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Glutamate receptor antagonist infusions into the basolateral and medial amygdala reveal differential contributions to olfactory vs. context fear conditioning and expression

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The basolateral amygdala’s involvement in fear acquisition and expression to visual and auditory stimuli is well known. The involvement of the basolateral and other amygdala areas in fear acquisition and expression to stimuli of other modalities is less certain. We evaluated the contribution of the basolateral and medial amygdala to olfactory and to context fear and fear conditioning by infusing into these areas the NMDA receptor antagonist AP5, the AMPA/kainate receptor antagonist NBQX, or vehicle prior to either odor-shock pairings or fear-potentiated startle testing. Pre-training AP5 infusions into the basolateral amygdala disrupted fear conditioning to the odor but not the context conditioned stimulus (CS). Pre-test NBQX infusions disrupted fear-potentiated startle to the odor but not context CS. Neither compound blocked fear conditioning when infused into the medial amygdala prior to training, but pre-test NBQX infusions did block fear-potentiated startle. The results confirm and extend recent findings suggesting a role for the basolateral amygdala in olfactory fear and fear conditioning, reveal an unexpected dissociation of the basolateral amygdala’s involvement in discrete cue versus context fear and fear conditioning, and implicate for the first time the medial amygdala in fear-potentiated startle.

For auditory and visual stimuli, the amygdala plays a key role in fear conditioning and fear expression as assessed with several behavioral measures (cf., Fendt and Fanselow 1999; Davis 2000; LeDoux 2000). We have found, for example, that pre-test infusions into the basolateral amygdala (i.e., the lateral, basolateral, and basomedial nuclei) of α-amino-3-hydroxy-5-methylisoxazole-4-propionic acid (AMPA)/kainate receptor antagonists block fear-potentiated startle to visual and auditory conditioned fear stimuli (CSs), and that pre-training (i.e., light-shock or tone-shock) infusions of either N-methyl-D-aspartate (NMDA) receptor antagonists or AMPA/kainate receptor antagonists block fear conditioning (cf., Walker and Davis 2002b). In most (e.g., Miserendino et al. 1990; Campeau et al. 1992; Gewirtz and Davis 1997), but not all (e.g., Fendt 2001; see also Lee et al. 2001) cases, pre-test intra-amygdala infusions of NMDA receptor antagonists have not been found to disrupt fear expression to auditory and visual CSs. Similar results have been obtained from other laboratories using different treatments and other fear-related behaviors (e.g., LeDoux et al. 1990; Muller et al. 1997; Groenink et al. 2000; Rodrigues et al. 2001).

Visual and auditory information reach the basolateral amygdala indirectly via intermediary structures such as auditory and visual thalamus and perirhinal cortex. Olfactory information also reaches the basolateral amygdala indirectly. Uniquely, however, the main and accessory olfactory bulbs also send direct projections to several amygdala areas including the medial and cortical nuclei, the nucleus of the lateral olfactory tract, and the periamygdaloid cortex (cf., Alheid et al. 1995; McDonald 1998; Pitkanen 2000). For olfactory fear and fear conditioning, the relative contribution of direct versus indirect inputs is unknown.

Using freezing as a behavioral measure of fear, Otto and colleagues have begun to assess the contribution of several brain areas. Lesion results suggest that the basolateral amygdala mediates fear responses to olfactory CSs (Cousens and Otto 1998) perhaps via afferents from the perirhinal cortex (Herzog and Otto 1997). Accompanying Fos data support this view and suggest also that the medial nucleus of the amygdala nucleus may participate in olfactory fear conditioning (Schettino and Otto 2001).

For the present study, there were three primary goals. First, we wished to confirm with an alternative behavioral measure (i.e., with fear-potentiated startle) the involvement of the basolateral amygdala in fear expression to an olfactory CS. Second, we wished to determine if the basolateral amygdala also participates in olfactory fear learning. And third, we wished to evaluate the contribution of the medial amygdala nucleus to both. The general strategy was to infuse glutamate receptor antagonists (either AMPA/kainate or NMDA receptor antagonists) into the area of interest either immediately prior to fear conditioning or immediately prior to testing. A disruption of fear-potentiated startle, assessed 48 h after training, would suggest an involvement of these areas either in olfactory fear learning (i.e., with pre-training infusions) or in fear expression (i.e., with pre-test infusions). The experimental design also allowed us to measure fear-potentiated startle to the context, and to evaluate the impact of these treatments on context fear and fear conditioning.

Results

Experiments IA and IB: Effect on fear conditioning and fear-potentiated startle of NMDA receptor blockade in the basolateral amygdala

Experiment IA (Acquisition)

Fifteen rats were infused with either artificial cerebrospinal fluid (ACSF; N = 8) or DL-2-amino-5-phosphonopentanoic acid (DL-

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Amygdala and odor vs. context conditioning

APS; 2.5 µg/side; \( N = 7 \). This dose was selected based on previous results from Campeau et al. (1992) and Miserendino et al. (1990). Fifteen minutes after the infusions were completed, rats were placed into the test chamber and 5 min later received the first of five odor-shock pairings (ITI = 4 min). Three ACSF-infused rats were later found not to have bilateral basolateral amygdala cannulations (as determined based on inclusion criteria described in the Methods section). The data from these rats were excluded from the statistical analyses (paired \( t \)-tests) that follow. Placements for the remaining animals are shown in Figure 1 as filled circles. As shown in Figure 2 (leftmost bars), fear-potentiated startle was significantly lower in rats that were trained following APS infusions into the basolateral amygdala compared to rats that were trained following ACSF infusions, \( t(11) = 2.92 \). Mean footshock reactivity (± S.E.M.), measured using the same procedure used to measure startle amplitude during training, was similar in the ACSF (1188 ± 295) and APS (1048 ± 301) groups.

Because the primary goal of Experiment 1A was to demonstrate an involvement of the basolateral amygdala in olfactory fear conditioning, and because this was accomplished using APS, we did not also evaluate the involvement of the basolateral amygdala in olfactory fear conditioning with the other compounds used in this study, 1,2,3,4-tetrahydro-6-nitro-2,3-dioxobenzof[\( f \)]quinoxa-line-7-sulfonamide (NBQX).

**Experiment 1B (Expression)**

Using a counterbalanced design with test days separated by 48 h, 10 rats received pre-test infusions of ACSF and, on another day, D,L-AP5 (2.5 µg/side). Immediately after the infusion procedure was completed, rats were placed into the test chamber. Collection of test data for odor-potentiated startle began 20 min later. Data from two rats that did not have bilateral cannulations in the basolateral amygdala were excluded from the statistical analysis that follows. Placements for the remaining animals are indicated in Figure 1 by filled triangles (point up). As shown in Figure 2 (middle bars), APS did not significantly influence fear-potentiated startle, \( t(7) = 0.44 \), when infused into the basolateral amygdala prior to testing.

Although extinction from the first to the second test is a potential concern in counterbalanced designs of this sort, there was no evidence of extinction in this experiment, probably due to the small number (i.e., 10) of test/extinction trials. Mean percent change scores for rats that received ACSF on day 1 followed by APS on day 2 (\( N = 3 \)) were 25.1 ± 4.9 and 56.4 ± 27.7, respectively. For rats that received APS on day 1 followed by ACSF on day 2 (\( N = 5 \)), mean percent change scores were 88.7 ± 26.2 and 91.0 ± 50.2, respectively.

**Experiment 2: Effect on fear-potentiated startle of AMPA/kainate receptor blockade in the basolateral amygdala**

Using a counterbalanced design with test days separated by 48 h, 14 rats were infused with ACSF and NBQX (3 µg/side—selected on the basis of previous findings from Walker and Davis 1997) disodium salt prior to fear-potentiated startle testing. Data from two rats that did not have bilateral cannulations in the basolateral amygdala were excluded from the statistical analyses that follow. Placements for the 12 remaining animals are indicated in Figure 1 by filled triangles (point down). As shown in Figure 2 (rightmost bars), NBQX significantly disrupted fear-potentiated startle, \( t(11) = 4.65 \).

As with Experiment 1, there was little evidence of between-session extinction. The mean percent change score for rats that received ACSF on day 1 (\( N = 7 \)) was 127.6 ± 39.3, and for rats that received ACSF on day 2 (\( N = 5 \)), 89.6 ± 34.4. The difference was not significant (\( P = 0.41 \)).

**Experiment 3: Effect on olfactory fear conditioning of AMPA/kainate and NMDA receptor blockade in the medial nucleus of the amygdala**

Thirty rats were infused with ACSF, NBQX (3 µg/side) disodium salt, or D,L-AP5 (2.5 µg/side) prior to odor-shock pairings. Data from 10 rats that did not have bilateral cannulations in or immediately adjacent to the medial amygdala nucleus were excluded from the statistical analysis (single factor ANOVA) that follows. Placements for the remaining animals (ACSF \( N = 7 \), APS \( N = 8 \), NBQX \( N = 6 \)) are indicated in Figure 1 by open circles. As shown in Figure 3 (leftmost bars), the antagonists did not disrupt fear-conditioning to the olfactory CS. In fact, mean percent potentiation scores were higher in these groups.
infused with NBQX prior to training.

The NMDA receptor antagonist AP5 disrupted fear conditioning to an olfactory CS when infused prior to training (leftmost bars). The AMPA/kainate receptor antagonist NBQX did disrupt fear-potentiated startle when infused prior to testing (rightmost bars). Baseline startle amplitude (i.e., from noise-alone test trials), from which percent-potentiation scores are derived, is indicated immediately below each bar.

**Experiments 4A and 4B: Effect on fear-potentiated startle of AMPA/kainate receptor antagonist infusions into the medial nucleus of the amygdala**

**Olfactory CS (Experiment 4A)**

Twenty-nine rats were infused with ACSF or NBQX (3 μg/side) disodium salt prior to fear-potentiated startle testing. Data from 12 rats that did not have bilateral canulations in or immediately adjacent to the medial amygdala nucleus were excluded from the statistical analysis (independent sample *t*-test) that follows. Placements for the remaining animals (ACSF [N = 7], NBQX [N = 10]) are indicated in Figure 1 by open triangles (point up). As shown in Figure 3 (middle bars), NBQX significantly disrupted fear-potentiated startle to the olfactory CS, *t*(19) = 2.26.

Because rats were trained and tested in the same chamber, we considered the possibility that startle amplitude was elevated in ACSF-infused rats due to context conditioning and that this effect was blocked by NBQX. For the purest assessment of context conditioning (i.e., one not contaminated by intermixed CS test trials), we compared startle amplitudes on the 30 noise-alone test trials that occurred at the beginning of the pre-shock acclimation session with startle amplitude on the 30 noise-alone test trials that occurred at the beginning of the post-conditioning test session (see Methods section for details). Indeed, the startle amplitude of ACSF- but not NBQX-infused rats increased markedly from the noise-alone test trials that preceded the first CS presentation; see Methods section for details. Statistically, an ANOVA on the between-session percent change scores indicated a significant Treatment effect, *F*(1,34) = 4.93, but not a significant Modality or interaction effect. The absolute level of startle prior to conditioning was somewhat, although not significantly, lower in the groups that later received NBQX (i.e., vs. the groups that later received ACSF). We do not believe that this in any way accounts for the apparent disruption of context conditioning insofar as we have previously found that percent

**Visual CS (Experiment 4B)**

Because the medial nucleus of the amygdala receives a prominent olfactory input relative to other modalities (cf., DeOlmos et al. 1985; Pitkanen 2000), we wondered if the disruptive effect of NBQX was modality-specific. To this end, an additional group of rats (N = 31) was trained as previously described, this time using a visual rather than olfactory stimulus. Data from 10 rats that did not have bilateral canulations in or immediately adjacent to the medial amygdala nucleus are excluded from the statistical analysis that follows. Placements for the remaining animals (ACSF [N = 8], NBQX [N = 13]) are indicated in Figure 1 by open triangles (point down). As shown in Figure 3 (rightmost bars), NBQX significantly disrupted fear-potentiated startle to the visual CS, *t*(19) = 2.26.

Because we were able to demonstrate an involvement of the medial amygdala in fear-potentiated startle using NBQX and because there was not an APS-induced disruption of fear conditioning with pre-training infusions, we did not also assess the involvement of the medial amygdala in fear-potentiated startle with APS.

**Treatment effects on “baseline” startle amplitude**

As also indicated in Figure 3, baseline startle amplitude (i.e., on noise-alone trials) was markedly lower in rats that received pre-test NBQX infusions into the medial nucleus of the amygdala compared to rats that received ACSF infusions. In fact, an ANOVA on baseline startle amplitude using rats from Experiments 4A together with those of 4B indicated a significant effect of drug treatment (ACSF vs. NBQX), *F*(1,34) = 16.02, but not modality (rats from Experiment 4A vs. 4B).

Because rats were trained and tested in the same chamber, we considered the possibility that startle amplitude was elevated in ACSF-infused rats due to context conditioning and that this effect was blocked by NBQX. For the purest assessment of context conditioning (i.e., one not contaminated by intermixed CS test trials), we compared startle amplitude on the 30 noise-alone test trials that occurred at the beginning of the pre-shock acclimation session with startle amplitude on the 30 noise-alone test trials that occurred at the beginning of the post-conditioning test session (i.e., the noise-alone test trials that preceded the first CS presentation; see Methods section for details). Indeed, the startle amplitude of ACSF- but not NBQX-infused rats increased markedly from the pre-conditioning acclimation session to the post-conditioning test session (see Fig. 4, leftmost bars). Statistically, an ANOVA on the between-session percent change scores indicated a significant Treatment effect, *F*(1,34) = 4.93, but not a significant Modality or interaction effect. The absolute level of startle prior to conditioning was somewhat, although not significantly, lower in the groups that later received NBQX (i.e., vs. the groups that later received ACSF). We do not believe that this in any way accounts for the apparent disruption of context conditioning insofar as we have previously found that percent

Figure 2. Effect on fear conditioning and fear-potentiated startle of glutamate receptor antagonist infusions into the basolateral amygdala. The NMDA receptor antagonist AP5 disrupted fear conditioning to an olfactory CS when infused prior to training (leftmost bars). The AMPA/kainate receptor antagonist NBQX did disrupt fear-potentiated startle when infused prior to testing (rightmost bars). Baseline startle amplitude (i.e., from noise-alone test trials), from which percent-potentiation scores are derived, is indicated immediately below each bar.

* * *
conditioning acclimation session, from which percent potentiation scores did not disrupt these increases, nor did pre-training infusions of AP5 are derived, is indicated immediately bars). In contrast, pre-test NBQX infusions into the basolateral amygdala (treatments shown in Experiments 1A and 2 to disrupt fear conditioning changes in baseline startle amplitude. Pre-test infusions of NBQX into the medial nucleus of the amygdala disrupted pre- to post-conditioning increases of startle amplitude on noise-alone trials (leftmost bars). In contrast, pre-test NBQX infusions into the basolateral amygdala did not disrupt these increases, nor did pre-training infusions of APS (rightmost bars). Baseline startle amplitude from the final pre-acclimation session, from which percent potentiation scores are derived, is indicated immediately below each bar.

For comparison, we also evaluated the effect of pre-test NBQX and pre-training APS infusions into the basolateral amygdala (treatments shown in Experiments 1A and 2 to disrupt fear conditioning and fear-potentiated startle to the discrete olfactory CS) on pre- to post-conditioning changes in baseline startle amplitude. Surprisingly, although these treatments potently disrupted olfactory fear conditioning and fear-potentiated startle, they had no obvious effect on the pre- to post-conditioning increases (see Fig. 4, right bars).

To ensure that these increases do in fact reflect context conditioning as opposed, for example, to a nonspecific shock-induced sensitization of startle amplitude, other rats were tested as before, but received training in a distinctively different chamber (see Methods section for details). Although these rats received the same number, intensity, and schedule of shocks as in our other experiments, they did not show the same pre-training to post-training increase. In fact, mean startle amplitude remained remarkably stable (i.e., 239.0 ± 37.6 prior to conditioning and 235.4 ± 46.7 after conditioning).

NBQX infused into the medial nucleus of the amygdala might have specifically disrupted the influence of context on startle, but might alternatively have depressed startle nonselectively and simply masked the influence of context. To discriminate between these alternatives, we evaluated the effect of NBQX infusions in untrained animals. Nine rats were implanted with cannula aimed at the medial nucleus of the amygdala. One week later, these rats were placed into the startle chamber whereupon they received, after a 5 min acclimation period, 30 95-dB noise bursts and 30 intermixed 110-dB noise bursts. Rats received infusions of either ACSF or 3 µg/side NBQX (the order of treatments was counterbalanced) 24 and 72 h later and were retested using the same protocol. The 110-dB noise bursts were included to ensure that a negative finding could not be attributed to baseline-dependent NBQX effects (e.g., an effect that was only apparent at higher startle amplitudes such as those that would have occurred to 95-dB stimuli in conditioned rats). Five rats were found to have misplaced cannula. Results from the remaining animals are shown in Figure 5. NBQX infusions did not decrease startle amplitude to either 95- or 110-dB noise bursts. Thus, the lower startle levels in NBQX- compared with ACSF-infused rats from Experiments 4A and 4B do not appear to reflect a general decrease of startle amplitude but reflect, instead, a specific disruption of the pre- to post-conditioning increase.

### Influence of cannula placement on the magnitude of the NBQX effect

Because the medial amygdala lies just medial to the basomedial and cortical amygdala nuclei, and because drugs infused into the medial nucleus might diffuse dorsally along the cannula track where they might also influence the central nucleus of the amygdala (where NBQX infusions have previously been shown to disrupt fear-potentiated startle to a visual CS (Walker and Davis 1997), we considered the possibility that the effects of pre-test infusions into the medial nucleus might be mediated by actions elsewhere.

For fear-potentiated startle to the olfactory stimulus (Experiment 4A), there were 10 NBQX-infused rats with bilateral placements in or sufficiently near the medial nucleus of the amygdala to be scored as hits, and four rats with only a single cannula in the medial amygdala and a second cannula elsewhere. As shown in Figure 6 (left), fear-potentiated startle to the olfactory CS in rats with bilateral hits was significantly lower, t(12) = 3.21, than in rats that did not have bilaterally accurate placements. Thus, the effect of NBQX on fear-potentiated startle to an olfactory CS was anatomically specific and appeared to require bilateral inactivation of the medial nucleus of the amygdala per se.

We also looked for placement-effect correlations using data from all of the NBQX-infused rats in Experiment 4A. For these analyses, the distance of one cannula tip to either the medial, basomedial, cortical, or central amygdala nuclei was added to the distance of the other cannula to the same structure on the opposite side of the brain, and this summed distance was entered into a correlation analysis (i.e., one Pearson Product-Moment Correlation for each of four structures) using percent potentiation as the correlated variable. Percent potentiation scores were significantly correlated (r = 0.82) with cannula tip distance to the medial nucleus of the amygdala (see Fig. 6, right), but were not significantly correlated with cannula tip distance from the other three structures (r’s = 0.15, −0.29, and −0.45 for the central, cortical, and basomedial nuclei, respectively).

For fear-potentiated startle to the visual stimulus (Experiment 4B), percent potentiation scores were not significantly correlated with proximity to any of the four structures considered (there were too few animals with missed placements to mean-

### Figure 5. Pre-test infusions of NBQX into the medial nucleus of the amygdala did not influence startle amplitude in untrained rats.
right of fear-potentiated startle to the olfactory CS (nucleus of the amygdala was significantly correlated with the magnitude of fear-potentiated startle to the olfactory CS (right). *P < 0.05 vs. ACSF control infusions.

Figure 6. Pre-test NBQX infusions disrupt fear-potentiated startle to an olfactory CS via actions within the medial nucleus of the amygdala. For all NBQX-infused rats from Experiment 4, percent potentiation scores were significantly lower for rats that received bilateral infusions into the medial nucleus of the amygdala (MeA) compared to rats that received one infusion into the medial nucleus and a second infusion elsewhere (left). For these same animals, the distance of the infusion sites from the medial nucleus of the amygdala was significantly correlated with the magnitude of fear-potentiated startle to the olfactory CS (right). *P < 0.05 vs. ACSF control infusions.

The three goals of the present study were (1) to evaluate the contribution of the medial nucleus of the amygdala (MeA) to fear-potentiated startle to an olfactory CS, as well as fear expression to a visual and contextual CS. We did not, however, find evidence for a role of the basolateral amygdala in fear conditioning. Because the same CS pathway presumably participates in both, the disruption of performance by pre-test NBQX infusions cannot readily be attributed to a disruption of neural transmission in CS pathways. A more plausible view is that the medial nucleus is either an embedded component of the conditioned response pathway, or an indirect modulator of that pathway. With regard to the former possibility, the medial nucleus projects to the bed nucleus of the stria terminalis (Canières et al. 1995; Dong et al. 2001) and the bed nucleus of the stria terminalis projects to the central nucleus of the amygdala (e.g., Dong et al. 2000, 2001). Both project to the primary acoustic startle circuit (Rosen et al. 1991; Koch and Ebert 1993; Fort et al. 1994, 1997; Frankland and Yekmons 1995; Koch and Schnitzerl 1997; Shi and Davis 2002) and have been implicated in the potentiation of startle by fear and anxiogenic stimuli (e.g., Koch and Schnitzler 1997; Lee and Davis 1997; Rosen and Davis 1998). As a recipient of projections from the basolateral amygdala (cf., Pitkanen 2000), the medial nucleus is potentially well-positioned to mediate the influence of fear on startle. The medial nucleus might also influence anxiety more generally. Duxon et al. (1997) found that infusions of a 5-HT2B receptor agonist into the medial amygdala nucleus reduced indicators of anxiety in the social interaction test although, in the same study, 5-HT2B activation was not anxiolytic in the Vogel punished drinking test.

Analyses of the relation between cannula placement and the effect of these infusions suggested that the disruption of fear-potentiated startle to the olfactory CS was indeed attributable to actions within or very near to the medial amygdala nucleus. First, whereas bilateral placements in the medial nucleus nearly abolished fear-potentiated startle to the olfactory CS, unilateral placements (one cannula in the medial nucleus and a second cannula elsewhere that did not satisfy our inclusion criteria) had no ob-
vious effect. Thus, the behavioral effects were anatomically specific. Moreover, for odor-potentiated startle, a significant correlation was observed between the behavioral effect and the distance of the cannula from the medial nucleus. Perhaps significantly, this was not the case for fear-potentiated startle to the visual CS, or for the increases in startle amplitude that occurred from the pre-conditioning acclimation session to the post-conditioning test session. For these stimuli, infusions elsewhere into the ventromedial amygdala were also effective and there was no such correlation. Thus, the role of the medial nucleus in fear-potentiated startle to nonolfactory CSs remains uncertain.

In light of a previous finding that NBQX infusions directly into the central nucleus of the amygdala disrupt fear-potentiated startle to visual CSs (Walker and Davis 1997), the possibility that the NBQX effects on odor-potentiated startle observed in the present study were due to actions within the central nucleus merits additional consideration. Recent unpublished findings from our laboratory are relevant. Using procedures similar to those used here (see Shi and Davis 1999 for details), C.-J. Shi and M. Davis (unpubl.) found that unilateral NBQX infusions into the medial nucleus of the amygdala did not disrupt fear-potentiated startle to a visual CS (see Fig. 7, left). In contrast, Z. Zhao and M. Davis (unpubl.), using the same behavioral procedures and a similar startle apparatus (see Meloni and Davis 2000 for details), found that unilateral infusions into the central nucleus of the amygdala did abolish fear-potentiated startle to a visual CS (see Fig. 7, right). Hence, if the blockade of fear-potentiated startle to a visual CS by NBQX infused into the medial nucleus was due to diffusion to the central nucleus, one might expect that this blockade would also have occurred with unilateral infusions into the medial nucleus of the amygdala. Because this did not occur, we are doubtful that the effects of NBQX in Experiment 4B are attributable to actions within the central nucleus of the amygdala. We have also found that NBQX infusions into the central amygdala nucleus disrupt fear conditioning (pre-training infusions) to a visual CS as well as fear-potentiated startle (pre-test infusions). In the present study, NBQX infusions into the medial nucleus disrupted fear-potentiated startle but did not disrupt fear conditioning. Again, this dissociation is most compatible with the view that our NBQX effects on odor-potentiated startle are not due to diffusion to the medial nucleus.

A comparison with results from lesion studies may also be informative insofar as the behavioral effects of lesions can be attributed with great precision to damage to within a circumscribed area. Pre- as well as post-training excitotoxic lesions of the medial nucleus blocked freezing to a context previously paired with predator odor in Takahashi et al. (2004) but the same group found only a nonsignificant effect of pre-training lesions on freezing to a context that had been paired with footshock (Li et al. 2004). Pre-training excitotoxic lesions of the medial amygdala nucleus also blocked freezing to a context previously paired with predator odor in Markham et al. (2004) but, again, did not significantly disrupt freezing to a context previously paired with footshock. Pre-training electrolytic lesions incompletely disrupted freezing to a footshock-paired context in Holahan and White (2002), and pre-training excitotoxic lesions had no effect on conditioned freezing to an auditory CS in Nader et al. (2001). Although the designs of these studies are not strictly comparable to our own, the general pattern of results appears to be similar. Whether an odor is used as a CS, as in our study, or as a unconditioned stimulus (US), as in Takahashi et al. (2004) and Markham et al. (2004), there is considerable consistency regarding the importance of the medial amygdala nucleus in fear-conditioning paradigms that use odor. For fear-conditioning paradigms that use nonolfactory stimuli, the results appear to be less consistent.

As indicated elsewhere, we also observed that baseline startle amplitude increased markedly from the pre-conditioning acclimation session to the post-conditioning test session and that these increases were context-dependent (see also McNish et al. 1997). Context-potentiated startle was not observed in rats that received pre-test NBQX infusions into the medial amygdala nucleus. Most surprisingly, however, context conditioning and context-potentiated startle appeared to be completely intact in rats that received pre-test NBQX or pre-training AP5 infusions into the basolateral amygdala. The disruption of odor-potentiated startle in the same animals and in the same test session following basolateral amygdala infusions provides an important positive control and limits the range of possible interpretations (i.e., insufficient dose or inaccurate placement) for the failure to also disrupt context-potentiated startle. Although many studies have shown that contextual fear conditioning and conditioned fear to contextual cues can be disrupted by treatments that interfere with basolateral amygdala function (e.g., Fanselow and Kim 1994; Helmstetter and Bellgowan 1994; Maren et al. 1997; Goosens and Maren 2001, 2003; Rodriguez et al. 2001; Wallace and Rosen 2001) our results are not wholly without precedent.

Maren (1999) found that although basolateral amygdala lesions slowed the emergence of freezing to a context CS, asymptotic levels after additional training were comparable to those observed in sham-lesioned controls. In the same animals, basolateral amygdala lesions completely blocked freezing to an auditory CS and this deficit was not overcome by additional training (see also Cahill et al. 2000; Berlau and McGaugh 2003). Selden et al. (1991) found that pre-training basolateral amygdala lesions disrupted lick suppression produced by an auditory CS, but did not disrupt avoidance by the same animals of the compartment in which they had previously received shocks. In our own laboratory, we recently noted that rats that received AP5 infusions into the basolateral amygdala prior to 30 light-shock pairings showed no evidence of fear conditioning to the visual cue (i.e., on CS-noise trials) and initially showed no evidence of fear conditioning to the context (i.e., on noise-alone trials). During testing however, startle amplitude on both trial types increased abruptly and to a comparable level once the first light stimulus was presented (see Walker and Davis 2002b). We believe that the visual stimulus in that study may have acted as a salient element.

![Figure 7](https://www.learnmem.org)
of the context rather than as a CS, triggering latent contextual fear that had been disrupted but not blocked by pre-training AP5 infusions. Even in those studies in which amygdala manipulations have been shown to impair context conditioning or fear expression, when context and discrete cue-elicited fear have been measured in the same study, the effects on the former are often weaker than on the latter (e.g. Helmstetter and Bellgowan 1994; Rodrigues et al. 2001).

We are uncertain what contributes to the variable effects of amygdala manipulations on context conditioning and fear expression. One possibility, for example, is that context trained as a foreground stimulus (i.e., paired with shock in the absence of an explicit discrete CS) may be more or less susceptible to amygdala manipulations than context trained as a background stimulus. However, amygdala manipulations have been found to disrupt fear behaviors elicited by contexts trained as foreground (Fanselow and Kim 1994) as well as background (e.g., Muller et al. 1997) stimuli and, in other studies, have been found not to disrupt fear behaviors elicited by contexts trained as foreground (Maren 1999) or as background (present results) stimuli.

With specific reference to our findings, it is also possible that receptors within the amygdala other than those targeted mediate contextual processes. Perhaps, for example, short-duration responses to short-duration CSs are mediated by ionotropic glutamatergic receptors within the basolateral amygdala, whereas longer duration responses to longer duration CSs (e.g., context) are mediated by peptide or metabotropic receptors.

A differential involvement of the amygdala in context versus discrete cue conditioning may be relevant to the long-standing debate as to whether the amygdala is a critical site of neural plasticity in aversive learning paradigms (e.g., Fanselow and LeDoux 1999) or, instead, a brain area that modulates memory storage elsewhere (e.g., Cahill et al. 1999). Evidence for the latter view comes largely from studies that have found incomplete effects of amygdala manipulations on tasks that have assessed retention using avoidance of shock-associated contexts as the primary behavioral measure (e.g. Berlau and McGaugh 2003; Vazdarjanova and McGaugh 1998). From the standpoint of the present discussion, it would be interesting to compare directly the influence of amygdala manipulations on avoidance of shock-paired contexts with avoidance elicited by discrete unimodal CSs.

In summary, our results indicate that the basolateral subdivision of the amygdala plays an important role in fear-potentiated startle and in fear conditioning to olfactory cues. The results also implicate the medial nucleus of the amygdala in fear expression, at least to olfactory cues. We also found that context conditioning and fear-potentiated startle to a shock-associated context was not disrupted by basolateral amygdala manipulations that did disrupt fear conditioning and expression to an olfactory CS.

Materials and Methods

Animals

Male Sprague-Dawley rats (Charles River) were maintained on a 12-h light/12-h dark cycle (lights on at 0800 h) with food and water available ad libitum. Rats were initially group-housed in 45 × 20 × 24-cm (depth × width × height) polycarbonate cages (four rats per cage) and individually housed, following surgery, in 20 × 19 × 24-cm hanging wire-mesh cages. At surgery, rat weights ranged from 275 to 350 g.

Apparatus

Rats were trained and tested in two identical 9 × 14 × 10-cm (depth × width × height; internal dimensions) Plexiglas and wire-mesh cages, each suspended between compression springs within a steel frame and located, together, within a custom-designed 60 × 79.5 × 59.5-cm sound-attenuating chamber lined with 6.3 mm thick Plexiglas. The floor of each cage consisted of four 6.0-mm diameter stainless steel bars spaced 18 mm apart.

Affixed to the bottom of each cage was an Endevco accelerometer (Model 2217E). Cage movement (e.g., produced by the rat’s startle response) resulted in displacement of the accelerometer that, in turn, produced a voltage output proportional to the velocity of cage movement. The accelerometer’s output was amplified by an Endevco Model 104 amplifier and recorded on a scale of 0–2500 units by an InstruNET device (GW Instruments, Model 100B) interfaced to a Macintosh G3 computer. Startle amplitude was defined as the maximal peak-to-peak voltage that occurred during the first 300 msec after onset of the startle-eliciting noise burst.

Startle responses were evoked by 50-msec 95-dB white-noise bursts (5-msec rise-decay time, 0–22 kHz) generated by a Macintosh G3 computer soundfile, amplified by a Radio Shack amplifier (Model MPA-200), and delivered through high frequency speakers (Radio Shack Supertweeter) located 8 cm in front of each cage. Background noise (60-dB wideband) was produced by an ACO Pacific white-noise generator (Model 3024) and was delivered through the same speakers as those used to provide white-noise bursts. Sound level measurements were made with a Brüel & Kjær model 2235 sound-level meter (A scale; random input) with the microphone (Type 4176) located 10 cm from the center of the speaker, which approximates the distance of the rat’s ear from the speaker during testing.

The olfactory CS and the means for delivering it have been described in detail elsewhere (Paschall and Davis 2002). In brief, a continuous flow of air was delivered from a compressed-air cylinder at a rate of 1.0 L/min through a small port (1.3-mm lumen diameter) positioned just above a 1.25-mm diameter opening in the top of each cage. For delivery of the olfactory stimulus, a computer-controlled solenoid (Model H15–03, Coulbourn Instruments) was opened for 4 sec, thereby diverting clean air from the compressed-air cylinder into and through a scaled 135-cm³ glass jar that contained 20 ml of 5% (vol/vol) amyl acetate (i.e., the odorant) in propylene glycol solution. The inlet and outlet ports of the glass jar were positioned above the solution such that clean air from the tank mixed with the amyl acetate-containing vapor. The output was then mixed in a 3:5 ratio with clean air before flowing into the cage. To deflect air that would otherwise flow directly onto the rat’s back, a 2.5 × 2.5-cm sheet-metal plate was positioned just inside the test cage, slightly below the port through which air was delivered.

The chamber (0.284 m³ total volume) was actively exhausted into the building’s ventilation system at a rate of 0.0114 m³/sec. Thus, a volume of air equal to the chamber volume was exhausted into the building’s ventilation system at a rate of 0.0114 m³/sec. Thus, a volume of air equal to the chamber volume was vented every 25 sec. Previous results with fear-conditioned rats indicate that with these procedures startle amplitude returns to baseline levels within 30 sec of solenoid closure (Paschall and Davis 2002). Cages were cleaned daily with warm tap water and 95% alcohol, and were air dried overnight.

A 4-sec visual CS (140 lux) was used for Experiment 4B. This CS was produced by an 8-W fluorescent bulb (100-μsec rise time) located 10 cm behind each cage. Luminosity was measured using a VWR light meter.

A context switch experiment was also performed. For this experiment, rats were trained in a 24 × 31 × 26 cm with 16 0.5-cm diameter floor bars (spaced 18 mm apart) with 24-cm metal-link chains hanging freely from a position 11.0 cm diagonal to each corner. During training, background noise was off and a 150 lux houselight was on. The olfactory stimulus was delivered through Pharmed Tygon tubing (3.2 mm inner diameter) that was routed through the ceiling of the training chamber. The end of the tubing hung approximately 7 cm above the floor bars.

The unconditioned stimulus for all experiments was a 0.5-sec 0.4-mA scrambled floorbar shock. Shock intensity was measured with a 1-kW resistor across a differential channel of an oscilloscope in series with a 100-KW resistor connected between

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adjacent floor bars within each cage. Current was defined as the root-mean-square voltage across the 1-kW resistor where mA = 0.707 × 0.5 × peak-to-peak voltage. Shocks were produced by LeHigh Valley shock generators (SGS-004).

The presentation and sequencing of all stimuli was under the control of the Macintosh G3 computer using custom-designed software (The Experimenter; Glassbeads Inc.).

Surgery
Rats were anesthetized with Nembutal (sodium pentobarbital; 50 mg/kg ip) and placed in an ASI, Inc. stereotaxic frame with the nosebar set to −3.8 mm (flat-skull position). The skull was exposed and a 22-gauge guide cannulae (Plastic Products, Model C313G) were lowered bilaterally into the basolateral amygdala (AP = −3.3, DV = −8.2, ML = ± 5.4 mm from bregma) or the medial nucleus of the amygdala (AP = −3.0, DV = −9.5, ML = ± 3.6 mm from bregma). Dummy cannulae (Plastic Products, Model C313DC/l) were inserted into each guide cannula to prevent clogging. The tip of each extended approximately 1 mm past the end of the guide cannula. Jeweler screws were anchored to the skull and the entire assembly was cemented in place using CranioPlastic Powder (Plastic Products). A minimum of 8 d elapsed between surgery and the behavioral procedures.

Drugs and infusion procedure
Rats were infused with the NMDA receptor antagonist D,L-AP5 (Sigma-Aldrich), the AMPA/kainate receptor antagonist NBQX (Tocris Cookson Inc.) disodium salt, or artificial cerebrospinal fluid (ACSF). Infusions (0.25 µL/min, 0.5 µL total volume) were made through 28-gauge injection cannulae (Plastic Products, Model C313I) that were attached by polyethylene tubing to a Hamilton microsyringe. Injection cannulae were left in place for 2 min after the infusion was completed.

Pre-infusion trials were completed immediately before the animals were placed into the test cages. The CS-noise and noise-alone test trials, from which fear-potentiated scores were derived, began 2 min later. For pre-training administration, infusions were completed 15 min before animals were placed into the cages, and CS-shock pairings began 5 min later. Thus, both testing and training began approximately 20 min after the infusion.

Behavioral procedures
Acclimation session
On each of two consecutive days, rats were placed into the test cages and after 5 min, presented with 30 95-dB startle-eliciting noise bursts (onset-to-onset interstimulus interval [ISI] = 30 sec). On the basis of these results, rats were sorted into different treatment groups such that each treatment group began with equivalent mean baseline startle levels.

Fear conditioning
Within three days of the final acclimation session, rats were returned to the same chamber whereupon they received either five odor-shock (Experiments 1, 2, 3, and 4A) or five light-shock (Experiment 4B) pairings. The first pairing occurred 5 min after placement into the startle chamber and successive shocks occurred every 4 min. For each pairing, the 0.5-s sec shock was delivered 3.5 sec after onset of the 4.0-sec CS.

Fear-potentiated startle test
Approximately 48 h later, rats were again returned to the same chamber in which they had previously received shocks. Five minutes later, the first of 30 95-dB startle-eliciting noise bursts was presented. Successive stimuli occurred every 30 sec. Rats then received 40 test trials consisting of 10 repetitions of a single CS test trial followed by three noise-alone test trials. For CS test trials, the 95-dB noise burst was presented 3.5 sec after onset of either the odor (Experiments 1, 2, 3, and 4A) or light (Experiment 4B) CS. For noise-alone test trials, the 95-dB noise burst was presented alone. The ISI for all stimuli was 30 sec.

Statistical analyses
For each rat, the mean startle amplitude on noise-alone and on CS test trials was determined, and a percent change score was calculated (mean startle amplitude on CS-noise test trials divided by the mean startle amplitude on noise-alone test trials × 100 minus 100). Percent change scores were used (i.e., vs. absolute difference scores) because we previously found that they remain stable across large variations in baseline startle amplitude (up to 10-fold in Walker and Davis 2000a). These scores were then analyzed using ANOVA and/or two-tailed t-tests as indicated in the Results section.

Changes in baseline startle from the pre-conditioning acclimation session to the post-conditioning test session, shown elsewhere to reflect context conditioning, were similarly calculated (i.e., mean startle amplitude of the 30 noise-alone trials which occurred at the beginning of the test session divided by the mean startle amplitude of the 30 noise-alone trials of the second acclimation session × 100 minus 100).

For all comparisons, the criterion for significance was P < 0.05.

Histology
Rats were sacrificed by chloral hydrate overdose and perfused intracardially with 0.9% saline (wt/vol) followed by 10% formalin (vol/vol). The brains were removed and immersed in a 30% sucrose-formalin solution (wt/vol) for at least three days, after which 40-µm coronal sections were cut through the area of interest. Every fourth section was mounted and stained with cresyl violet.

Cannula placements, and the determination as to whether the cannula was within or sufficiently near the intended target to be scored as a hit, were judged by a scorer blind to the animal’s group assignment and behavioral data. To be scored as an accurate placement, cannula tips were required to be within or no further than 0.5 mm from the intended target (i.e., either the basolateral amygdala complex [e.g., the lateral, basolateral, and basomedial nuclei] for Experiments 1 and 2, or the medial amygdala nucleus for Experiments 3 and 4). Because white matter tracts have previously been shown to be barrier to diffusion (e.g., Morris et al. 1989), data from animals with cannula tips located on the opposite side of a white matter tract (i.e., cannula intended for the medial amygdala that were medial to the optic tract, or cannula intended for the basolateral amygdala that were lateral to the external capsule) were always excluded unless the guide cannula sheared through the tract thereby allowing for diffusion to the intended target.

With the dose and infusion parameters used in the present study and similar inclusion criteria, we previously reported that NBQX infusions into the basolateral but not the central amygdala disrupt the facilitatory effects of sustained bright light on startle (Walker and Davis 1997). We have also observed that NBQX infusions into the ventral cochlear nucleus disrupt tone-evoked compound action potentials only when the cannula tips are within 1.0 mm of the ventral cochlear nucleus (E.G. Meloni and M. Davis, unpubl.). Similarly, with the same dose of AP5 as that used here, albeit with a nominally smaller infusion volume (0.5 µl vs. 0.5 µl), Fanselow and Kim (1994) reported that D,L-AP5 disrupted fear conditioning when infused into the basolateral amygdala but not when infused into the central amygdala nucleus. Thus, we believe that these procedures and inclusion criteria are appropriate for evaluating the contribution of the basolateral and medial amygdala to fear acquisition and potentiated startle.

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