Sex differences in fear regulation and reward seeking behaviors in a fear-safety-reward discrimination task

Abbreviated title: Sex differences in conditioned safety

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Number of figures: 4
Number of tables: 1

Abstract

Reward availability and the potential for danger or safety potently regulates emotion. Despite women being more likely than men to develop emotion dysregulation disorders, there are comparatively few studies investigating fear, safety and reward regulation in females. Here, we show that female Long Evans rats do not suppress conditioned freezing in the presence of a safety cue, nor do they extinguish their freezing response, whereas males do both. Females were also more reward responsive during the reward cue until the first footshock exposure, at which point there were no sex differences in reward seeking to the reward cue. Darting analyses indicate females might be able to regulate this behavior in response to the safety cue, suggesting they might be able to discriminate between fear and safety cues but do not demonstrate this with conditioned suppression of the freezing behavior. However, levels of darting in this study were too low to make any clear conclusions. In summary, females showed a significantly different behavioral profile than males in a task that tests the ability to discriminate among fear, safety and reward cues. This paradigm offers a great opportunity to test for mechanisms that are generating these behavioral sex differences in learned safety and reward seeking.

Keywords: sex differences; fear; safety; reward

1. Introduction

Clinical disorders arising from maladaptive emotion regulation present a large burden on society worldwide. Many of these disorders show comorbidity, for example, addiction with anxiety disorders (Grant et al., 2016). Cues predicting something aversive elicit avoidance and fear behaviors whereas cues predicting reward elicit approach and reward-seeking behaviors. Cues signifying safety have the power to modulate fear and reward-seeking behaviors by informing the organism whether or not the environment is safe (Walasek, Wesierska, & Zieliński, 1995). Thus, safety, fear and reward behaviors, and the circuitries governing these behaviors, are intertwined. The majority of studies on reward and fear processing have been conducted in parallel, investigating the circuitries separately in primarily male subjects. If we hope to understand and treat comorbid disorders resulting from maladaptive emotion regulation, increased efforts in investigating how these circuitries integrate their functions to influence behavior is needed in both male and female subjects.

Our laboratory has designed and validated a behavioral task in which fear, safety and reward cues are learned within the same session allowing us to assess the animal's ability to discriminate among these cues (Ng et al., 2018; Sangha et al., 2013; Sangha, Greba et al., 2014; Sangha, Robinson et al., 2014). Rats are exposed to cues associated with safety, fear (fear cue paired with footshock), and reward (reward cue paired with sucrose). Male rats consistently learn to discriminate among safety, fear and reward cues to 1) suppress conditioned freezing in the presence of a safety cue (fear+safety cue), and 2) increase reward seeking when reward is available (reward cue) (Ng et al., 2018; Sangha et al., 2013; Sangha, Greba, et al., 2014; Sangha, Robinson, et al., 2014). This paradigm also allows us to investigate how safety cues can regulate both fear and reward behaviors. Evidence suggests that reward learning mechanisms overlap with safety learning (Leknes et al., 2011; Pollak et al., 2008; Rescorla, 1969; Rogan et al., 2005; Sangha et al., 2013; Tanimoto et al., 2004; Walasek et al., 1995). For example, learned safety can act as a behavioral antidepressant in mice (Pollak et al., 2008), and animals will perform certain behaviors in order to turn on a safety signal (Rescorla, 1969; Rogan et al., 2005). Within the amygdala we have shown a subpopulation of neurons responding with the same level of excitation or inhibition during both the reward and safety cues (Sangha et al., 2013). Additionally, we have shown inactivation of the prelimbic or infralimbic cortices of the ventromedial prefrontal cortex have differential effects on reward and safety discrimination, respectively (Sangha, Robinson, et al., 2014). Thus, in male rats, our prior work

has already shown a critical involvement of the corticoamygdalar circuit in learning this fear-safety-reward cue discrimination.

Much of the research investigating emotion regulation mechanisms have exclusively used male subjects. In a study using male Vietnam veterans, PTSD patients show impairments in suppressing their fear response in the presence of a safety cue (Jovanovic et al., 2009). But, women are more than twice as likely to develop Post-Traumatic Stress Disorder (PTSD) than men, with females having a lifetime prevalence of 8.5% in contrast to 3.4% in males (Mclean et al., 2011). In fear studies that have included female rats, it has been shown that females exhibit lower levels of freezing behavior than male rats after repeated fear cue presentations (Daviu et al., 2014). These findings have been thought to indicate a difficulty in fear conditioning learning in female rats. A more recent experiment has identified that approximately 40% of female rats tested exhibit an alternate fear behavior in the form of fast paced movements called ‘darting’; this was only seen in approximately 10% of male rats tested (Gruene et al., 2015). There is also evidence of sex differences in the seeking of natural rewards, where it has been reported that female rats consume more sucrose pellets than males and are willing to work harder for them (Tapia, Lee, Weise, Tamasi, & Will, 2019). Dopamine signaling during reward tasks has also been demonstrated to be different between sexes. For example, Conway et al (2019) showed females continue to perform intracranial self-stimulation for brain stimulation reward while under the influence of a kappa-opioid receptor agonist, which suppresses dopamine release, whereas males decrease this behavior. Their data suggest that female rats may have an increased capacity to produce and release dopamine compared to males, under these conditions. Our prior work has shown, in males, that dopamine signaling in the basolateral amygdala contributes to effective discrimination among fear, safety and reward cues (Ng et al., 2018).

Taken together, we hypothesized there would be sex differences in the ability to express clear discrimination among fear, safety and reward cues. The inability of male PTSD patients to learn safety signaling has been labeled a biomarker of the disorder (Jovanovic et al., 2012). Due to sex-related differences in human diagnosis of PTSD, with women diagnosed at rates twice that of men (Glover et al., 2015), any differences female rats have in the learning or retention of safety signals could steer towards further research on the neurological processes underlying these variations.

2. Materials and Methods

2.1 Subjects

A total of 24 adult male (215-375g) and 28 adult age-matched female (198-230g) Long Evans rats (Blue Spruce; Envigo, Indianapolis), were single-housed and handled for 1 week prior to testing. All procedures were performed during the light cycle and approved by the Purdue Animal Care and Use Committee. Rats had *ad libitum* access to food and water prior to the start of the experiment. After experiment onset, they were maintained on a food restricted diet (20g per day for males; 16g per day for females) until the last day of the experiment.

2.2 Apparatus

The rats were trained in operant conditioning chambers consisting of Plexiglas boxes (32cm length x 25cm width x 30cm height) encased in sound-attenuating chambers (Med Associates, ST Albans, VT). 10% liquid sucrose was delivered through a recessed port 2cm above the floor in the center of one wall. Two lights (28V, 100mA) were located 10.5cm from floor on either side of the port. A light (28V, 100mA) 27cm above the floor on the wall was on throughout the entire session. Auditory cues were delivered via a speaker (ENV-224BM) located 24cm from the floor on the same wall as the port. Footshocks were delivered through a grid floor via a constant current aversive stimulator (ENV-414S). An overhead video camera and side video camera recorded the sessions for subsequent offline video scoring.

2.3 Behavioral Procedures

<u>Reward pre-training</u> (5 sessions): An auditory cue is paired with 10% sucrose solution delivery (100µl) and serves as the reward cue (25 trials; ITI, 90-130s).

<u>Habituation</u> (1 session): Rats continue to receive 25 reward cue-sucrose pairings (ITI, 90-130s) in addition to 5 unreinforced presentations each of the future fear and safety cues in order to habituate the rats to their presentation, thereby reducing any baseline freezing to these novel cues.

<u>Discriminative conditioning (DC)</u> (4 sessions): Reward cue-sucrose pairings continue (15 trials). Another auditory cue is paired with a mild 0.5mA, 0.5s footshock and serves as the fear cue (4 trials). In separate trials the 20s fear cue is presented at the same time as a 20s safety light cue resulting in no footshock ('fear+safety', 15 trials). Trials in which the safety cue is presented alone without any footshock are also included to assess whether freezing develops to the safety cue as well as providing the animal with additional trials that contains a safety cue-no shock contingency (10 trials). Trials are presented pseudorandomly (ITI, 100-140 s). Cues were

145 counterbalanced across subjects with the caveat that the fear+safety compound cue was
146 composed of one auditory cue and one light cue. Eight of the male rats and 12 of the female
147 rats underwent DC training in which the reward cue was a continuous auditory cue (3 kHz, 20s
148 cue; 70dB), the fear cue a pulsing auditory cue (11 kHz, 20s; 70dB), and the safety cue was the
149 presentation of two lights (28V, 100mA located on both sides of the port). The remaining eight
150 male rats and eight female rats underwent training in which the fear and safety cue stimuli were
151 counterbalanced: the light served as the fear cue and the pulsing auditory cue served as the
152 safety cue.
153 Extinction Training (1 session): One day after the last DC session, both the reward cue and fear
154 cue were presented 20 times each in a pseudorandomized order without sucrose or footshock.
155 Extinction Test (1 session): One day after extinction training, rats were presented with the
156 reward (10 trials), fear (10 trials), fear+safety (5 trials) and safety (5 trials) cues in a
157 pseudorandomized order (ITI, 60-120s). None of the cues were presented with sucrose or
158 footshock.
159
160 To exclude possible sex differences in pain sensitivity and foot shock perception a separate
161 group of male (n=8) and age-matched female (n=8) rats was presented with a series of
162 unsignalled foot shocks of increasing intensities (0.3 mA, 0.35 mA, 0.4 mA, 0.45 mA, 0.5 mA,
163 0.55 mA, 0.6 mA, 0.7 mA, 0.8 mA, 0.9, 1.0 mA) with an inter-stimulus interval of 2 min. The
164 session was flanked with 5 min intervals in which no stimulus occurred.
165
166 *2.4 Data analyses*
167 Our experimental groups to directly compare males and females on discrimination behavior
168 consisted of 16-20 rats. Cohorts of 4 or 8 female rats were trained alongside cohorts of 4 male
169 rats for a total of 4 replications. Fear behavior was assessed manually offline from videos by
170 measuring freezing, defined as complete immobility with the exception of respiratory
171 movements, which is an innate defensive behavior (Blanchard & Blanchard, 1969; Fendt &
172 Fanselow, 1999). The total time spent freezing during each 20s cue was quantified and
173 expressed as a percentage. Measuring the total time the animal spent inside the reward port
174 and at the entrance of the port with nose positioned at port entrance during each cue assessed
175 reward seeking behavior and was expressed as a percentage. Darting behavior was detected
176 and quantified offline from videos recorded from overhead cameras via a custom MatLab
177 program, with movements of a velocity of 23.5cm/s or faster qualifying as a single dart (Gruene
178 et al., 2015); these were also confirmed manually. Darting was expressed as the averaged # of

179 darts per cue (sum of darts/ # trials) or trial (sum of darts). Since there were different number of
180 trials per reward, fear, fear+safety and safety cue in each DC session and test for extinction,
181 this was expressed as the sum of darts across trials divided by the number of trials for each cue
182 (sum of darts/ # trials). During extinction training, data for each individual trial is shown and the
183 dart rate is expressed the averaged sum of darts for each individual trial (sum of darts). Three
184 individuals performed manual offline behavioral scoring. Pearson's correlations of behavioral
185 values between scorers were greater than $r = 0.80$. Behavioral data were analyzed with one-
186 way or two-way repeated measures ANOVAs, with sex as the independent factor and condition
187 as the repeated factor, followed by *post hoc* Sidak's, Tukey's or Dunnett's multiple comparisons
188 tests with GraphPad Prism 8. P values were adjusted for multiple comparisons.
189
190 For shock sensitivity testing, freezing duration in the 2-min intervals between shock
191 presentations was scored manually as an indicator of fear, as well as darting and jumping as an
192 immediate shock response. For the freezing durations, a two-way repeated measures ANOVA
193 was carried out via GraphPad Prism 7, with sex as the independent factor and shock intensity
194 as the repeated factor. Darting and jumping were assessed as dichotomous variables with
195 darting/no darting and jumping/no jumping, respectively. For both, a Cochran test was
196 performed.
197
198 **3. Results**
199 *3.1 Female rats spent more time reward seeking during reward pre-training*
200 All rats first underwent 5 reward pre-training sessions in which the reward cue was paired with
201 sucrose delivery. The percent time spent at or in the reward port during each reward cue across
202 each reward session was quantified (Figure 1b). Two-way repeated-measures ANOVAs showed
203 main effects of session ($F(4,136)=5.395$, $p=0.0005$) and sex ($F(1,34)=10.83$, $p=.0023$), but no
204 significant interaction ($F(4,136)=0.9031$, $p=0.4641$). *Post hoc* Sidak's multiple comparisons test
205 showed females spent significantly more time reward seeking during the reward cue than males
206 for sessions R2 ($p=0.0274$), R3 ($p=0.0151$) and R5 ($p=0.0041$). The latency, in seconds, to
207 enter the port post-cue onset was also calculated for each reward cue presentation across all
208 sessions (Figure 1c). Two-way repeated-measures ANOVAs showed a main effect of sex
209 ($F(1,34)=20.37$, $p<.0001$), but no significant interaction ($F(4,136)=1.684$, $p=0.1571$) or main
210 effect of session ($F(4,136)=0.7755$, $p=0.5429$). *Post hoc* Sidak's multiple comparisons test
211 showed females were significantly faster to enter the port than males during the last 3 reward
212 sessions (R3, $p=0.001$; R4, $p=0.0391$; R5, $p=0.0014$). Taken together, female rats consistently

spent more time than males in the reward port during the reward cue in reward pre-training sessions.

3.2 Female rats did not show conditioned inhibition of freezing

After reward pre-training, rats were then exposed to sessions also consisting of reward, fear and safety cues. The reward cue and sucrose reward were the same as the reward pre-training sessions. The fear cue was paired with a 0.5mA footshock, and both the safety cue and fear+safety cue did not result in footshock or sucrose.

The percent time spent at or in the reward port during each cue across session was quantified for each DC session (Figure 2b). Two-way repeated-measures ANOVAs showed a significant cue by sex effect, as well as main effects of cue and sex for DC1 (Table 1). *Post hoc* Sidak's multiple comparisons test showed that, during DC1, females spent significantly more time reward seeking during the reward cue compared to males ($p<0.001$), consistent to what was seen in reward pre-training. For the remaining DC2-4 sessions, a main effect of cue was observed (Table 1) and *post hoc* Sidak's multiple comparisons test showed that both male and female rats spent significantly more time reward seeking during the reward cue compared to all other cues ($p<0.0001$), with no significant differences between the males and females. Thus, the noticeable increase in reward seeking in the females, that was seen during reward pre-training, dissipated by the 2nd DC session.

The percent time freezing during each cue across session was quantified for each DC session (Figure 2c). Two-way repeated-measures ANOVAs showed a significant cue by sex effect for sessions DC2-4, as well as main effects of cue and sex for every session (Table 1). *Post hoc* Sidak's multiple comparisons tests showed that, for every session, females displayed significantly more freezing to the fear+safety cue compared to males (DC1, $p=0.0313$; DC2, $p=0.007$; DC3, $p=0.0007$; DC4, $p<0.0001$). Females also showed significantly higher freezing levels to the fear cue compared to males during DC2 ($p=0.0111$). Males showed a significant reduction in freezing levels to the fear+safety cue compared to the fear cue during sessions DC3 ($p=0.0156$) and DC4 ($p<0.0001$), thus showing significant conditioned inhibition of freezing. Females did not show a significant inhibition of freezing during any session.

The number of darts during each cue was also quantified for each DC session and expressed as a dart rate (Figure 2d; sum of darts/ # trials). Darting behavior during cue presentation was

247 largely absent until DC3 and DC4. Two-way repeated-measures ANOVAs showed a significant
248 cue by sex effect for DC4, as well as main effects of cue, for DC2-4, and sex, for DC1 and DC4
249 (Table 1). *Post hoc* Sidak's multiple comparisons test showed that, during DC4, females
250 expressed more darting behavior compared to males during both the fear cue (p=0.0017) and
251 the fear+safety cue (p=0.0186). The females reduced their darting rate from 0.51 during the fear
252 cue to 0.32 during the fear+safety cue, but this was not statistically significant.
253
254 *3.3 Female rats did not show significant extinction of freezing*
255 The day after the last DC session all rats underwent fear and reward extinction within the same
256 session. That is, both the fear and reward cues were presented within the same training
257 session, without footshocks or sucrose presentations.
258
259 During extinction of reward, there was no main effect of reward trial (F(19,646)=1.526,
260 p=0.0704) or sex (F(1,34)=1.31, p=0.2603) and no interaction (F(19,646)=0.8927, p=0.5924);
261 there was also no significant difference between male and female groups for any trial (Figure
262 3Bi). One day later when rats were re-tested for extinction memory (Figure 3Bii), there was a
263 main effect of cue (2-way RM ANOVA; F(3,102)=134.7, p<0.0001) and sex (F(1,34)=6.217,
264 p=0.0177). *Post hoc* Sidak's multiple comparisons test showed females had significantly more
265 port activity than males just during the safety cue (p=0.0452), although this difference did not
266 reflect a large increase in port activity as females spent 6.38% +/- 0.86 of the safety cue in the
267 port compared to 2.66% +/- 0.86 in males. Overall, there appeared to be no differences in the
268 ability of males and females to extinguish their reward seeking responses.
269
270 To assess fear extinction the averaged percent time freezing during each trial of fear extinction
271 training was calculated (Figure 3Ci). There was a main effect of fear trial (2-way RM ANOVA;
272 F(19, 646)=7.69, p<0.0001) and sex (2-way RM ANOVA; F(1, 34)=4.607, p=0.0391), but no
273 significant interaction (F(19, 646)=1.566, p=0.059). Compared to trial 1, males showed
274 significantly reduced freezing in extinction trials 8, 9 and 11-20 (*post hoc* Tukey's multiple
275 comparisons test, p<0.05), demonstrating good fear extinction beginning around the 8th trial. In
276 contrast, females only showed a significant reduction in freezing during trial 19 compared to the
277 first trial (*post hoc* Tukey's multiple comparisons test, p=0.0394), demonstrating relatively
278 absent fear extinction. One day later when rats were retested for extinction memory (Figure
279 3Cii), there was a main effect of cue (F(3,102)=134.7, p<0.0001) and sex (F(1, 34)=6.217,
280 p=0.0177), as well as a significant interaction of cue X sex (F(3, 102)=3.481, p=0.0187). *Post*

281 *hoc* Sidak's multiple comparisons test showed that females froze significantly more than males
282 to the fear (p=0.0146) and fear+safety (p=0.0091) cues. This indicates the continued absence of
283 any extinction of freezing in females.
284
285 In response to each fear cue presentation across extinction, we also assessed darting levels
286 (Figure 3Di). There was a main effect of sex (F(1,34)=4.816, p=0.0351), but no effect of trial
287 (F(6.957, 236.6)=0.6941, p=0.6762) and no significant interaction (F(19, 646)=1.083, p=0.3640).
288 *Post hoc* Sidak's multiple comparisons test showed no significant differences between males
289 and females for any trial. For the extinction memory test one day later (Figure 3Dii), there was a
290 significant cue X sex interaction (F(3,102)=4.447, p=0.0056), as well as a main effect of both
291 cue (F(2.013, 68.44)=4.248, p=0.018) and sex (F1, 34)=4.834, p=0.0348). Females showed a
292 significantly higher dart rate than males during the fear cue (*post hoc* Sidak's multiple
293 comparisons test, p<0.05), which was also significantly higher than the dart rate to the reward
294 and safety cues in the females (*post hoc* Sidak's multiple comparisons test, p<0.05). However,
295 even though statistically significant, the amount of darting during the fear cue in females was
296 very low (0.05-0.4 in extinction training and 0.065 in the extinction memory test), and therefore
297 no clear conclusions can be made regarding darting and extinction in this study.
298
299 *3.4 Shock reactivity in males versus females*
300 To exclude possible sex differences in pain sensitivity and footshock perception, a separate
301 cohort of 8 male and 8 age-matched female rats received 11 unsignaled footshocks of
302 increasing intensities (0.3 mA, 0.35 mA, 0.4 mA, 0.45 mA, 0.5 mA, 0.55 mA, 0.6 mA, 0.7 mA,
303 0.8 mA, 0.9, 1.0 mA) with an inter-stimulus interval of 2 min. Freezing increased as a function of
304 shock intensities (Figure 4A; 2-way RM ANOVA; F(11,121)=25.9, p<0.0001). No main effects of
305 sex (F(1,121)=0.2871, p=0.6027) or sex by shock (F(11,121)=1.413, p=0.1754) were observed.
306 Our experiments utilized a shock intensity of 0.5mA throughout this study. For this particular
307 intensity, we also noted the number of rats that jumped or darted in response to a 0.5mA shock
308 (Figure 4B,C). No sex differences in the number of rats jumping in response to the 0.5mA
309 footshock were observed (χ2: p>0.9). The number of female rats darting after the 0.5mA
310 footshock was higher than males, but not significantly (χ2: p =0.0769), with five of the eight
311 female rats tested exhibiting the behavior. A higher number of females darting in response to
312 the footshock in this test would still not explain the lack of conditioned inhibition of freezing in
313 the females, as freezing levels at 0.5mA was slightly lower than the males (Figure 4A). Our

314 results do not definitively show, but do suggest, that females may be more likely to respond to a
315 footshock with a darting response.
316
317 **4. Discussion**
318 In this study, we show females exhibit a significantly different behavioral profile than males in a
319 task that tests for reward, fear and safety cue discrimination, as well as conditioned inhibition
320 and extinction. Female Long Evans rats showed more reward seeking early in training and
321 persistently high freezing levels to the fear cue when in the presence of a safety cue or after
322 fear extinction. Darting behavior in the females late in training showed conditioned inhibition of
323 this behavior in the presence of a safety cue, suggesting the females are able to discriminate
324 between the fear and safety cues but do not suppress their freezing response. This data adds to
325 the growing body of evidence of sex differences in fear regulation and highlights the advantages
326 of using more complex learning paradigms with additional behavioral measurements.
327
328 Even though studies including female subjects have been proportionally low, several studies
329 have reported clear sex differences in fear regulation. Most of these are consistent with our
330 findings of reduced discrimination between fear and safety signals. For instance, female mice
331 show more generalization of fear to novel and safe contexts compared to males, and with this
332 generalization there is a concurrent increase in basal amygdala activity (Keiser et al., 2017).
333 Male and female rats also respond differently to the controllability of a stressor. Males display
334 reduced fear during escapable stress versus inescapable stress whereas females exhibit no
335 beneficial effects of perceiving a stressor as escapable and controllable (Baratta et al., 2018).
336 The buffering effects seen in these males were linked to prelimbic cortical neurons projecting to
337 the dorsal raphe nucleus, which do not appear to be engaged in females. Females displaying a
338 similar fear response to both inescapable and escapable stress is similar to our findings of
339 females showing equivalent freezing levels to the fear cue in the presence or absence of a
340 safety cue, in that there were no buffering effects seen by the safety cue. It appears that
341 females do not downregulate their fear response in situations cued as safe.
342
343 Our data showing an increase in darting behavior in female rats as the number of fear cue-
344 footshock trials increase is consistent with another report using female rats in a fear conditioning
345 and extinction paradigm (Gruene et al., 2015). Like us, Gruene et al (2015) also show darting
346 levels increase as learning about the fear cue advances. Compared to us, Gruene et al (2015)
347 report notably higher darting frequencies, which is most likely due to the differences in shock

intensities and number of trials; our study used 4 trials of 0.5mA per day for 4 days compared to their study using 7 trials of 0.7mA on one day. Our study also includes reinforced reward trials within the same sessions as the fear cue-footshock trials, which could alter the contextual expectations of the training session and reduce overall darting levels. It would be interesting in future studies to identify what leads a female to become a 'darter' versus 'non-darter'. As darting is a more active response compared to freezing, the circuits engaged during potential threats would likely be different in these two populations.

Our findings showing a lack of conditioned inhibition of freezing in females appear to be inconsistent with a recent study demonstrating a lack of sex differences in conditioned inhibition of freezing (Foilb et al., 2018). This is likely due to differences in our respective protocols. First, their footshock intensity was 1.2mA, resulting in freezing levels >90% during the fear cue. As footshock intensity and number of trials are consistently inconsistent across studies, it would be interesting to assess if freezing and darting levels in females follow a linear trend with increasing training intensity, or if there is instead a possibly U-shaped relationship. Foilb et al (2018) also used separate presentations of the fear cue and safety cue throughout training and employed the fear+safety cue summation test during recall, whereas we include fear+safety trials as part of the training. In contrast, another study has shown females discriminate equally to males early in training but then generalize their fear response to the safety cue with continued training (Day et al., 2016). While the females in our study clearly showed equivalent freezing levels to both the fear and fear+safety cues at all time points throughout training, they did not increase their freezing levels to the safety cue when presented alone. And, lastly, our paradigm, unlike others, includes reinforced reward trials during the training of fear and safety cues, which would change the context from a 'threat-no threat' situation to a 'threat-no threat-reward' situation, inducing approach behaviors on top of defensive behaviors.

Altogether, the data paints a consistent picture of females showing heightened fear responses to cues signaling safety, mimicking the clinical picture in women (Gamwell et al., 2015; Lonsdorf et al., 2015). The presentation of a safety signal not only decreases fear, but also stimulates opposing neuronal activity. Field potential recordings in the striatum during safety signal presentation has shown that brain regions dealing with approach and reward become activated (Rogan et al., 2005). These findings have also been translated to using safety signals to overcome anhedonia in rats (Pollak et al., 2008), showing that safety signals may also be regulating emotion in addition to conditioned behavior (Foilb & Christianson, 2018).

382

383 In our study, females consistently showed elevated reward seeking behavior during the reward
384 cue compared to males beginning in the second reward pre-training session. This data appears
385 consistent with reward studies showing significant sex differences in response to sucrose, with
386 females willing to work more for sucrose in a progressive ratio paradigm (Tapia et al., 2019),
387 and in response to drugs of abuse, with female rats consistently self-administering drugs more
388 rapidly than males (Becker & Koob, 2016). The increased reward seeking in females seen in our
389 study remained until the end of the first DC session at which point they were equivalent to the
390 males. Interestingly, DC1 is the first time the animals are exposed to footshock. Taking into
391 account the lack of conditioned inhibition of freezing in the females, the females may no longer
392 be as motivated to seek rewards in the face of adverse footshocks. This would be consistent
393 with the report that female rats sacrifice their metabolic needs in order to avoid shocks more
394 than males (Pellmanet al., 2017).
395
396 Numerous sex differences have been reported in the functioning of the stress neuropeptide,
397 corticotropin-releasing factor (CRF), with differences in receptor expression, distribution,
398 trafficking and signaling (reviewed in (Bangasser & Wiersielis, 2018)). The majority of these
399 differences lead to enhanced CRF efficacy in females which may lead to heightened sensitivity
400 to stressors in females. Recently, the gene for CRH receptor 1 (*CRHR1*) has been identified as
401 a possible candidate gene for mood and anxiety disorders. Weber et al. (2016) have shown that
402 carrying the *CRHR1* minor rs17689918 allele increases the risk for panic disorders in women.
403 Patients carrying this risk allele also demonstrate more generalization of fear to a safety cue,
404 increased amygdala activation during the safety cue and decreased frontal cortex activation with
405 discriminative fear conditioning. Thus, aberrant CRF signaling can lead to sustained fear under
406 conditions cued as safe and can be manifested by changes in neural activity in the amygdala
407 and frontal cortex.
408
409 Neural activity in the amygdala and prefrontal cortex has been shown by our lab to also play a
410 critical role in effective discriminative conditioning in male rats. We have previously identified
411 neurons in the basolateral amygdala (BLA) that discriminate among safety, fear and reward
412 cues in male rats (Sangha et al., 2013); our future experiments will test if females show the
413 same discriminative neurons. Using reversible pharmacological inactivations in male rats, we
414 have also demonstrated that the infralimbic prefrontal cortex (IL) is necessary for suppression of
415 conditioned fear during a safety cue and the prelimbic prefrontal cortex (PL) is necessary for

fear expression and discriminatory reward seeking (Sangha, Robinson, et al., 2014). These results indicate that activating the IL in the females may improve conditioned inhibition to the combined fear and safety cues. Our results with male rats also show that manipulating D1-receptor mediated dopamine activity in the BLA disrupts suppression of conditioned fear (Ng et al., 2018), implicating dopaminergic ventral tegmental area (VTA) neurons projecting to the BLA in safety-fear-reward discrimination.

Our findings are consistent with human studies where females show less discrimination between the fear and safety signals than males (Gamwell et al., 2015; Lonsdorf et al., 2015), which may reflect underlying mechanisms of increased prevalence for anxiety and stress-related disorders in women. For example, a deficiency in effective safety signal processing has been linked to Post-traumatic Stress Disorder (Jovanovic et al., 2009, 2010), panic disorder (Gorka et al., 2014), and anxiety (Lissek et al., 2005), all disorders with a higher incidence in women than men (Mclean et al., 2011). In our paradigm, females show a significantly different behavioral profile than males that is consistent with the clinical picture, thus making it a great tool to test for the neurobiological mechanisms underlying these sex differences.

5. Acknowledgements:

We thank Yolanda Jonker, Signe Hobaugh, Jessica Kerns, Emma Speckelson, Emily Willis, and Mackenzie McIntosh for excellent animal care. IM was supported by the Alexander von Humboldt foundation. This work was supported by the National Institutes of Health [NIMH R01MH110425 to SS].

6. References

Bangasser, D. A., & Wiersielis, K. R. (2018). Sex differences in stress responses : a critical role for corticotropin-releasing factor. *Hormones*, *17*(1), 5–13.

Baratta, M. V, Leslie, N. R., Fallon, I. P., Dolzani, S. D., Chun, L. E., Tamalunas, A. M., … Maier, S. F. (2018). Behavioural and neural sequelae of stressor exposure are not modulated by controllability in females. *European Journal of Neuroscience*, *47*, 959–967. http://doi.org/10.1111/ejn.13833

Becker, J. B., & Koob, G. F. (2016). Sex Differences in Animal Models : Focus on Addiction. *Pharmacological Reviews*, *68*, 242–263.

Blanchard, R. J., & Blanchard, D. C. (1969). Crouching as an index of fear. *Journal of Comparative and Physiological Psychology*, *67*(3), 370–5.

Christianson, J. P., Fernando, a. B. P., Kazama, a. M., Jovanovic, T., Ostroff, L. E., & Sangha, S. (2012). Inhibition of Fear by Learned Safety Signals: A Mini-Symposium Review. *Journal of Neuroscience*, *32*(41), 14118–14124. http://doi.org/10.1523/JNEUROSCI.3340-12.2012

Conway, S. M., Puttick, D., Russell, S., Potter, D., Roitman, M. F., & Charto, E. H. (2019). Neuropharmacology Females are less sensitive than males to the motivational- and dopamine- suppressing e ff ects of kappa opioid receptor activation, *146*(November 2018), 231–241. http://doi.org/10.1016/j.neuropharm.2018.12.002

Daviu, N., Andero, R., Armario, A., & Nadal, R. (2014). Sex differences in the behavioural and hypothalamic-pituitary-adrenal response to contextual fear conditioning in rats. *Hormones and Behavior*, *66*(5), 713–723. http://doi.org/10.1016/j.yhbeh.2014.09.015

Day, H. L. L., Reed, M. M., & Stevenson, C. W. (2016). Sex differences in discriminating between cues predicting threat and safety. *Neurobiology of Learning and Memory*, *133*, 196–203. http://doi.org/10.1016/j.nlm.2016.07.014

Fendt, M., & Fanselow, M. S. (1999). The neuroanatomical and neurochemical basis of conditioned fear. *Neuroscience & Biobehavioral Reviews*, *23*(5), 743–760. http://doi.org/10.1016/S0149-7634(99)00016-0

Foilb, A. R., Bals, J., Sarlitto, M. C., & Christianson, J. P. (2018). Sex differences in fear discrimination do not manifest as differences in conditioned inhibition. *Learning and Memory*, *25*(1), 49–53. http://doi.org/10.1101/lm.045500.117

Foilb, A. R., & Christianson, J. P. (2018). Chapter 11 Brain Mechanisms for Learning and Using Safety Signals. In S. Sangha & D. Foti (Eds.), *Neurobiology of Abnormal Emotion and Motivated Behaviors: Integrating Animal and Human Research* (1st editio, pp. 205–224). Cambridge, Massachusetts: Academic Press.

Gamwell, K., Nylocks, M., Cross, D., Bradley, B., Norrholm, S. D., & Jovanovic, T. (2015). Fear Conditioned Responses and PTSD Symptoms in Children : Sex Differences in Fear-Related Symptoms. *Developmental Psychobiology*, *57*, 799–808. http://doi.org/10.1002/dev.21313

Glover, E. M., Jovanovic, T., & Norrholm, S. D. (2015). Estrogen and extinction of fear memories: Implications for posttraumatic stress disorder treatment. *Biological Psychiatry*, *78*(3), 178–185. http://doi.org/10.1016/j.biopsych.2015.02.007

Gorka, S. M., Lieberman, L., Nelson, B. D., Sarapas, C., & Shankman, S. A. (2014). Aversive responding to safety signals in panic disorder: The moderating role of intolerance of uncertainty. *Journal of Anxiety Disorders*, *28*(7), 731–736. http://doi.org/10.1016/j.janxdis.2014.06.004

Grant, B. F., Saha, T. D., June Ruan, W., Goldstein, R. B., Patricia Chou, S., Jung, J., … Hasin, D. S. (2016). Epidemiology of DSM-5 drug use disorder results from the national epidemiologic survey on alcohol and related conditions-III. *JAMA Psychiatry*, *73*(1), 39–47. http://doi.org/10.1001/jamapsychiatry.2015.2132

Gruene, T. M., Flick, K., Stefano, A., Shea, S. D., & Shansky, R. M. (2015). Sexually divergent expression of active and passive conditioned fear responses in rats. *ELife*, *4*, 1–9. http://doi.org/10.7554/eLife.11352

Hernandez-Avila, C. A., Rounsaville, B. J., & Kranzler, H. R. (2004). Opioid-, cannabis- and alcohol-dependent women show more rapid progression to substance abuse treatment. *Drug and Alcohol Dependence*, *74*(3), 265–272. http://doi.org/10.1016/j.drugalcdep.2004.02.001

Jovanovic, T., Kazama, A., Bachevalier, J., & Davis, M. (2012). Impaired safety signal learning may be a biomarker of PTSD. *Neuropharmacology*, *62*(2), 695–704. http://doi.org/10.1016/j.neuropharm.2011.02.023

Jovanovic, T., Norrholm, S. D., Blanding, N. Q., Davis, M., Duncan, E., Bradley, B., & Ressler, K. J. (2010). Impaired fear inhibition is a biomarker of PTSD but not depression. *Depression and Anxiety*, *27*(3), 244–251. http://doi.org/10.1002/da.20663

Jovanovic, T., Norrholm, S. D., Fennell, J. E., Keyes, M., Fiallos, A. M., Myers, K. M., … Duncan, E. J. (2009). Posttraumatic stress disorder may be associated with impaired fear inhibition: Relation to symptom severity. *Psychiatry Research*, *167*(1–2), 151–160. http://doi.org/10.1016/j.psychres.2007.12.014

Keiser, A. A., Turnbull, L. M., Darian, M. A., Feldman, D. E., Song, I., & Tronson, N. C. (2017). Sex Differences in Context Fear Generalization and Recruitment of Hippocampus and

Amygdala during Retrieval. *Neuropsychopharmacology*, *42*(2), 397–407. http://doi.org/10.1038/npp.2016.174

Keyes, K. M., Grant, B. F., & Hasin, D. S. (2008). Evidence for a closing gender gap in alcohol use, abuse, and dependence in the United States population. *Drug and Alcohol Dependence*, *93*(1–2), 21–29. http://doi.org/10.1016/j.drugalcdep.2007.08.017

Leknes, S., Lee, M., Berna, C., Andersson, J., & Tracey, I. (2011). Relief as a reward: Hedonic and neural responses to safety from pain. *PLoS ONE*, *6*(4), e17870. http://doi.org/10.1371/journal.pone.0017870

Lissek, S., Powers, A. S., McClure, E. B., Phelps, E. A., Woldehawariat, G., Grillon, C., & Pine, D. S. (2005). Classical fear conditioning in the anxiety disorders: A meta-analysis. *Behaviour Research and Therapy*, *43*(11), 1391–1424. http://doi.org/10.1016/j.brat.2004.10.007

Lonsdorf, T. . B., Haaker, J., Schümann, D., Sommer, T., Brassen, S., Bunzek, N., … Kalisch, R. (2015). Sex differences in CS-discrimination during context-dependent fear learning and its retrieval - the role of biological sex, contraceptives and menstrual cycle phases. *Journal of Psychiatry and Neuroscience*, *40*(6), 368–375. http://doi.org/10.1503/jpn.140336

Mclean, C., Asnaani, A., Litz, B., & Hofmann, S. (2011). Gender differences in anxiety disorders: Prevalence, course of illness, comorbidity and burden of illness. *Journal of Psychiatric Research*, *45*(8), 1027–1035. http://doi.org/10.1016/j.jpsychires.2011.03.006.Gender

Ng, K. H., Pollock, M. W., Urbanczyk, P. J., & Sangha, S. (2018). Altering D1 receptor activity in the basolateral amygdala impairs fear suppression during a safety cue. *Neurobiology of Learning and Memory*, *147*, 26–34. http://doi.org/10.1016/j.nlm.2017.11.011

Pellman, B. A., Schuessler, B. P., Tellakat, M., & Kim, J. J. (2017). Sexually Dimorphic Risk Mitigation Strategies in Rats. *Eneuro*, *4*(1), e.0288-16.2017. http://doi.org/10.1523/ENEURO.0288-16.2017

Pollak, D. D., Monje, F. J., Zuckerman, L., Denny, C. a., Drew, M. R., & Kandel, E. R. (2008). An Animal Model of a Behavioral Intervention for Depression. *Neuron*, *60*(1), 149–161. http://doi.org/10.1016/j.neuron.2008.07.041

Rescorla, R. A. (1969). Conditioned inhibition of fear resulting from negative CS-US contingencies. *Journal of Comparative and Physiological Psychology*, *67*(4), 504–509. http://doi.org/10.1037/h0027313

Rogan, M. T., Leon, K. S., Perez, D. L., & Kandel, E. R. (2005). Distinct neural signatures for safety and danger in the amygdala and striatum of the mouse. *Neuron*, *46*(2), 309–20. http://doi.org/10.1016/j.neuron.2005.02.017

Sangha, S., Chadick, J. Z., & Janak, P. H. (2013). Safety encoding in the basal amygdala. *The Journal of Neuroscience*, *33*(9), 3744–51. http://doi.org/10.1523/JNEUROSCI.3302-12.2013

Sangha, S., Greba, Q., Robinson, P. D., Ballendine, S. A., & Howland, J. G. (2014). Heightened fear in response to a safety cue and extinguished fear cue in a rat model of maternal immune activation. *Frontiers in Behavioral Neuroscience*, *8*, 168. http://doi.org/10.3389/fnbeh.2014.00168

Sangha, S., Robinson, P. D., Greba, Q., Davies, D. A., & Howland, J. G. (2014). Alterations in reward, fear and safety cue discrimination after inactivation of the rat prelimbic and infralimbic cortices. *Neuropsychopharmacology*, *39*(10), 2405–13. http://doi.org/10.1038/npp.2014.89

Tanimoto, H., Heisenberg, M., & Gerber, B. (2004). Event timing turns punishment to reward. *Nature*, *430*, 983. http://doi.org/10.1038/nature02816

Tapia, M. A., Lee, J. R., Weise, V. N., Tamasi, A. M., & Will, M. J. (2019). Sex di ff erences in hedonic and homeostatic aspects of palatable food motivation. *Behavioural Brain Research*, *359*(November 2018), 396–400. http://doi.org/10.1016/j.bbr.2018.11.023

Walasek, G., Wesierska, M., & Zieliński, K. (1995). Conditioning of fear and conditioning of safety in rats. *Acta Neurobiologiae Experimentalis*, *55*(2), 121–132.

Weber, H., Richter, J., Straube, B., Lueken, U., Domschke, K., Schartner, C., … Reif, A. (2016). Allelic variation in CRHR1 predisposes to panic disorder: Evidence for biased fear processing. *Molecular Psychiatry*, *21*(6), 813–822. http://doi.org/10.1038/mp.2015.125

562 **7. Figure Legends**

563 **Figure 1**. **Females show increased reward seeking in response to the reward cue**. **A)**
564 Schematic depicting experimental outline. During reward pre-training, rats (16 males, 20
565 females) received 25 cue-sucrose pairings across 5 separate sessions. **B)** Averaged percent
566 time spent in the reward port during the five reward pre-training sessions (R1-5). Females spent
567 significantly more time in the port compared to males during R2, R3 and R5. **C)** Averaged
568 latency to enter the port after cue onset (in seconds). Females entered the port significantly
569 sooner than males during R3-5. Means +/- SEM. *p<0.05, **p<0.01.
570

571 **Figure 2. Females do not show inhibition of conditioned freezing in the presence of the**
572 **safety cue. A)** Schematic depicting experimental outline. During the 4 DC sessions, rats (16
573 males, 20 females) were presented with four types of cued trials: reward cue-sucrose, fear cue-
574 shock, fear+safety cue with no footshock and the safety cue presented alone without footshock.
575 **B)** Averaged percent time spent in the port during each cue across the 4 DC sessions. Both
576 males and females showed significantly higher reward seeking during the reward cue compared
577 to all other cues during every DC session. During DC1, females showed significantly higher
578 reward seeking to the reward cue compared to males. **C)** Averaged percent time spent freezing
579 during each cue across the 4 DC sessions. During DC3 and DC4, males showed significantly
580 lower freezing to the fear+safety cue (and reward and safety cues) when compared to the fear
581 cue. Females did not show significant inhibition of conditioned freezing to the fear+safety cue
582 compared to the fear cue during any DC session. Females also showed significantly higher
583 freezing to the fear+safety cue compared to males during every session. **D)** Darting behavior
584 during each cue across the 4 DC sessions. During DC4 females showed significantly more darts
585 than males during the fear and fear+safety cues. Means +/- SEM. # p<0.05, ####p<0.0001
586 within sex, between cue comparison; * p<0.05, **p<0.01, ***p<0.001, ****p<0.0001 within cue,
587 between sex comparison.
588

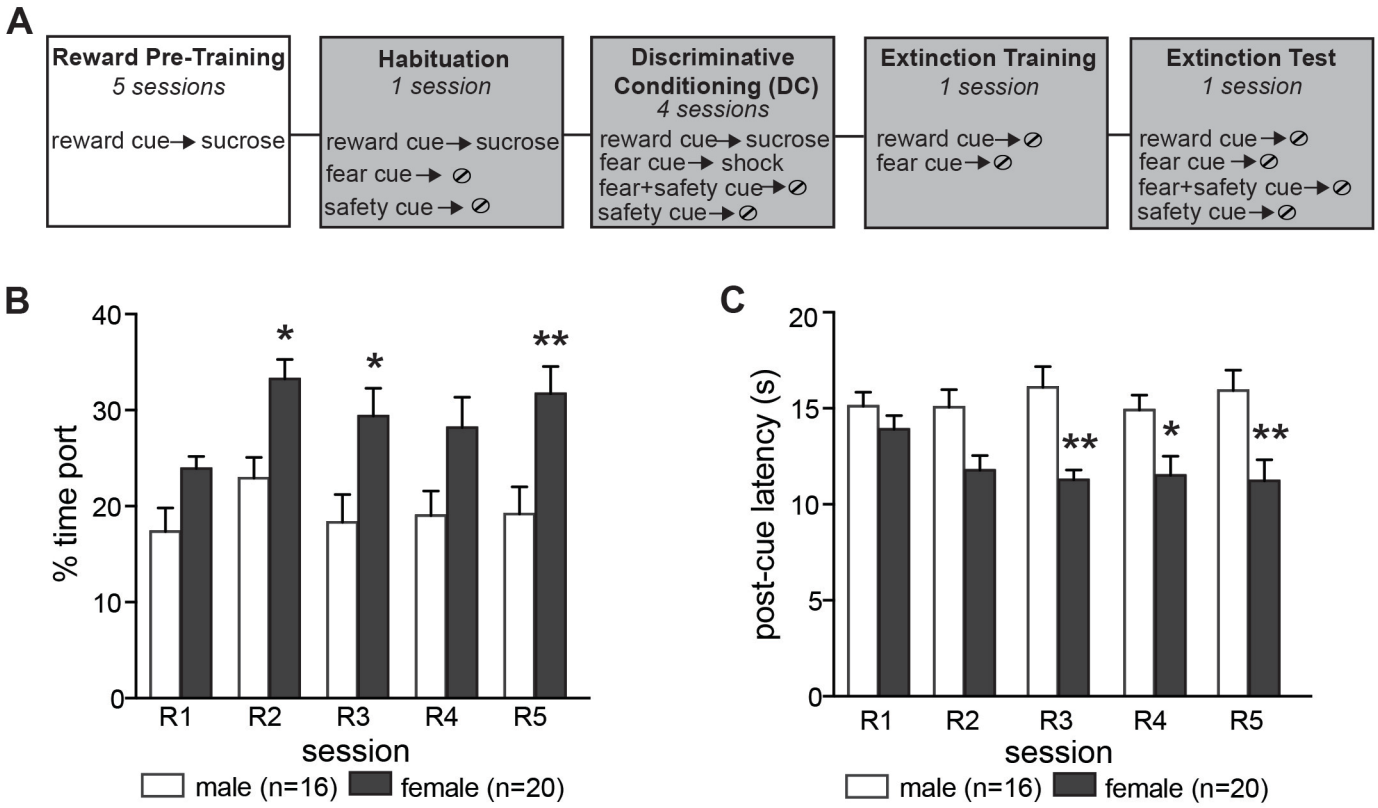
589 **Figure 3. Females do not show significant extinction of fear. A)** Schematic depicting
590 experimental outline. During extinction training both the reward and fear cues are presented in
591 the same session without sucrose or footshock. During the test for extinction memory 1 day
592 later all cues are presented without sucrose or footshock. **Bi)** Averaged percent time spent in
593 the port during each reward cue presentation during extinction training. No significant
594 differences were found between males and females during extinction training. **Bii)** Averaged
595 percent time spent in the port during each cue 1 day after extinction training. Females spent

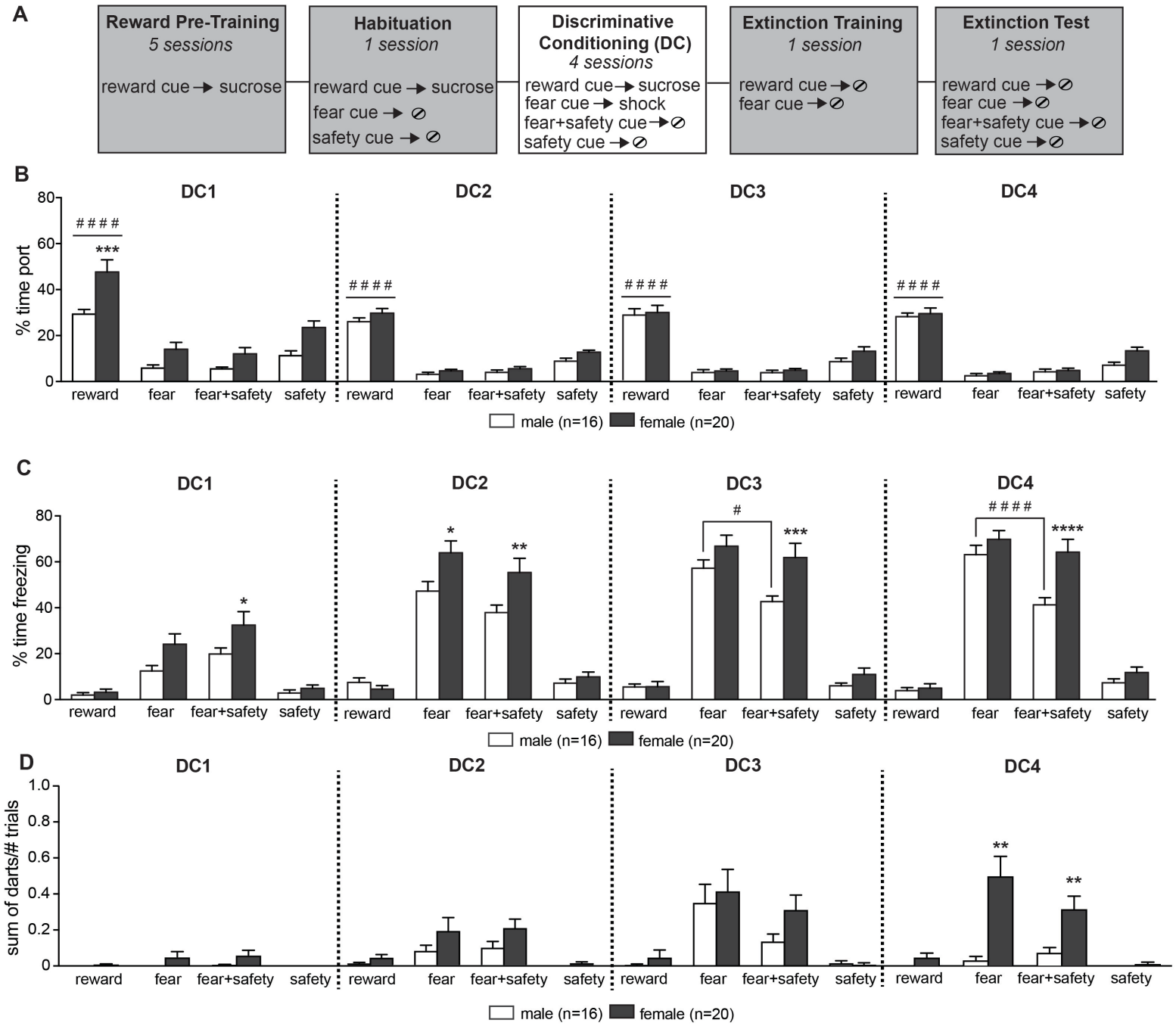
596 significantly more time in the port than males during the safety cue. **Ci)** Averaged percent time
597 spent freezing during each fear cue presentation during extinction training. Compared to the first
598 trial of extinction, males showed significantly reduced freezing during trials 8, 9 and 11-20.
599 Freezing levels for females did not significantly decrease at any point in extinction training, with
600 the exception of trial 19. #p<0.05, compared to trial 1. **Cii)** Averaged percent time spent freezing
601 during each cue 1 day after extinction training. Males showed evidence of fear cue extinction
602 retention. Females froze significantly more than males during the fear and fear+safety cues. **Di)**
603 Averaged darting during each fear cue presentation during extinction training. No significant
604 post hoc differences found between males and females during extinction training. **Dii)** Averaged
605 darting during each cue 1 day after extinction training. Females had a significantly higher dart
606 rate than males during the fear cue, which was also significantly higher than the reward and
607 safety cues in females. Means +/- SEM. #p<0.05, ####p<0.0001 within sex, between cue/trial
608 comparisons. *p<0.05, **p<001, ****p<0.0001 within cue, between sex comparisons.
609
610 **Figure 4. No significant differences in shock reactivity between age-matched male and**
611 **female rats. A)** Male and female rats (n=8 each) were subjected to increasing footshock
612 intensities from 0.3mA to 1.0mA. No significant differences in freezing levels (means +/- SEM)
613 were detected between males and females after each shock presentation. The box around the
614 data at 0.5mA indicates the intensity used for the experiments in this study. There were no
615 significant differences in the number of males or females who jumped **(B)** or darted **(C)** in
616 response to the 0.5mA shock.

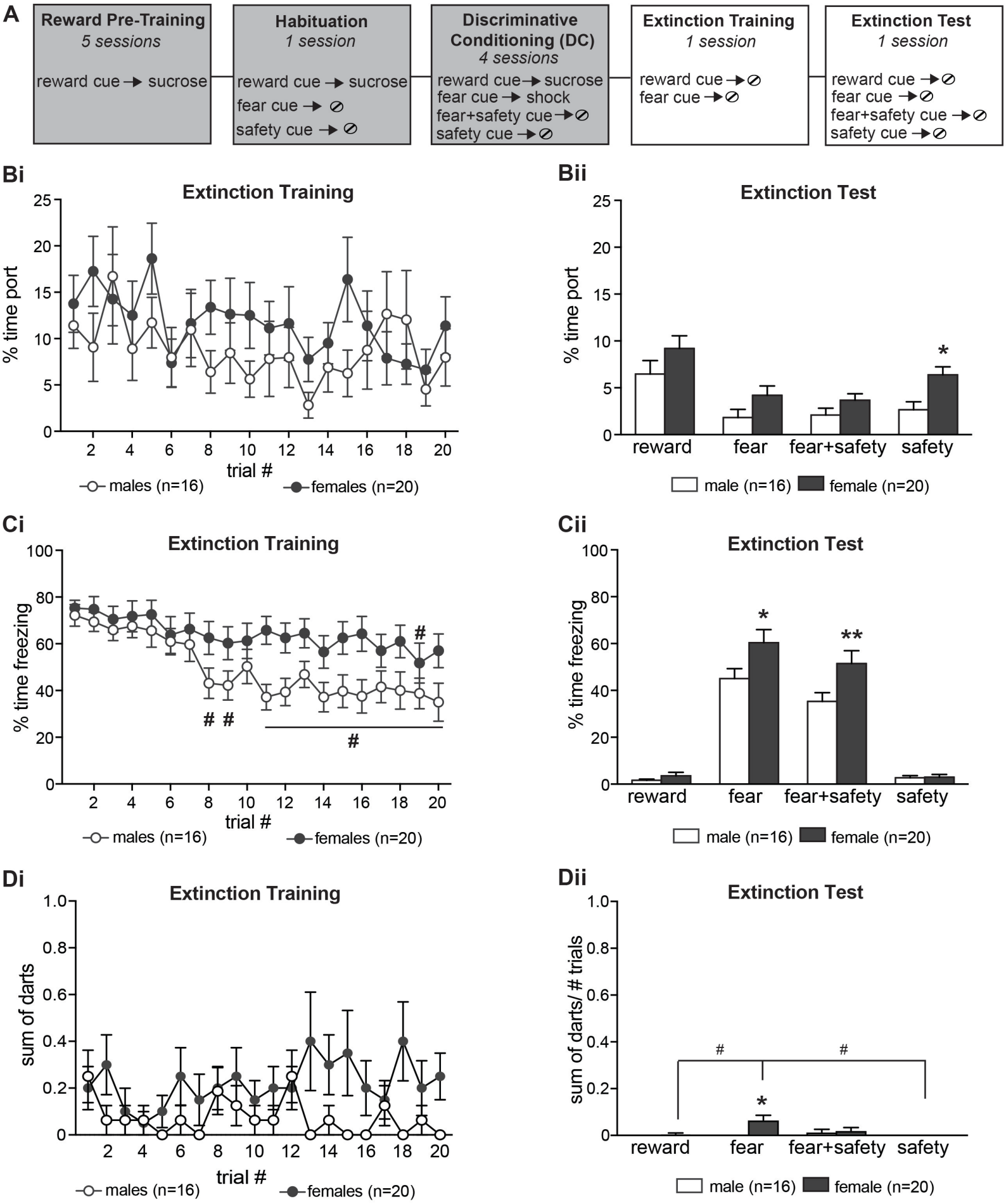
Reward seeking			
Session	*Cue x Sex effects*	*Main effect of cue*	*Main effect of sex*
DC1	F(3,102) = 3.472, **p=0.0189**	F(3,102) = 95.16, **p<0.0001**	F(1,34) = 9.827, **p=0.0035**
DC2	F(3,102) = 0.7742, p=0.5110	F(3,102) = 227.9, **p<0.0001**	F(1,34) = 4.69, **p=0.0374**
DC3*	F(3,90) = 0.6512, p=0.5843	F(3,90) = 117, **p<0.0001**	F(1,30)=1.041, p=0.3157
DC4	F(3,102) = 2.255, p=0.0864	F(3,102) = 181.2, **p<0.0001**	F(1,34) = 2.453, p=0.1266
Freezing			
Session	*Cue x Sex effects*	*Main effect of cue*	*Main effect of sex*
DC1	F(3,102) = 2.245, p=0.0876	F(3,102) = 31.82, **p<0.0001**	F(1,34) = 5.045, **p=0.0313**
DC2	F(3,102) = 4.075, **p=0.0089**	F(3,102) = 103.4, **p<0.0001**	F(1,34) = 6.621, **p=0.0146**
DC3*	F(3,90) = 2.9, **p=0.0393**	F(3,90) = 151.3, **p<0.0001**	F(1,30)=9.719, **p=0.0040**
DC4	F(3,102) = 4.889, **p=0.0032**	F(3,102) = 198.9, **p<0.0001**	F(1,34) = 8.294, **p=0.0068**
Darting			
Session	*Cue x Sex effects*	*Main effect of cue*	*Main effect of sex*
DC1	F(3,102) = 1.98, p=0.1216	F(2.022,68.73) = 2.388, p=0.0989	F(1,34) = 4.146, **p=0.0496**
DC2	F(3,102) = 1.134, p=0.3390	F(1.923, 65.38) = 9.377, **p=0.0003**	F(1,34) = 3.667, p=0.0640
DC3	F(3,102) = 0.9158 p=0.4361	F(1.586,53.93) = 18.96, **p<0.0001**	F(1,34)=0.9579, p=0.3346
DC4	F(3,102) = 10.65, **p<0.0001**	F(1.684, 57.26) = 15.65, **p<0.0001**	F(1,34) = 13.34, **p=0.0009**

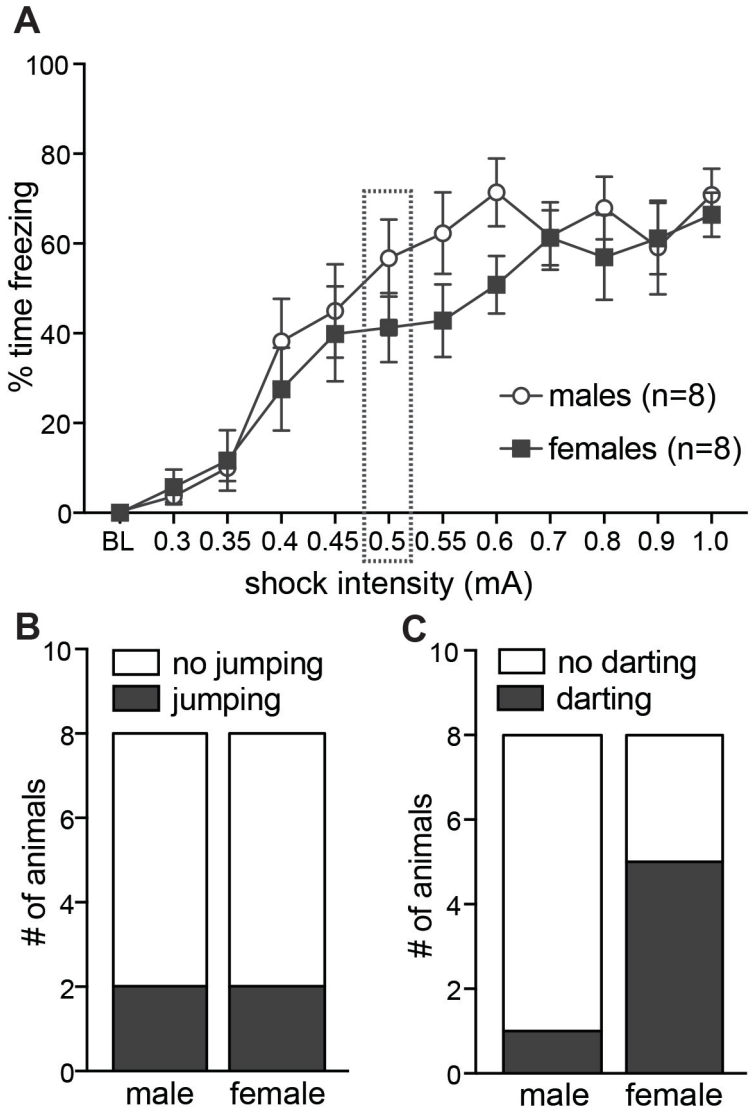
*video files for 4 females were corrupted for this session (n=16 females, 16 males)

Table 1. Summary of two-way repeated-measures ANOVA analyses for reward seeking, freezing and darting behaviors during the four discriminative conditioning (DC) sessions.


A
Reward Pre-Training
5 sessions
reward cue → sucrose
Habituation
1 session
reward cue → sucrose
fear cue → ⊘
safety cue → ⊘
Discriminative Conditioning (DC)
4 sessions
reward cue → sucrose
fear cue → shock
fear+safety cue → ⊘
safety cue → ⊘
Extinction Training
1 session
reward cue → ⊘
fear cue → ⊘
Extinction Test
1 session
reward cue → ⊘
fear cue → ⊘
fear+safety cue → ⊘
safety cue → ⊘
B
% time port
40
30
20
10
0
*
*
**
R1
R2
R3
R4
R5
session
male (n=16)
female (n=20)
C
post-cue latency (s)
20
15
10
5
0
**
*
**
R1
R2
R3
R4
R5
session
male (n=16)
female (n=20)


A
Reward Pre-Training
5 sessions
reward cue → sucrose
Habituation
1 session
reward cue → sucrose
fear cue → ⊘
safety cue → ⊘
Discriminative Conditioning (DC)
4 sessions
reward cue → sucrose
fear cue → shock
fear+safety cue → ⊘
safety cue → ⊘
Extinction Training
1 session
reward cue → ⊘
fear cue → ⊘
Extinction Test
1 session
reward cue → ⊘
fear cue → ⊘
fear+safety cue → ⊘
safety cue → ⊘
B
DC1
DC2
DC3
DC4
% time port
reward
fear
fear+safety
safety
male (n=16)
female (n=20)
C
% time freezing
D
sum of darts/# trials


A
Reward Pre-Training
5 sessions
reward cue → sucrose
Habituation
1 session
reward cue → sucrose
fear cue → ⊘
safety cue → ⊘
Discriminative Conditioning (DC)
4 sessions
reward cue → sucrose
fear cue → shock
fear+safety cue → ⊘
safety cue → ⊘
Extinction Training
1 session
reward cue → ⊘
fear cue → ⊘
Extinction Test
1 session
reward cue → ⊘
fear cue → ⊘
fear+safety cue → ⊘
safety cue → ⊘
Bi
Extinction Training
% time port
trial #
males (n=16)
females (n=20)
Bii
Extinction Test
% time port
reward
fear
fear+safety
safety
male (n=16)
female (n=20)
Ci
Extinction Training
% time freezing
trial #
males (n=16)
females (n=20)
Cii
Extinction Test
% time freezing
reward
fear
fear+safety
safety
male (n=16)
female (n=20)
Di
Extinction Training
sum of darts
trial #
males (n=16)
females (n=20)
Dii
Extinction Test
sum of darts/ # trials
reward
fear
fear+safety
safety
male (n=16)
female (n=20)


A
100
80
60
40
20
0
% time freezing
males (n=8)
females (n=8)
BL
0.3
0.35
0.4
0.45
0.5
0.55
0.6
0.7
0.8
0.9
1.0
shock intensity (mA)
B
no jumping
jumping
10
8
6
4
2
0
# of animals
male
female
C
no darting
darting
10
8
6
4
2
0
# of animals
male
female