA, or regression analyses. The means, standard deviations, and N values were collected from each study. In general, Cohen's d values were calculated using the following formula for paired t using means and standard deviations:

$$d_{av} = \frac{M_1 - M_2}{\frac{SD_1 + SD_2}{2}}$$

This equation is described in detail in Cumming (2012) as an alternative to the traditional calculation of d for paired samples t, wherein the denominator is the standard deviation of the difference scores:

$$d_z = \frac{M_1 - M_2}{SD_{diff}}$$

This equation for d_{av} not only allows for calculations from published articles that do not include SD_{diff} (i.e., most articles included), but also has been shown to be less upwardly biased than d_z . Alternative formulas include controlling for r between paired levels, as described in Lakens (2013); however, these values were not available in the selected articles, and Lakens also recommends d_{av} as an effect size for paired designs. When only mean differences and standard deviation of the difference scores were available, the second equation for d_z was used.

We planned to use traditional and newer methods of meta-analysis, following guidelines from Cooper, Hedges, and Valentine (2009) and Borenstein et al. (2007), as well as van Aert et al. (2016). Sampling variance of the effect sizes were estimated using the *escalc()* function from the *metafor* package in R (Viechtbauer, 2010). The variance formula was originally published in Morris and DeShon (2002) and is shown below:

$$v = \frac{1}{n} \left(\frac{n-1}{n-3}\right) \left(1 + n * d^2\right) - \frac{d^2}{[c(n-1)]^2}$$

In this formula, n is the number of paired observations, d is the calculated effect size, and c is a correction factor, wherein df are n-1 (Hedges, 1982):

$$c = 1 - \frac{3}{4 * df - 1}$$

We used the metagen() function in the metafor package to calculate both fixed and 363 random effects models, which uses standard error of the effect to calculate overall estimates 364 of an effect and their confidence intervals. Thus, we took the square root of the variance 365 estimate for standard error. Given these calculations, the goal of this analysis was to 366 calculate a combined effect size, along with a confidence interval for study planning and an 367 assessment of the literature. A fixed effects model requires the assumption that there is a 368 true population effect size across all studies. By including multiple measures of psychological 369 outcomes, this assumption may be tenuous, and therefore, a random effects model was also 370 calculated. In random effects models, the true effect is assumed to vary across studies 371 (Borenstein et al., 2007). For a fixed effects model, the effect sizes are weighted by their inverse variance (v; Sánchez-Meca & Marín-Martínez, 2008), which is calculated automatically in *metafor* by:

$$w_i^{FE} = \frac{1}{v}$$

The advantage to this procedure is that analyses are weighted by their precision, that is, that studies with more information (often, larger samples), are given larger weights in the overall estimated effect size (Borenstein et al., 2007). Random effects models are also weighted by inverse variance, with an additional correction for variance between studies, τ_{DL}^2 , as described by DerSimonian and Laird (1986):

$$w_i^{RE} = \frac{1}{v + \tau_{DL}^2}$$

Confidence intervals were calculated in two ways for this study. Cumming (2012), 380 Kelley (2007), and Smithson (2001) have shown that the distribution of d values are 381 non-normal, and thus, CIs should be estimated using the non-centrality parameter and a 382 non-normal distribution. These values were calculated using the functions in the MOTE 383 library which iteratively estimates the appropriate non-centrality parameter and converts 384 back to d values (i.e., non-centrality parameter divided by the square root of n; Buchanan, Valentine, & Scofield, 2017; Smithson, 2001, 2003). However, the metafor package in R uses central distributions to estimate CIs for each study and overall effect sizes. Therefore, we present both sets of values for the interested reader, as meta-analytic procedures have not 388 implemented non-central distributions of effect sizes. 389

390 Additional Meta-Analytic Techniques

p-Curve and p-Uniform. We used p-curve.com to conduct a p-curve analysis 391 (Simonsohn et al., 2014). The purpose of this type of analysis is to detect true effects. 392 Specifically, p-curve is used to reveal possible p-hacking in published literature in order to 393 decipher whether or not a true effect exists. Broadly, p-hacking occurs when researchers use 394 questionable research practices to create significant results by manipulating dependent 395 variables or covariates. Additionally, authors may add participants if the initial findings are 396 not significant (Bruns & Ioannidis, 2016). Researchers may also decide to exclude 397 participants for final analyses if that exclusion leads to a significant difference (John, Loewenstein, & Prelec, 2012). Thus, it is necessary to distinguish between true and false effects in order to effectively interpret effect sizes corresponding to those p-values. p-curve accomplishes this task by examining the distributions of the published p-values. If an effect 401 exists, or rather the results should be interpreted as presented, the distribution of p-values 402 will be positively skewed (Simonsohn et al., 2014). If, however, no effect exists, then the

distribution of p-values will be flat. p-curve analyses ultimately provide evidence of 404 p-hacking in groups of studies and has become an important tool for interpreting 405 meta-analyses. In order to accurately estimate effect sizes because of scrutiny associated 406 with effect size estimation of p-curve, we also conducted p-uniform. p-uniform analyses, too, 407 are interpreted by examining the distribution of p-values in a set of studies (van Aert et al., 408 2016). However, it is assumed that the population effect size equals the effect size from the 400 dataset. Because of this assumption, the population effect size is referred to as uniform. This 410 analysis also examines for publication bias and presents the researcher with a corrected effect 411 size. Publication bias occurs when only select studies are published, usually only significant 412 studies, although many factors can bias a study's publication (McShane, Böckenholt, & 413 Hansen, 2016). p-uniform was calculated from code provided by van Aert (2017) on GitHub. 414

PET-PEESE. Originally, meta-analyses relied on the calculation of Egger's 415 regression test which examined the relationship of the standard error (predictor) to the effect 416 size estimates (criterion). In this regression, the intercept values were used to determine if 417 effect size measures were different than zero, by providing a meta-analytic estimate (Egger, 418 Davey Smith, Schneider, & Minder, 1997; Stanley, 2005). PET-PEESE analyses examine for 419 publication bias by adapting parts from Egger's traditional regression tests: PET (Precision Effect Test) and PEESE (Precision Effect Estimate with Standard Error, Carter & 421 McCullough, 2014). PET is a more reliable test of publication bias with effect size estimates of zero, $b_0=0$, while PEESE is more accurate with non-zero effect size estimates, $b_0\neq 0$ 423 (Stanley & Doucouliagos, 2014). PET-PEESE was calculated using Hilgard's (2016) code 424 provided on GitHub. 425

Selection Models. Selection model analyses provide the researcher with a test of publication bias and effect size estimates using maximum likelihood estimation (Vevea & Hedges, 1995; Vevea & Woods, 2005). Using selection models, researchers are able to discover effect size estimates as well as evidence of publication bias (McShane et al., 2016) by using a mixed general linear model to estimate these values. Selection models were

calculated with the weightr package in R (Coburn & Vevea, 2017).

Trim and Fill. Trim and Fill analyses, in contrast to PET-PEESE, regress standard 432 error (criterion) and effect size estimates (predictor). Specifically, the purpose of Trim and 433 Fill techniques is to examine whether or not publication bias may influence the regression equation (Carter & McCullough, 2014). Effect sizes and standard error terms are graphically displayed on x and y-axes, respectively, in a funnel plot. If this graphical representation indicates asymmetry, considered a gap of missing data points in the lower center area of the plot, the study set can be assumed to have studies that are both non-significant and small in 438 sample size (van Assen, van Aert, & Wicherts, 2015). This funnel is then trimmed until 439 symmetry is achieved. Missing studies from the symmetrical graph are imputed (filled) while 440 maintaining the given symmetry (Duval & Tweedie, 2000). The meta-analytic effect size is 441 then estimated from the trimmed and filled funnel plot. Trim and fill analyses, as well as 442 funnel plots included below, were calculated with the *metafor* package. 443

Results

445 **PTS**

Overall Effect Size. As described above, both fixed effects and random effects 446 models with centralized confidence intervals are presented in Table 1. Studies were examined 447 for potential outliers using the metafor package in R. This package calculates traditional 448 regression influence values, such as Cook's and hat values (Cohen, 1988). These values indicate change in overall meta-analytic model with and without the effect; thus, 450 determining their impact on the pooled effect size (Viechtbauer, 2010). Because published 451 studies likely represent the range of the sampling distribution of effect sizes, we included the 452 analyses with and without outliers to present evidence for both paths a researcher might 453 take when examining an overall effect.

2 outliers were detected with this procedure, all showing very large effect sizes, average d=2.81. The fixed and random effects estimates without these points are also included in

Table 1. Figures 1, 2, 3, and 4 portray the effect sizes for PTS studies, separated by 457 intrusions, avoidance, hyperarousal, and total scores for easier viewing (i.e., over 100+ effect 458 sizes did not fit easily on one combined graph). Although these categories are not reflective 459 of updated DSM-5 criteria, researchers have not yet conducted enough studies using 460 expressive writing on PTS with updated PTSD criteria to warrant a meta-analysis. Name 461 acronym coding can be found in the data online. This forest plot includes the non-centralized 462 confidence interval calculated from the MOTE library (Buchanan et al., 2017). Shape size 463 indicates study weight, and these values were taken from the overall random effects 464 meta-analysis and normalized by dividing by the mean weight. The dashed lines indicate the 465 average non-weighted lower and upper confidence interval limit for the non-centralized 466 estimates. Overall, PTS studies include a small effect size that appears to be significantly 467 greater than zero across all estimate types (fixed, random, with or without outliers).

We further calculated the overall effect sizes by PTSD diagnosis category using a random effects model. Studies only including individuals with a diagnosis of PTSD exhibited a medium effect size (before and after outlier exclusion): with outliers d=0.64 [0.48, 0.79]; without outliers d=0.55 [0.41, 0.69], while studies not requiring (or listing) a PTSD diagnosis showed a small to medium effect size: d=0.32 [0.24, 0.40]. Similarly, the mixed category showed a small to medium effect size: d=0.35 [0.16, 0.54]. Complete estimates of all the following analyses split by diagnosis are included online at https://osf.io/4mjqt/, and their pattern of results is similar to the overall pattern here.

Homogeneity. A prerequisite for newer meta-analytic techniques includes the
assessment of homogeneity of the effects (van Aert et al., 2016). Using the *metafor* package
in R, we calculated the Q-statistic and the I² index (Cochran, 1954; Huedo-Medina,
Sánchez-Meca, Marín-Martínez, & Botella, 2006). Significant values imply inconsistencies
across the variable or variables of interest and are represented by Q. In contrast, I² indicates
the percentage of heterogeneity along with a 95% CI. Both can, however, be biased with a
small number of experiments included for analyses (Higgins, Thompson, Deeks, & Altman,

⁴⁸⁴ 2003; Huedo-Medina et al., 2006). Thus, we sought to calculate an overall level of ⁴⁸⁵ heterogeneity after examining each variable separately before and after excluding outliers. ⁴⁸⁶ For PTS studies including outliers, we found significant heterogeneity, Q(162) = 776.74, p <⁴⁸⁷ .001 and $I^2 = 79.1$, 95% CI[75.9 - 81.9]. These values were reduced slightly with the ⁴⁸⁸ exclusion of outliers, Q(160) = 677.98, p < .001 and $I^2 = 76.4$, 95% CI[72.6 - 79.7].

Power was calculated in two different ways using the pwr package in R 489 (Champely, 2016). Post hoc power was first calculated using sample size and effect size 490 statistics from each individual study. Additionally, we calculated power using the study 491 sample size and estimated overall effect size from the random effects model with and without 492 outliers, as explained by Francis (2012) and Francis (2014). The first estimate indicates the 493 likelihood of finding an effect from our sample statistics, while the second indicates the 494 likelihood of finding the true population effect size. If each study had been conducted on 495 only the change in the experimental group, 46.6% of studies would have been considered significant at $\alpha < .05$. The average power of these studies based on their original study 497 characteristics was .48 (SD = .36). Power for the random-effects meta-analytic effect size 498 with outliers was .52 (SD = .25) and without outliers was .49 (SD = .25). Therefore, power 499 consistently was around 40-50% for studies examining PTS, regardless of outlier effects. In these studies, only 28.8% achieved recommended 80% power for their found effect size, a smaller 24.5% for the random-effect outlier effect size, and even smaller 20.2% for power 502 calculations on the random-effect size without the outliers. 503

Other Meta-Analytic Estimates. As noted in van Aert et al. (2016), p-curve and p-uniform analyses are upwardly biased when heterogeneity is high. Therefore, we use caution when interpreting these analyses on PTS outcomes. As seen in Table 1, the estimates for p-uniform were higher than other techniques, likely because of the focus on significant p-values and the great degree of heterogeneity described earlier. p-curve pictures can be found at https://osf.io/4mjqt/ online, and this analysis indicated evidentiary value at p < .001. Additionally, the p-uniform analysis indicated that there was likely no publication

bias present, Z = -5.71, p = 1.000. When examining the PET analysis, we found that the 511 intercept was significant, which indicated that PEESE was likely a better estimator of the 512 meta-analytic effect size. PEESE estimates were lower than the original meta-analytic 513 estimate, but confidence intervals indicated that the effect is small to medium, and still 514 larger than zero. Selection models indicated a larger effect size, especially with the 515 random-effects models, and these effects were influenced by the outliers found in the 516 published studies. Trim and fill models are shown in Table 1, and figures are included online. 517 Nineteen missing studies were imputed for both models with and without outliers. Across all 518 these effect size estimates, we found that expressive writing was likely to decrease PTS 519 symptoms in a small to moderate way. The correlation of effect size with time between 520 measurement times was r = -.01, 95% CI [-.17, .14], t(161) = -0.15, p = .879, and 521 r = -.08, 95% CI [-.23, .08], t(159) = -1.00, p = .320 without outliers. This result indicated that the effect of expressive writing slightly decreased across time.

524 **PTG**

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Overall Effect Size. Both fixed and random effects models with centralized 525 confidence intervals for PTG are presented in Table 2. When examining expressive writing 526 on PTG, no outliers were detected. Fixed and random effects estimates are included in Table 527 2, while Figure 5 shows effect sizes for PTG studies where shape size indicates the 528 normalized weight of the study. Dashed lines indicate non-weighted lower and upper 529 confidence intervals for non-centralized estimates. Overall, PTG studies indicated a 530 negligible to small effect size across both random and fixed effects models, and the 531 non-centralized confidence intervals indicated an effect that crossed zero. **Homogeneity.** Using the *metafor* package in R, we calculated both a Q statistic 533 and I^2 index. Since PTG studied did not contain any outliers, we did not calculate two 534

separate analyses to examine heterogeneity both with and without outliers. We did not find

significant heterogeneity across PTG studies, Q(20) = 14.18, p = .821 and $I^2 = 0.0$, 95%

CI[0.0 - 25.3].

Power. First, we calculated *post hoc* power using both sample and effect size statistics from individual studies. Individual studies examining change in experimental groups showed that 9.5% of studies would have been considered significant at $\alpha < .05$.

Average power of PTG studies was .15 (SD = .16). 0.0% achieved recommended 80% power for their found effect size. Additionally, we calculated power using study sample size and estimated effect size from our random effects model. Power for the true effect size was .08 (SD = .02). Again, 0.0% achieved recommended 80% power.

Other Meta-Analytic Estimates. Due to no heterogeneity across PTG studies, 545 we can use both p-curve and p-uniform analyses with more confidence. A pictorial 546 representation of p-curve can be found at https://osf.io/4mjqt/. This analysis did not 547 indicate evidentiary value, p = .75, as only two of the results would be considered significant 548 at $\alpha < .05$. p-uniform estimates are presented in Table 2. Specifically, these analyses 549 indicated that there was no publication bias present, Z = 0.70, p = .243. The p-uniform 550 estimates of the effect size for PTG were negative, in contrast to the fixed and random 551 effects overall model. The confidence interval for this analysis indicates a wide range of 552 possible effects. In examining PET-PEESE analyses, we did not find a significant intercept, 553 indicating that PET is most likely a better effect size estimator. PET analyses indicated 554 that the effect size is negligible to small, with our confidence interval crossing zero. These 555 results corroborated our original effect size calculations. Selection models indicated negligible to small effect sizes, again wherein the confidence interval includes zero effect. Trim and fill models are shown in Table 2, and figures are included online. Zero studies were imputed for our model, and thus, the effect size estimate is the same as the overall model. Across 559 techniques, we found that expressive writing has little to no effect on PTG. The correlation 560 of effect size across measurement times in PTG studies at subsequent time points was 561 r = .09, 95% CI [-.36, .50], t(19) = 0.38, p = .707, and no change over time was found.

563 **QOL**

564

###Overall Effect Size

Finally, for QOL, both fixed and random effects models with centralized confidence 565 intervals are presented in Table 3. Two outliers were detected with this procedure, average d 566 = -0.07. While the average effect of these outliers indicates a small number, it is important 567 to note that these two outliers were the largest positive and negative effects found from the 568 Possemato, Ouimette, and Geller (2010) study. Fixed and random effects estimates without 569 these points are also included in Table 3, while Figure 6 shows effect sizes for QOL studies. 570 Overall, QOL studies indicated a negligible to small effect that showed a non-significant 571 decrease in quality of life as a result of expressive writing. 572

Homogeneity. For QOL studies including outliers, we found significant heterogeneity from our random effects model, $Q(36)=200.09,\ p<.001$ and $I^2=82.0,\ 95\%$ CI[75.9 - 86.5]. After excluding outliers, our random effects model still indicated heterogeneity, $Q(34)=93.18,\ p<.001$ and $I^2=63.5,\ 95\%$ CI[47.6 - 74.6].

Power. In conducting post hoc power using sample and effect size statistics from 577 individual studies, we found that 21.6% of studies would have been considered significant at 578 $\alpha < .05$. Average power based on actual study characteristics was .33 (SD = .32). Power for 579 the random effects meta-analytic effect size with outliers was .05 (SD = .00) and without 580 outliers was .05 (SD = .00). Unfortunately, power was around 5% for both random effects 581 models with and without outliers. In these studies, 18.9% achieved adequate power of 80% 582 on their found effect size, while 0.0% achieved 80% power for our random effects model with 583 outliers. Finally, without outliers, 0.0% achieved 80% power. 584

Other Meta-Analytic Estimates. We exert caution in interpreting p-curve and p-uniform analyses on QOL outcomes with and without outliers due to heterogeneity. As seen in Table 1, p-uniform estimates were stronger and positive than other techniques because of the high degree of heterogeneity recently described. p-curve pictures can be found at the following OSF Link: https://osf.io/4mjqt. Eight studies were significant at $\alpha < .05$,

and the studies indicated evidentiary value, p = .004. p-uniform analyses did not indicate 590 publication bias, Z = -2.75, p = .997. In PET-PEESE analyses, we found that the intercept 591 was not significant, and therefore, PET was a better estimator of the meta-analytic effect. 592 Table 1 indicates that both of these analyses estimate the effect size around zero, with a 593 confidence interval that includes zero. Selection models correspondingly show small effects 594 crossing zero, except for random effects models with outliers, that appear to be heavily 595 influenced by the outliers. Trim and fill models are shown in Table 3, and figures are 596 included online. No studies were imputed for these analyses, and therefore, the effect size 597 estimates match the original meta-analysis. Overall, these results appear to point to no 598 effects, ranging across zero with several negative estimates. Interestingly, the correlation of 599 effect sizes across measurement times with outliers was $r=-.37,\,95\%$ CI $[-.62,\,-.05],$ 600 t(35) = -2.33, p = .026 and r = -.64, 95% CI [-.80, -.39], t(33) = -4.75, p < .001 without outliers. The effect of expressive writing appears to be positive at short time intervals and decreases into negative effects at longer time intervals.

Discussion

In examining pre- to post-test comparisons across each variable separately, we found 605 that PTS studies indicated a small effect size across all meta-analytic estimates. 606 Interestingly, those studies requiring a diagnosis of PTSD for inclusion resulted in a medium 607 effect size, while those studies not requiring a PTSD diagnosis resulted in a small to medium 608 effect size. These results suggest that those with clinical symptoms of PTSD may benefit 609 more from expressive writing interventions. Further, these results are in contrast to recently-conducted studies, which suggest that those with subclinical symptoms benefit more from expressive writing tasks (Di Blasio et al., 2015; Sloan et al., 2011). Both QOL and 612 PTG studies indicated a negligible to small effect size using random effects models. 613 Although the PTG effect in our overall meta-analysis estimate was significant, other methods 614 indicate this small effect is likely not different from zero. We also examined the relationship 615

of time between measurements of the dependent variables and the corresponding effect size to determine if effects change over time. For both PTS and PTG, there was no relationship between effect size and time. For QOL studies, a medium to large negative correlation was found. A negative relationship between time and effect size implies that writing tasks were more effective in the initial time points, and effects decreased over longer time spans.

The authors note several limitations. Generally, ineffective emotional expression may 621 be a contributing factor. If participants/clients are not deeply engaged with the material, an 622 expressive writing task may not be effective, as Pennebaker and Graybeal (2001) imply that 623 connectedness is an important factor for the task. However, it may be difficult to implement 624 a check for engagement in these types of research designs. Doing so may also set a context 625 that will inhibit emotional processing and general responses. Research on expressive writing 626 has found a wide range of outcomes for different variables (Frattaroli, 2006), and these 627 various results may explain the large heterogeneity found in this study. Encouragingly, we 628 did not find much evidence of publication bias, and therefore, these estimates may represent 629 a true population effect size. Regardless, methodology of expressive writing studies is 630 variable, as it is applied in different forms across different contexts. Ideally, it would be 631 possible to control for these varied instructions and protocols. However, this is simply not feasible, as most studies do not use measures that examine how engaged an individual is with the material. As such, this current meta-analysis sought to provide readers with a 634 global effect of expressive writing on the aforementioned outcome variables. More studies are 635 needed to examine potential moderating effects of participant engagement. 636

The authors also note limitations in regards to the specific outcome variables. The
nature of the construct of PTG makes it difficult to analyze rigorously. For example, on the
Posttraumatic Growth Inventory (commonly used to study PTG), one could respond 0 to
the item "I have a greater appreciation for the value in my own life" because they already
had a high level of appreciation in their life (i.e., ceiling effect). This conceptual issue may
account for the non-effect of expressive writing on PTG. Logically, it would be difficult to

determine whether or not an individual experiences growth from trauma without having experienced trauma. In conducting the literature search for the present meta-analysis, an 644 insufficient number of studies requiring a diagnosis of PTSD employed PTG as an outcome 645 variable. Thus, it was difficult to determine whether participants in the studies employed 646 had experienced trauma in line with DSM-IV criteria. For PTS, studies not specifying 647 whether or not participants had a diagnosis of PTSD were included. It is possible that 648 studies included in the subclinical symptom category did in fact include participants without 649 PTSD diagnosis (perhaps it was simply not assessed by means of a structured clinical 650 interview). It is also crucial to consider mainstream issues not specific to expressive writing 651 and the outcome variables utilized in the present study. 652

The psychological scientific community has shifted focus to reproducibility and 653 research design in the last several years (Nelson, Simmons, & Simonsohn, 2018), and much of 654 this discussion has focused on adequately powering studies for publication (Bakker et al., 655 2016; Maxwell, Lau, & Howard, 2015). Maxwell et al. (2015) and Open Science 656 Collaboration (2015) have shown that the "replication crisis" may be attributed to low power 657 in published studies. The power found in the current meta-analysis was very poor, with very 658 few studies reaching the suggested 80% criterion to adequately power their study. This result was the same when considering individual study characteristics or the estimate true population effect size. Research by Bakker et al. (2016) indicates that researchers' intuitions 661 about power are particularly poor, and many studies could benefit from more informed 662 power analyses. Anderson, Kelley, and Maxwell (2017) recently published a primer on power, 663 with an online application to help with sample size planning for many types of research 664 designs. Additionally, we encourage researchers to report power analyses of studies in order 665 to better understand methodology for replication and reproducibility. 666

Meta-analyses, while useful tools to pool for population effect sizes, contain various limitations to their usefulness (van Elk et al., 2015). As mentioned previously, these analyses can be affected by high heterogeneity, which was found in this study (van Aert et al., 2016).

Selection models have been criticized when using a smaller number of studies (van Assen et 670 al., 2015), and trim and fill analyses may not always estimate accurate confidence intervals 671 and funnel plots may be biased with heterogeneity (Terrin, Schmid, Lau, & Olkin, 2003). 672 When focusing on improving the psychological sciences, van Elk et al. (2015) suggest that 673 the reliability and size of effects may be best elucidated by conducting large preregistered 674 studies. This suggestion will also improve the outlook for power in published studies, and 675 projects such as Many Labs can aide in subsidizing large samples (Klein et al., 2014). Even 676 with limitations, meta-analyses allow researchers to examine the state of a research area, and 677 we find potential with expressive writing on reducing PTS symptoms, and an overall need for 678 better sample size and power planning for studies. 679

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Model	Fixed Effects	Random Effects	Fixed No Outliers	Random No Outliers
Overall Effects	0.36 [0.34, 0.39]	0.42 [0.35, 0.49]	0.36 [0.33, 0.38]	0.40 [0.33, 0.46]
Z Values	24.64, p < .001	12.35, p < .001	23.97, p < .001	12.38, p < .001
<i>p</i> -Uniform	0.63 [0.54, 0.72]	-	$0.61 \ [0.52, \ 0.70]$	-
PET	0.09 [0.01, 0.18]	-	$0.14 \ [0.06, \ 0.22]$	-
PEESE	0.24 [0.20, 0.29]	-	$0.26 \ [0.22, \ 0.31]$	-
Selection Models	0.33 [0.28, 0.37]	$0.45 \ [0.33, \ 0.57]$	$0.29 \ [0.24, \ 0.33]$	$0.39 \ [0.27, \ 0.50]$
Trim and Fill	$0.28 \ [0.25, \ 0.31]$	0.28 [0.21, 0.36]	$0.28 \ [0.25, \ 0.31]$	$0.28 \ [0.21, \ 0.35]$

Note. [] indicates the 95 percent confidence interval for each effect size estimate.

 $\begin{tabular}{ll} Table 2 \\ Effect Size Estimates for PTG Results \end{tabular}$

Model	Fixed Effects	Random Effects	
Overall Effects	0.10 [0.02, 0.17]	0.10 [0.02, 0.17]	
Z Values	2.45, p = .014	2.45, p = .014	
<i>p</i> -Uniform	-0.11 [-1.43, 0.42]	-	
PET	0.06 [-0.20, 0.32]	-	
PEESE	0.08 [-0.04, 0.20]	-	
Selection Models	0.09 [-0.01, 0.18]	0.09 [-0.03, 0.20]	
Trim and Fill	$0.10 \ [0.02, \ 0.17]$	$0.10 \ [0.02, \ 0.17]$	

Note. [] indicates the 95 percent confidence interval for each effect size estimate.

Table 3 ${\it Effect Size Estimates for QOL Results}$

Model	Fixed Effects	Random Effects	Fixed No Outliers	Random No Outliers
Overall Effects	-0.01 [-0.07, 0.05]	-0.01 [-0.16, 0.13]	-0.01 [-0.07, 0.05]	-0.01 [-0.11, 0.09]
Z Values	-0.33, p = .745	-0.18, p = .860	-0.25, p = .805	-0.20, p = .838
<i>p</i> -Uniform	0.79 [0.33, 1.61]	-	$0.62 \ [0.10, \ 0.96]$	-
PET	0.05 [-0.26, 0.36]	-	0.05 [-0.29, 0.38]	-
PEESE	0.00 [-0.17, 0.17]	-	0.00 [-0.19, 0.19]	-
Selection Models	-0.06 [-0.12, 0.01]	0.51 [-0.09, 1.12]	-0.04 [-0.11, 0.03]	$0.05 \ [-0.15, \ 0.24]$
Trim and Fill	-0.01 [-0.07, 0.05]	-0.01 [-0.16, 0.13]	-0.01 [-0.07, 0.05]	-0.01 [-0.11, 0.09]

Note. [] indicates the 95 percent confidence interval for each effect size estimate.

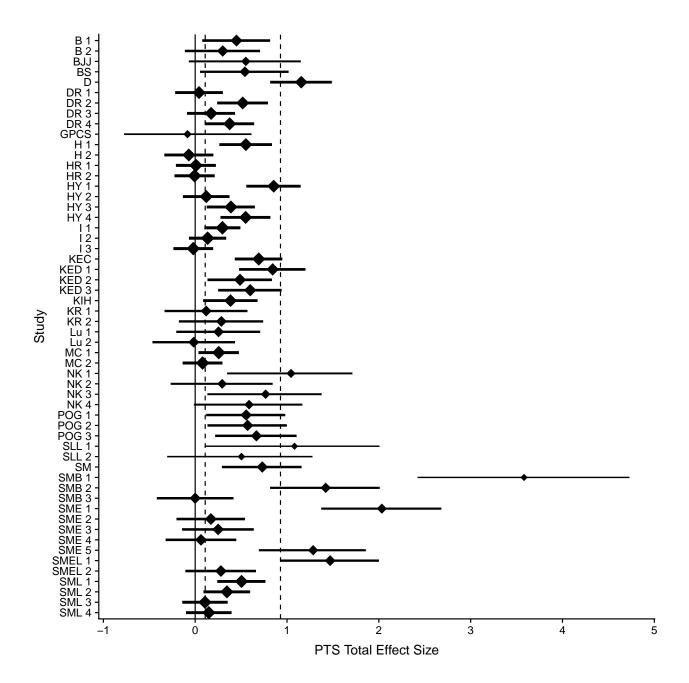


Figure 1. Effect sizes and their non-centralized confidence interval for PTS total scores. Dashed lines indicated average non-weighted lower and upper confidence interval limits. Diamond size indicates normalized study weight from a random effects model.

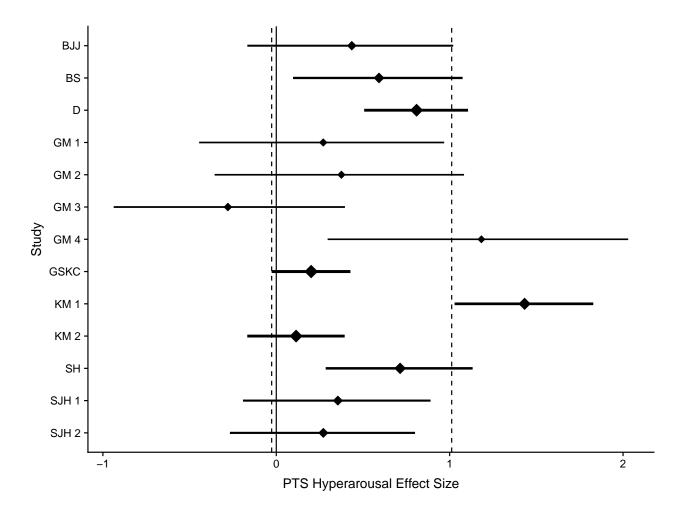


Figure 2. Effect sizes and their non-centralized confidence interval for PTS Hyperarousal. Dashed lines indicated average non-weighted lower and upper confidence interval limits. Diamond size indicates normalized study weight from a random effects model.

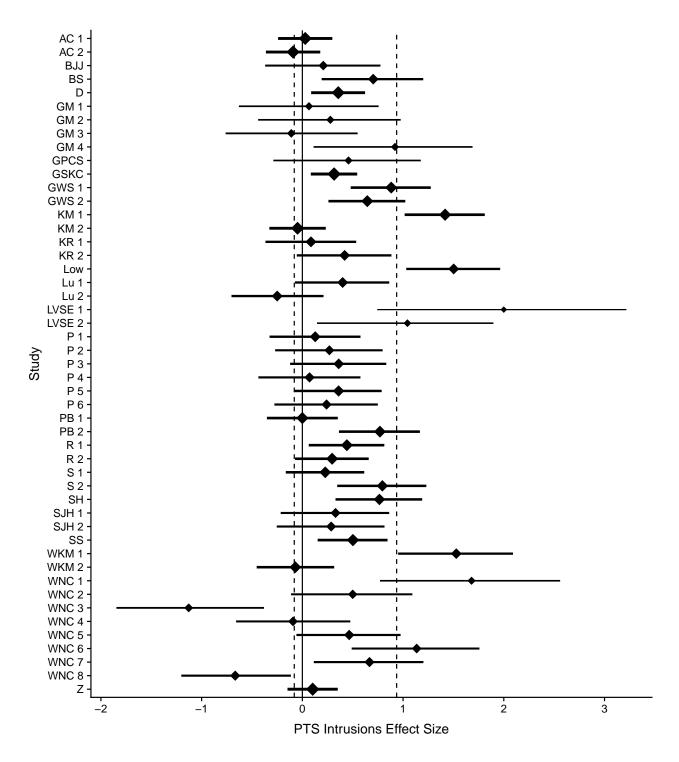


Figure 3. Effect sizes and their non-centralized confidence interval for PTS Intrusion scores. Dashed lines indicated average non-weighted lower and upper confidence interval limits. Diamond size indicates normalized study weight from a random effects model.

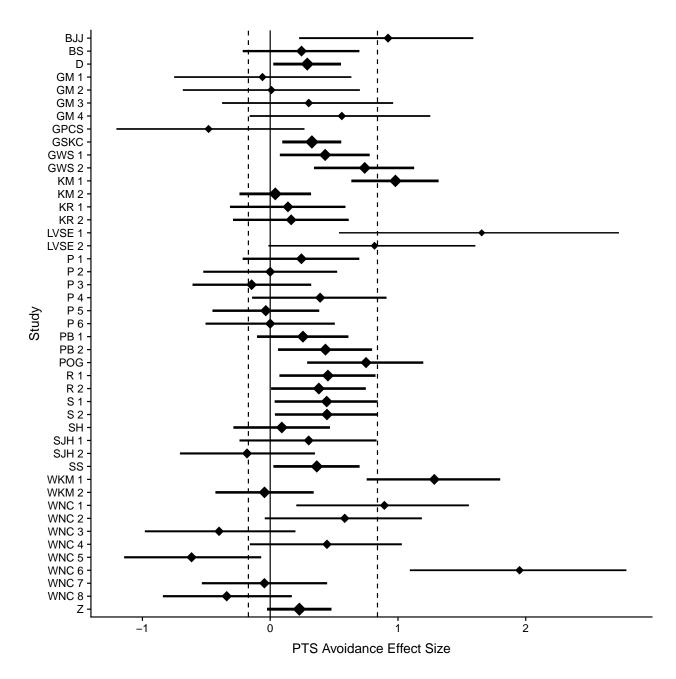


Figure 4. Effect sizes and their non-centralized confidence interval for PTS Avoidance Scores. Dashed lines indicated average non-weighted lower and upper confidence interval limits. Diamond size indicates normalized study weight from a random effects model.

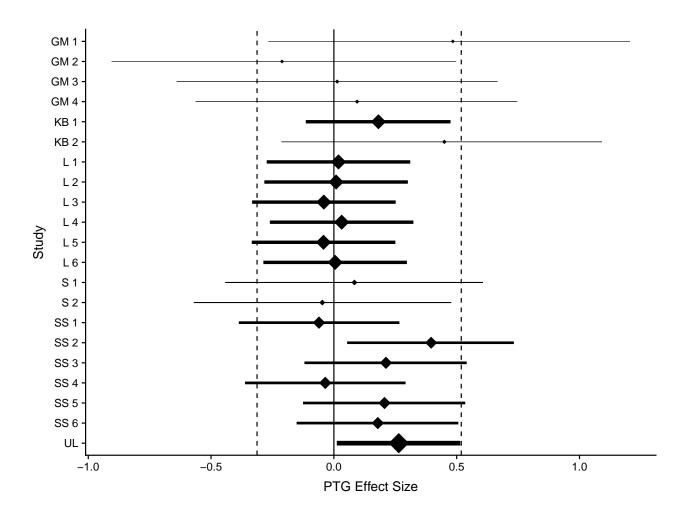


Figure 5. Effect sizes and their non-centralized confidence interval for PTG outcome variables. Dashed lines indicated average non-weighted lower and upper confidence interval limits. Diamond size indicates normalized study weight from a random effects model.

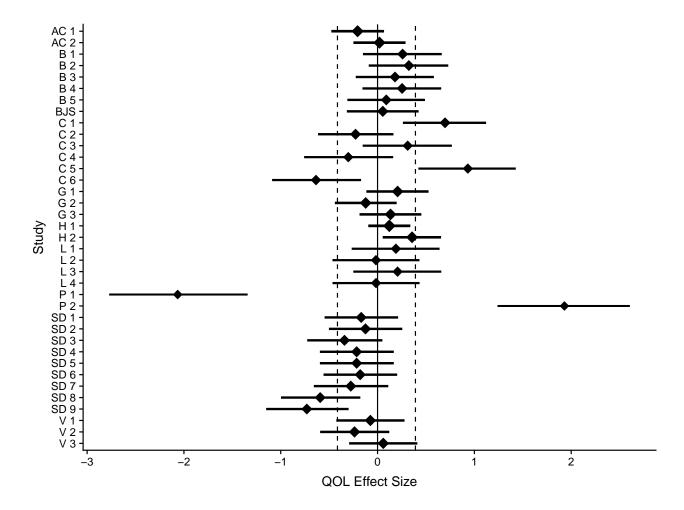


Figure 6. Effect sizes and their non-centralized confidence interval for QOL outcome variables. Dashed lines indicated average non-weighted lower and upper confidence interval limits. Diamond size indicates normalized study weight from a random effects model.