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Context and explicit threat cue modulation of the startle reflex: Preliminary evidence of distinctions between adolescents with principal fear disorders versus distress disorders

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Abstract

Anxiety and depression are prevalent, impairing disorders. High comorbidity has raised questions about how to define and classify them. Structural models emphasise distinctions between “fear” and “distress” disorders while other initiatives propose they be defined by neurobiological indicators that cut across disorders. This study examined startle reflex (SR) modulation in adolescents with principal fear disorders (specific phobia; social phobia) ($n = 20$), distress disorders (unipolar depressive disorders, dysthymia, generalized anxiety disorder; post-traumatic stress disorder) ($n = 9$), and controls ($n = 29$) during (a) baseline conditions, (b) threat context conditions (presence of contraction pads over the biceps muscle), and (c) an explicit threat cue paradigm involving phases that signalled safety from aversive stimuli (early and late stages of safe phases; early stages of danger phases) and phases that signalled immediate danger of an aversive stimulus (late stages of danger phases). Adolescents with principal fear disorders showed larger SRs than other groups throughout safe phases and early stages of danger phases. SRs did not differ between groups during late danger phases. Adolescents with principal distress disorders showed attenuated SRs during baseline and context conditions compared to other groups. Preliminary findings support initiatives to redefine emotional disorders based on neurobiological functioning.

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Keywords

Fear disorders; Distress disorders; Startle reflexes; Adolescents

1. Introduction

Anxiety and depression during adolescence are highly prevalent emotional disorders that cause significant concurrent and long-term impairment and economic burden (Bittner et al., 2007; Verduin and Kendall, 2007; Mathews et al., 2011). Furthermore, there is high comorbidity across the life span among anxiety disorders, and between anxiety and depressive disorders (e.g., Kashani and Orvaschel, 2000; for a review, see Craske and Waters, 2005). Such comorbidity has raised questions about how best to classify and define psychiatric disorders, including proposed revisions to the Diagnostic and Statistical Manual of Mental Disorders (DSM-IV-TR) and the International Classification of Diseases (ICD-10). While the DSM and ICD generally emphasise self-reported or observable behaviours and emotional features as criteria for individual disorders, the aim of the recent Research Domain Criteria (RDoC) initiative of the National Institute of Health is to characterize psychiatric disorders in terms of neurobiological indicators that cut across disorders as traditionally defined (Insel and Cuthbert, 2009; Craske, 2012).

Comorbidity between anxiety and depressive disorders has been explained in various models of psychiatric disorders as reflecting a broad “internalizing” factor (Kendler et al., 2003; Watson, 2005; Krueger and Markon, 2006; Lahey et al., 2008; Seeley et al., 2011) which subsumes two related sub-factors: “fear” and “distress” disorders (e.g., Clark and Watson, 2006; Prenoveau et al., 2010). Specific phobia, social phobia, separation anxiety disorder, panic disorder, and agoraphobia form the “fear” disorders, while generalised anxiety disorder (GAD), depression, post-traumatic stress disorder (PTSD), dysthymia, and possibly obsessive-compulsive disorder (OCD) form the “distress” disorders (Clark and Watson, 2006; Prenoveau et al., 2010). The major sources of evidence for the distinction between fear and distress disorders have come from self-report and genetic data (e.g., Kendler, 1996). However, more recent evidence emerging from cognitive science suggests that fear disorders may be characterised by attention biases away from threat (i.e., threat avoidance) while anxiety-related distress disorders (i.e., GAD) are characterised by attention biases towards threat (i.e., threat vigilance) (e.g., Salum et al., 2013; Waters et al., 2014b). Moreover, recent reviews of event-related negativity and eye blink startle reflex (SR) data as a function of anxiety and depressive disorders (see below) have revealed distinct neurophysiological indicators that show some consistency with structural models (e.g., Vaidyanathan et al., 2012).

The SR is a widely used psychophysiological index of human defensive responding. The SR involves the contraction of the orbicularis oculi muscle in response to a sudden, unexpected stimulus and is one of many obligatory somatic and visceral changes that comprise the human startle reaction (Graham, 1979). A variety of procedures have been used to examine SR modulation in threatening emotional states, including aversive picture-viewing and imagery experiments, as well as fear-potential paradigms when SR magnitudes have been assessed during explicit threat conditions (e.g., a cue signalling immediate threat of shock),

when participants anticipate shocks, and during safe conditions that predict the absence of shocks. “Baseline” SR magnitudes have also been assessed either before and after fear-potential paradigms when threat of shock was explicitly absent or during non-cued phases throughout experiments, while context-potentiated SR refers to amplified SRs when a cue of an upcoming or unpredictable threat may be present.

Several studies have reported that individuals diagnosed with phobias (e.g., animal phobias; non-generalized social phobia), or with high levels of phobic symptoms or trait fearfulness, demonstrate greater SR magnitudes during aversive picture-viewing or imagery while showing relatively normal baseline SR magnitudes (de Jong et al., 1991; Vrana et al., 1992; Globisch et al., 1999; Cuthbert et al., 2003; Lang et al., 2007; McTeague et al. 2009; 2012; Vaidyanathan et al., 2009a). Inconsistent results have been found for panic disorder (Grillon et al., 1994; Vaidyanathan et al., 2009b; McTeague et al., 2011) and generalised social phobia (McTeague et al., 2009), suggesting they may be characterised by more generalised SR activation similar to the broader and more chronic anxiety disorders discussed below. In studies utilising fear-potential paradigms, SR may be larger overall in some fear disorders, such as panic disorder, throughout fear-potential paradigms (suggestive of anticipatory apprehension) but not during baseline phases before and after fear-potential paradigms when threat is explicitly absent (Grillon et al., 1994). Few studies of circumscribed fears, such as specific phobias and non-generalized social phobia have utilised fear-potential paradigms. However, the fear conditioning literature suggests that fear disorders, such as panic disorder and social phobia, are characterised by overgeneralized defensive responding to stimuli that are safe (i.e., CS–) but are associated with danger cues (i.e., CS+) (see Lissek et al., 2009; see Lissek, 2012, for review).

In contrast, distress-related anxiety disorders such as PTSD, OCD and GAD are more consistently associated with elevated baseline SR magnitudes and amplified context-potentiated SR relative to healthy controls (Morgan et al., 1995; Grillon et al., 1996; Kumari et al., 2001; Cuthbert et al. 2003; Grillon et al., 2008; Ray et al. 2009; Pole et al., 2009). Results regarding fear-potentiated SR have been more mixed (e.g., Kumari et al. 2001; Cuthbert et al., 2003; Kaviani et al., 2004; Lang et al., 2007). PTSD and GAD have also been linked to overgeneralized defensive responding to safe stimuli (i.e., CS–) in conditioning experiments (see Lissek, 2012; Lissek et al., 2013), and SR results for PTSD in particular have been considerably mixed compared with findings for other anxiety disorders (Grillon & Baas, 2003; Pole, 2007; Vaidyanathan et al., 2009b).

Other studies have shown that depression-related distress disorders have an attenuating effect upon affective and fear-potentiated SR relative to healthy controls (Allen et al., 1999; Kaviani et al., 2004; Forbes et al., 2005; McTeague et al., 2009) and anxious samples without depression (Melzig et al., 2007). Moreover, SR modulation during emotional picture viewing in anxiety disorders is blunted by comorbid depression (Taylor-Clift et al., 2011). Depression has also been linked to diminished physiological reactivity during the anticipation of threat in conditioning experiments in offspring of mothers with a principal depressive disorder relative to offspring of mothers with a principal anxiety disorder and low risk offspring (Waters et al., 2014a). These findings could suggest that depressive disorders are associated with blunted or context insensitive emotional responding (Rottenberg et al.,

2005). However, other research has found that depressive disorders are characterised by elevated SR throughout fear-potential experiments, contextual anxiety (i.e., placement of shock electrodes) and when shock is predictable (Grillon et al., 2013), while other findings suggest that depressive disorders have no additional effect on the elevated SR magnitudes of adults with panic disorder relative to controls during fear-potential experiments (Shankman et al., 2013). Differences in findings may be due to variation in methodology. Blunting in depression has been obtained during viewing of pictures and films, or emotional imagery (Allen et al., 1999; Taylor-Clift et al., 2011; McTeague et al., 2012) or in anticipation of aversive tones (Waters et al., 2014a). Thus, threat is mild, imaginary, and/or may lack personal relevance (Grillon et al., 2013). Therefore, it may be adaptive for depressed individuals to disengage from these types of threats but not from more explicit danger such as a shock in fear-potential paradigms (Nesse, 2000; Grillon et al., 2013; Shankman et al., 2013).

As is clear, wide variation in methodology exists across studies. Furthermore, experimental protocols that separate baseline, context and explicit threat phases might help to clarify the conditions under which neurophysiological markers cut across versus differentiate fear-related and distress-related disorders (Watson, 2005; Insel and Cuthbert, 2009; Seeley et al., 2011; Craske, 2012). Therefore, the current study utilised a SR modulation protocol involving a baseline condition when threat of an aversive muscle contraction was explicitly absent, a context condition when a cue of later threat was present (i.e., presence of muscle contraction pads over the biceps muscle), and an explicit threat cue paradigm involving phases that signalled safety from aversive stimuli (i.e., early and late stages of safe phases; early stages of danger phases) and phases that signalled immediate danger of an aversive stimulus (i.e., late stages of danger phases).

If fear disorders are characterised by overgeneralized defensive responding to safe stimuli associated with threat (e.g., Grillon et al., 1994; Lissek et al., 2009), then adolescents with a principal fear disorder (i.e., either specific phobia or non-generalised social phobia) were expected to show amplified SRs during safe phases of the explicit threat cue paradigm compared to healthy controls and adolescents with principal distress disorders. However, groups were not expected to differ in SR magnitudes in response to immediate danger of an aversive stimulus (i.e., late stages of danger phases), reflecting a biologically imperative defence response to explicit threat (Lissek et al., 2005; Craske et al., 2009). In contrast, if distress disorders are associated with elevated SRs during baseline and context conditions (i.e., placement of shock electrodes) within experiments involving fear-potential paradigms (Grillon et al., 1996; Pole et al., 2009; Grillon et al., 2013), i.e., the explicit threat cue paradigm in the present study, then adolescents with a principal distress disorder (i.e., either GAD, PTSD or a depressive disorder) were expected to show larger SR magnitudes during the baseline and context conditions in comparison with controls and those with a principal fear disorder.

2. Methods

2.1 Participants

Participants were high school juniors from schools in suburban Chicago, Illinois, and suburban Los Angeles, California who participated in the Northwestern University–University of California, Los Angeles (NUCLA) Youth Emotion Project (YEP) (see Craske et al., 2009 and Zinbarg et al., 2010 for further details on recruitment and overall study design). Of 1269 students who completed an initial screening measure of neuroticism, 627 participants (69% female) had parental consent and gave youth assent to participate in the project. They all completed a baseline diagnostic assessment and 185 of these participants completed the startle modulation experiment. Of these, 132 did not meet criteria for any psychiatric disorder and the effects of neuroticism upon SR modulation are reported in Craske et al. (2009). Of the remaining 53 participants, 29 met criteria for one or more emotional disorders (19 girls; 10 boys). The remaining 24 participants met criteria for other psychiatric disorders, or did not have usable SR data, and are not included in this study.

Thus, the present study compared SR data from the 29 participants with emotional disorders with SR data from 29 healthy controls matched for gender and age (in years) selected from the 132 participants without a psychiatric disorder originally reported in Craske et al. (2009). There were no significant differences in any socio-demographic, self-report or SR measures between the 29 control children and the larger sample from which they were selected (i.e., Craske et al., 2009). Of the 29 participants with emotional disorders, 20 had a current principal (i.e., most severe) fear disorder of either specific phobia ($n = 7$) or social phobia ($n = 13$). The other nine of the 29 participants had a current principal (i.e., most severe) distress disorder; five had a principal unipolar depression-related distress disorder (MDD), dysthymia, minor depressive disorder) and four had a principal anxiety-related distress disorder (GAD, PTSD). The 29 participants with emotional disorders had an average of 1.69 diagnoses ($SD = 0.85$), all of which were emotional disorders. Thus, participants in the Distress and Fear Disorder groups did not meet criteria for psychiatric disorders other than anxiety and depressive disorders. None were on medication or receiving psychological treatment at the time of assessment. See Table 1 for patterns of comorbidity.

Age and ethnic composition of the total sample ($n = 58$) were similar to the larger sample (see Craske et al., 2009; 2012); 50% of the participants were Caucasian, 14% Hispanic/Latino American, 12% African American, 7% Asian American/Pacific Islander, and 17% “other” or multi-ethnic. Age ranged from 16 to 18 years (mean=17.0, $SD=0.35$). Participants received monetary compensation for their time and transportation costs.

2.2 Materials

2.2.1. Diagnosis—Lifetime and current Axis I psychopathology was assessed using the Structured Clinical Interview for DSM-IV (SCID-Non-Patient Version) (First et al., 2002). After completing interviews, interviewers rated the severity of each current diagnosis in the past month using the Di Nardo and Barlow (1988) 0 to 8 clinician-severity rating (CSR) scale, in which scores of 4 or above indicate clinically significant impairment or distress have been present for the past month. The 29 participants with emotional disorders all met

diagnostic criteria for one or more current emotional disorders. Diagnostic reliability was adequate to good based on the larger YEP sample by having trained interviewers observe live SCIDs ($n = 69$) ($\kappa = 0.65 - 0.83$).

2.2.2 Symptoms—Anxiety and depression symptoms were assessed using the Multidimensional Anxiety and Mood Questionnaire which consists of five subscales: Mixed Symptoms, Anxiety Symptoms, Depressive Symptoms, Anxious Arousal, Anhedonic Depression (Watson et al., 1995a; 1995b). Each of the five scales had good internal consistency ($\alpha > 0.84$) (see Sutton et al. 2011).

2.2.3 Electrophysiological equipment and data acquisition—The equipment and data acquisition were the same as in Craske et al. (2009). Auditory startle stimuli (105 dB, zero rise time, 50-ms white noise bursts) were presented binaurally through stereophonic headphones (Sony, Model MDRV700). The muscle contraction, delivered by a Digital 807 Electrical Muscle Stimulation Device (Everyway Medical Instruments Co.), was a 20.4 mA peak current (i.e., equating to 50 V peak) for 0.5 s. The intensity level was pre-set on the basis of pilot testing to represent an uncomfortable but not painful intensity but was similar to mean voltage levels of shock intensity in studies using shock work-up procedures (Neumann and Waters, 2006); individualized work-up procedures were not chosen, because pre-exposure to the muscle contraction might have decreased anticipatory anxiety during the explicit threat cue paradigm and/or weakened its aversiveness due to habituation (Baker et al., 1981). Startle reflex was measured by electromyogram (EMG) activity of the orbicularis oculi (see Craske et al., 2009, for further details).

2.2.4 Subjective Ratings—Anxiety was rated on a 10-point Likert scale (“calm and relaxed” to “really nervous or scared”) after each baseline and context condition and the explicit threat cue paradigm. After the experiment, participants rated the intensity and unpleasantness of the biceps contraction on 20-point scales (higher scores reflecting higher values).

2.3 Procedure

Data from 25 participants at UCLA and 33 participants at Northwestern University (NU) were used in this study. The two laboratories used identical hardware, software, manualized procedures, and technician training procedures.

Detailed information on the experimental procedure (including a Figure depicting the experimental conditions, the timing of startle probes and the muscle contraction) is reported in Craske et al. (2009). Briefly, after a 5-min *resting period* for adaptation, participants were presented with a single startle stimulus to reduce initial reactivity (discarded from analyses). During the *first baseline condition*, participants received 8 startle probes while focusing on a white fixation cross on the computer. For the *first context condition*, they were fitted with two contraction pads to the biceps muscle and told they would be informed when the contractions would happen later in the experiment. Eight more startle probes were presented while participants focused on the fixation cross. Before the *explicit threat cue paradigm*, participants were told there would be no muscle contractions delivered while the words

'Safe: no contraction will be given' were on the green screen, and they might receive a contraction when the words 'Danger: contraction may be given' were on the red screen. Participants were also told that for both phases, they would see a progressing bar showing the time from 0 to 55 s, and that if a biceps contraction occurred during a Danger phase, it would occur when the bar turned from pink to red in the last 15 s. They were then told they might receive a muscle contraction up to three times of increasing intensity each time. Participants received only one contraction in the final 15s of the fourth danger phase, half way through the paradigm. There were 8 safe and 8 danger phases in alternating order. A total of 32 startle probes were presented: 16 in the Danger and 16 in the Safe phases with two trials per phase at 5, 15, 35 or 45 s (the final startle probe presented during the final 15 s when threat of contraction was imminent in the Danger phase).

The muscle contraction pads were removed for the *second baseline condition*, and reattached for the *second context condition*. Following electrode removal, startle stimuli and muscle contractions were rated, hearing was tested (all participants passed), and participants were debriefed.

2.4 Response definitions and data analyses

Startle magnitudes were defined using conventional methods as described in Craske et al. (2009). In short, EMG magnitudes were expressed as the difference between the mean amplitude of the 200 ms of EMG preceding the startle stimulus and the peak response, in microvolts (μV). Analyses were performed on natural log ($\ln$) transformed eye blink data as per Craske et al. (2009) using a linear mixed model for repeated measurements with Satterthwaite's Approximation for degrees of freedom.

For baseline and context phases, the fixed effects were Group (Fear; Distress; CON), Block (Pre; Post Explicit Threat Cue Paradigm), Condition (Baseline; Context), and Trial Half (Trials 1 to 4; Trials 5 to 8). For the explicit threat cue paradigm, the fixed effects were Group, Block (Pre; Post Muscle Contraction), Phase (Safe; Danger), and Probe Time (Early; Late). Consistent with Craske et al. (2009), "Early" referred to 5-, 15-, and 35-s probe times and "Late" referred to the 45-s probe time, when the muscle contraction could occur within danger but not safe phases.

3. Results

3.1 Group comparisons

There were no significant group differences in age, $F(2, 55) = 2.02, p = 0.14, n_p^2 = 0.07$, ethnicity, $\chi^2(2, n = 58) = 0.15, p = 0.93$, gender, $\chi^2(2, n = 58) = 0.86, p = 0.65$, or number of participants recruited from each site, $\chi^2(2, n = 58) = 0.41, p = 0.81$. There were no significant differences in the number of diagnoses, $t(27) = 0.25, p = 0.85$, or the severity of principal diagnoses between the clinical groups, $t(27) = 0.22, p = 0.83$. One-way analyses of variance (ANOVAs) of the MASQ subscales revealed significant group differences on the Mixed, $F(2, 55) = 14.62, p < 0.001, n_p^2 = 0.34$, Anxiety, $F(2, 55) = 7.55, p = 0.001, n_p^2 = 0.22$, Depressive, $F(2, 55) = 20.7, p < 0.001, n_p^2 = 0.43$, Anhedonic, $F(2, 55) = 13.6, p < 0.001, n_p^2 = 0.33$, and Anxious Arousal subscales, $F(2, 55) = 5.90, p = 0.004, n_p^2 = 0.18$ (see Table 2). The Fear and Distress Disorder groups had significantly higher scores than

CONs on all subscales (all $p < 0.048$), but did not differ significantly from each other on any subscale (all $p > 0.98$) (see Table 2).

3.2 Baseline and context conditions

Analysis revealed significant main effects for Block, $F(1, 354) = 62.64, p < 0.001$, SRs were significantly smaller in the post- compared with the pre-explicit threat cue paradigm, Condition, $F(1, 997) = 4.33, p = 0.038$, SRs were significantly larger in the context compared with the baseline phases, Trial Half, $F(1, 1734) = 16.73, p < 0.001$, SRs were significantly larger in the first compared with the last half of trials, and Group, $F(2, 211) = 10.00, p < 0.001$, SRs in the Distress Disorder group were significantly smaller compared with the CON group ($p < 0.001$) and the Fear Disorder group ($p < 0.001$), which did not differ significantly from each other ($p = 0.63$) (see Fig. 1, upper panel). There was a significant Block $\times$ Condition interaction, $F(1, 663) = 5.65, p = 0.018$. The effect of Condition before the explicit threat cue paradigm was not significant, $F(1, 699) = 2.34, p = 0.10$, whereas SRs were significantly larger during the context compared with the baseline phase post-explicit threat cue paradigm, $F(1, 431) = 13.07, p < 0.001$.

3.3 Explicit threat cue paradigm

Analysis revealed significant main effects of Phase, $F(1, 1171) = 44.85, p < 0.001$, Block, $F(1, 466) = 8.47, p = 0.004$, Probe Time, $F(1, 1410) = 52.60, p < 0.001$, and Group, $F(2, 206) = 8.04, p < 0.001$, which were subsumed by significant Phase $\times$ Probe Time, $F(1, 1352) = 35.20, p < 0.001$, Block $\times$ Phase $\times$ Probe Time, $F(1, 1373) = 6.67, p = 0.01$ and Phase $\times$ Probe Time $\times$ Group interactions, $F(2, 1348) = 3.54, p = 0.029$ (see Fig. 1, middle panel).

The three-way interaction between Phase, Probe Time and Group was driven by the simple 2-way Phase $\times$ Group interaction being larger for late probe times than for early probe times. The simple two-way interaction was not significant for the early probe times, $F(1, 1220) = 1.95, p = 0.14$ (however, significant main effects for Phase, $F(1, 1217) = 11.92, p < 0.001$, and Group, $F(2, 173) = 8.23, p < 0.001$), whereas the simple two-way interaction was significant for the late probe times, $F(2, 365) = 3.12, p = 0.045$. As expected, the simple two-way interaction for the late probe times was driven by significantly larger group differences in the safe phases, $F(2, 62) = 3.47, p = 0.037$, than in the danger phases, $F(2, 57) = 0.81, p = 0.45$. At late probe times during safe phases, the Fear Disorder group had significantly larger SRs than the Distress Disorder and CON groups (both $p < 0.036$).

3.4. Subjective ratings

Subjective anxiety ratings¹ differed significantly according to Condition, $F(4, 49) = 29.88, p < 0.001, \eta_p^2 = 0.71$, but not by Group, $F(2, 52) = 2.66, p = 0.07, \eta_p^2 = 0.09$, or in terms of a Group $\times$ Condition interaction, $F(8, 44) = 1.14, p = 0.20, \eta_p^2 = 0.05$. Anxiety was significantly higher during the explicit threat cue paradigm than all other conditions (all $p < 0.001$), and during the first baseline condition compared to the last baseline and context conditions (both $p < 0.015$) (see Table 2). There were no significant group differences in the ratings of the biceps contraction (intensity: $F(2, 55) = 2.12, p = 0.13, \eta_p^2 = 0.07$;

¹Subjective anxiety ratings were missing from one participant in the Fear Disorder Group and two participants in the CON group.

unpleasantness: $F(2, 55) = 0.40, p = 0.68, n_p^2 = 0.01$) or the startle stimulus (intensity: $F(2, 55) = 1.3, p = 0.29, n_p^2 = 0.04$; unpleasantness: $F(2, 55) = 0.19, p = 0.83, n_p^2 = 0.007$) (see Table 2).

4. Discussion

Consistent with hypotheses, adolescents with a principal fear disorder showed significantly *larger* SRs during safe phases and early danger phases of the explicit threat cue paradigm compared to healthy controls and adolescents with principal distress disorders. As expected, groups did not differ significantly in SR magnitudes during late danger phases when threat of the aversive contraction was immediate and all participants would be expected to show neurobiologically imperative defensive responses (Lissek et al., 2006; Craske et al., 2009). Contrary to expectations, adolescents with a principal distress disorder displayed *attenuated* SRs compared to controls and the fear disorder group during baseline and context conditions before and after the explicit threat cue paradigm.

The finding of amplified defensive responding during safe stages of the explicit threat cue paradigm is broadly consistent with evidence from the few previous studies of fear-related disorders employing fear-potential protocols that have found larger SR during but not before and after the fear-potential protocol and overgeneralized threat responding to stimuli resembling threat (i.e., CS-) in conditioning experiments (Grillon et al., 1994; Lissek et al., 2009; Lissek, 2012). Moreover, the finding that elevation in SR magnitudes in the fear disorder group emerged only during the explicit threat cue paradigm and quickly dissipated by the final baseline and context conditions highlights that phasic defensive responding to safe cues that are associated with threat is a highly time-locked state in fear disorders, leading to rapid dissipation of fear when threat is not explicit (Davis et al., 1989; de Jongh et al., 2003).

On the other hand, distress-related anxiety disorders such as GAD, PTSD and affiliated traits in the domain of negative affectivity have been associated in past research with larger baseline and context-potentiated SR (e.g., Kumari et al. 2001; Cuthbert et al. 2003; Grillon et al. 2008; Ray et al. 2009). Moreover, they have been less reliably associated with elevated fear-potentiated SR (see Kumari et al. 2001; Kaviani et al. 2004; Lang et al. 2007).

However, in contrast to expectations, we observed *attenuated* rather than amplified SR during baseline and context conditions both before and after the explicit threat cue paradigm in the distress disorder group. Consistent with some prior research (e.g., Allen et al., 1999; Kaviani et al., 2004; Forbes et al., 2005; Lang et al., 2007; Taylor-Clift et al., 2011; Waters et al., in press), this result could be due to the inhibiting effect of depressive disorders upon SR magnitudes; 5 of the 9 adolescents in this group had a principal diagnosis of a depressive disorder. However, Grillon et al. (2013) found enhanced rather than attenuated context-potentiated SR (i.e., placement of shock electrodes) in depression. As the aversive stimulus in the present study was an unpleasant “muscle contraction” rather than “shock”, threat following electrode placement may have been milder or more ambiguous than for shock, thereby attenuating SR due to disengagement/withdrawal (Grillon et al., 2013). The small sample of adolescents with a mix of anxiety and depressive disorders prevented potential distinctions in SR between anxiety-related and depression-related distress disorders from

being examined (e.g., see Vaidyanathan et al., 2012). Nevertheless, SR in the distress disorder group did not differ significantly from controls during the explicit threat cue paradigm; thus, the blunting effect of distress disorders on SR appeared to diminish when threat was explicit², consistent with the suggestion that emotional blunting in depression does not generalize to situations where strong defensive responses are evoked by actual threats (Grillon et al., 2013). In accord, we have shown previously that SR during safe phases of the explicit threat cue paradigm is a unique predictor of the onset of anxiety but not depressive disorders over an ensuing four year follow-up (Craske et al., 2012).

Unfortunately, due to the sample size in the present study, it was not possible to separate anxiety- from depression-related distress disorders to examine the effects of comorbid fear versus distress disorders or single diagnoses. Given that prior reviews of neurophysiological data suggest there may be some distinctions between (a) phobic and fear disorders, (b) non-phobic anxiety disorders and negative affect, and (c) depressive disorders (see Vaidyanathan et al., 2012), larger studies that can separate these diagnostic categories will be important for informing structural models of internalizing psychopathology (e.g., Watson and Clarke, 2006; Lahey et al., 2008; Seeley et al., 2011), as well as current initiatives, such as the RDoC, which aim to redefine the classification of emotional disorders along neurobiological lines (e.g., Insel and Cuthbert, 2009; Vaidyanathan et al., 2009; Craske, 2012). In contrast to studies of anxious adults (McTeague et al., 2012) but similar to studies with anxious children (Waters et al., 2014b), the present study did not find significant differences in the number of diagnoses or self-report symptom severity between the fear and distress disorder groups. These differences could reflect on developmental differences in the capacity to report on ones' own problems as well as the actual severity and impairment associated with emotional disorders that increases with advancing age. Nevertheless, that distinct patterns of SR modulation were observed in adolescents with principal fear versus distress disorders that cut across anxiety and depressive disorders as traditionally defined and exist in the presence of high rates of comorbidity suggests further research as a function of diagnostic category is warranted.

In summary, the present study found that adolescents with principal fear disorders showed elevated SRs during safe phases of an explicit threat cue paradigm, possibly reflective of overgeneralized defensive responding to safe stimuli under explicit threat conditions, whereas adolescents with principal distress disorders showed attenuated SRs during baseline and context conditions, perhaps reflective of withdrawal from the environment when threat is mild or not explicit. These findings support current initiatives to redefine emotional disorders in terms of patterns of neurobiological functioning and encourage further research along these lines.

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²Supplementary analyses comparing the seven fear disorder participants with comorbid depression with the remaining 13 fear disorder participants without comorbid depressive disorders revealed no significant group differences.

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Fig. 1.

Startle reflex magnitude during baseline and context phases prior to (left panel) during (middle panel) and after (right panel) the explicit threat cue paradigm as a function of diagnostic category (^ = significant Group main effect (collapsed across Block and Condition); * = significant Group main effect at Early Probe Times (collapsed across Safe and Threat Phases); # = significant interaction at Late Probe Times due to significant Group differences during Late Probe Times within Safe Phases).

Table 1

Summary of comorbid diagnoses as a function of diagnostic category

Principal diagnostic category (number of participants)	Comorbid diagnostic category (number of participants)	Type of comorbid diagnoses
Fear disorder (20)	Distress disorder-Dep only (4)	MDD
	Distress disorder-Anx only (3)	GAD, OCD
	Distress disorder-Dep & Anx (1)	GAD, MDD
	Distress disorder-Anx + Fear disorder (1)	PTSD, OCD, SP
	Distress disorder-Dep & Anx + Fear disorder (2)	PTSD, SP, Minor DD, GAD
	No comorbid diagnoses (9)	
Distress disorder (9)	Distress disorder-Dep + Fear disorder (1)	Dysthymia, SoP
	Fear disorder only (4)	SoP
	No comorbid diagnoses (4)	

Note: Dep = Depression, Anx = Anxiety; MDD = Major Depressive Disorder; GAD = Generalised Anxiety Disorder; OCD = Obsessive Compulsive Disorder; PTSD = Post Traumatic Stress Disorder; SP = Specific Phobia; Minor DD = Minor Depressive Disorder; SoP = Social Phobia

Table 2

Descriptive information as a function of diagnostic category

Measure	CON (<i>n</i> = 29)	Fear (<i>n</i> = 20)	Distress (<i>n</i> = 9)
Gender (male: female)	10:19	7:13	3:6
Age (years: months)	17.0 (.2)	16.9 (.4)	17.2 (.4)
Principal diagnosis severity		4.75 (1.02)	4.67 (.70)
Number of diagnoses		1.85 (.99)	1.66 (.70)
Contraction			
intensity	12.4 (4.3)	14.2 (3.9)	15.1 (2.1)
Unpleasantness	9.6 (4.3)	9.4 (4.1)	10.9 (3.3)
Startle stimulus			
intensity	10.6 (4.9)	12.4 (4.0)	12.2 (3.8)
Unpleasantness	8.5 (3.7)	9.1 (3.3)	8.4 (3.9)
MASQ			
Mixed	28.1 (9.1)	41.8 (11.3)	42.6 (8.1)
Anxiety	17.2 (6.5)	25.6 (7.3)	24.6 (8.9)
Depression	20.1 (7.4)	34.7 (11.7)	37.4 (8.0)
Arousal	23.2 (8.6)	32.8 (12.8)	32.7 (10.7)
Anhedonia	41.8 (18.3)	60.6 (18.1)	77.1 (11.99)
Anxiety ratings			
Baseline	2.8 (2.2)	4.1 (2.0)	4.2 (2.2)
Context	2.5 (2.1)	3.3 (2.1)	3.7 (1.9)
Explicit threat	4.3 (2.6)	6.4 (2.4)	5.1 (2.2)
Baseline	1.5 (1.6)	2.0 (1.4)	2.2 (1.9)
Context	2.0 (2.1)	3.0 (2.2)	2.3 (1.7)