Amygdala response to smoking-cessation messages mediates the effects of serotonin transporter gene variation on quitting

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A B S T R A C T

The amygdala is critically involved in detecting emotionally salient stimuli and in enhancing memory for emotional information. Growing evidence also suggests that the amygdala plays a crucial role in addiction, perhaps by strengthening associations between emotionally-charged drug cues and drug-seeking behavior. In the current study, by integrating functional MRI (fMRI), genetics, and outcome data from a large group of smokers who completed a smoking-cessation intervention and attempted to quit, we show that the amygdala also plays a role in quitting. Specifically, we demonstrate that the amygdala response to smoking-cessation messages in smokers trying to quit is a predictor of their post-intervention quitting outcome. We further show that the amygdala response is modulated by genetic variation in the serotonin transporter and mediates the impact of this genetic variation on quitting. These results point to a gene–brain–behavior pathway relevant to smoking cessation, and add to our understanding of the role of the amygdala in nicotine addiction.

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Introduction

Cigarette smoking continues to be a leading preventable cause of morbidity and mortality in the U.S. (CDC, 2003, 2008). Many smokers attempt to quit but a majority relapse within 6 months (Quaak et al., 2009), highlighting a vital need for more effective interventions. Identification of brain predictors of relapse susceptibility could inform the design and selection of smoking-cessation interventions in the clinic, while also adding to our understanding of brain processes underlying nicotine addiction.

The goal of the current study was to examine the role of amygdala response to smoking-cessation messages in quitting as well as the genetic sources of variability in amygdala response. The amygdala plays a critical role in emotion processing, including response to emotionally salient stimuli and acquisition of emotional learning (LeDoux, 2000; Phelps, 2006). Emotionally salient information tends to be remembered better than emotionally neutral information (Cahill and McGaugh, 1995), and both animal and human studies suggest that the amygdala is required for this enhanced learning (McGaugh, 2004). For drug users, drug-related cues are emotionally arousing and are known to activate the amygdala (Childress et al., 1999; Franklin et al., 2007; Kilts et al., 2001). Therefore, drug-related information is more likely to be remembered by them and influence their behavior. Consistent with this view, growing evidence suggests that the amygdala is involved in drug addiction, by signaling the salience of drug-related cues and creating strong associations with drug-seeking behaviors (Robbins et al., 2008; See et al., 2003).

Previous neuroimaging studies in smokers have shown the link between amygdala response to smoking-related cues on the one side, and craving intensity (Franklin et al., 2011; Goudriaan et al., 2010) and relapse susceptibility (Janes et al., 2010) on the other side. In contrast, amygdala response to smoking-cessation messages has not yet been examined. We hypothesized that the amygdala may also play an important role in how smokers respond to smoking-cessation messages and how these messages influence their smoking behavior. Specifically, we expected that a greater amygdala response to smoking-cessation messages should be associated with enhanced memory of these messages, and therefore increased odds of quitting success. To test our hypothesis, we integrated functional MRI (fMRI) and outcome data from a large group of smokers who completed a smoking-cessation intervention, in order to examine whether the amygdala response to smoking-cessation messages predicted subsequent quitting outcome in these smokers.

We also investigated the genetic factors that may account for individual differences in amygdala response to smoking-cessation messages, using an imaging genetics approach (Bigos and Weinberger,
nucleotide substitution (rs25531), with the LG allele being functional—Lesch et al., 1996). In addition, the 5-HTTLPR includes an A→G single nucleotide substitution (rs25531), with the long allele (L) less efficiently transcribed than the long allele (L) (Heils et al., 1996; Lesch et al., 1996). In addition, the STin2 polymorphism —modulated the amygdala response. The 5-HTTLPR is a 44-base pair insertion/deletion polymorphism in the promoter region, with the short allele (S) less efficiently transcribed than the long allele (L) (Heils et al., 1996; Lesch et al., 1996). Based on previous evidence (Dannlowski et al., 2010; Hariri et al., 2002, 2005), we hypothesized that low-transcription alleles of the 5-HTTLPR/rs25531 and STin2 would be associated with a relatively increased amygdala response compared to high-transcription alleles.

Materials and methods

Subjects

Ninety-one smokers interested in quitting (44 females and 47 males, mean age = 37.5 years, mean number of cigarettes per day = 16.7) were recruited for the study (Chua et al., 2011), which included a web-based tailored smoking-cessation intervention developed at the University of Michigan’s Center for Health Communications Research (Strecher et al., 2008). Smokers were eligible to participate if they smoked a minimum of 10 cigarettes per day and at least 100 cigarettes in their lifetime. Subjects were not enrolled in other smoking-cessation programs or taking pharmaceutical treatments for smoking cessation during study enrollment. All subjects were native English speakers, had normal vision and hearing, and had no history of head injury or mental illness. The study protocol was approved by the University of Michigan Medical School IRB and all subjects provided written informed consent.

Study design

The study involved 3 sessions plus a follow-up phone interview. In Session 1, subjects completed a baseline assessment of their smoking history and other health, demographic, and psychosocial characteristics relevant to smoking cessation. The responses were used to create tailored smoking-cessation messages for the subsequent intervention. In Session 2, subjects completed a Messages Task during functional MRI (fMRI). In Session 3, scheduled within one week of their fMRI session, subjects completed a web-based computer-tailored smoking-cessation intervention (Strecher et al., 2008) and started their quit attempt. All subjects also provided saliva for DNA extraction and genotyping, and received a 10-week supply of nicotine patches to help them quit. Four months after the intervention session, subjects were interviewed on the phone to determine their smoking-cessation outcome. The primary outcome measure was 7-day point-prevalence abstinence (cigarette free for the past 7 days), a metric that is highly correlated with both 24-hour point-prevalence abstinence and 30-day prolonged abstinence, as well as with physiological measures of smoking cessation (Velicer and Prochaska, 2004).

Messages Task

All subjects completed 5 runs of the Messages Task (Chua et al., 2011) during fMRI, with 2 blocks of each of the three types of messages (tailored smoking-cessation messages, untailored smoking-cessation messages, and neutral control messages) per run, 5 messages per block, for the total number of 150 messages. Each block lasted 24 s. The order of blocks in the Messages Task was pseudo-randomized within and across participants. Blocks were separated by fixations lasting 4–10 s (an average of 7 seconds). The messages were presented visually on the screen and simultaneously presented as an audio track. Subjects were instructed to pay attention to the messages but no response was required. Tailored smoking-cessation messages were created on the basis of the individual’s responses during a baseline assessment and made references to this person’s characteristics, experiences, and specific obstacles to quitting. Therefore, although of the same type, tailored smoking-cessation messages varied across subjects. Examples of tailored smoking-cessation messages include: You want to quit because you are tired of spending your money on cigarettes; You feel like your friend will help you stay on track once you quit; You have a very strong urge to smoke when you first wake. In contrast, untailored smoking-cessation messages were relevant to smokers in general and were identical for all subjects. Examples of untailored smoking-cessation messages include: Many people quit with another person so they can support each other; Many people relapse due to stress, alcohol, and cravings; Most people need to try more than once to quit smoking for good. Neutral control messages (i.e., world-knowledge messages not related to smoking) were also the same for all subjects. Examples of neutral messages include: The longest duration of a solar eclipse was 7 minutes, 31 seconds; Bali attracts more tourists than any other Indonesian island; Global warming caused the recent collapse of an Antarctic ice shelf.

Image acquisition

Scanning was performed on a 3T GE Signa Excite 2 scanner (Milwaukee, Wisconsin), beginning with a structural T1-overlay image (repetition time [TR]=250 ms, echo time [TE]=7 ms, flip angle [FA]=75 degree, field of view [FOV]=220 mm, 43 oblique axial slices, 256×256, slice thickness 3.0 mm). Functional scans were collected using a T2*—weighted spiral-in acquisition sequence (gradient echo, TR=2000 ms, TE=30 ms, FA=90°, FOV=220 mm, 64×64, slice thickness 3.0 mm) (Noll et al., 1998). High-resolution T1 scans were also obtained for precise anatomical localization (3D spoiled-gradient echo [3D-SPGR] with inversion recovery prep, time of inversion = 400 ms, TR=9.0 ms, TE=1.8 ms, FA=15°, FOV=260 mm, 128 slices, 256×256, 1.2 mm slice).

Image preprocessing

All functional scans were slice-time-corrected, motion-corrected, and realigned to the first scan using the MCFLIRT program (FSL Analysis Group, FMRIB, Oxford, UK). Subsequent processing was done using SPM (Wellcome Institute of Cognitive Neurology, London, UK). The T1-overlay was co-registered with a functional scan. The high-resolution 3D-SPGR image was co-registered to the T1-overlay...
and anatomically normalized to the Montreal Neurological Institute (MNI) 152 template. The resulting transformation parameters were then applied to the co-registered functional volumes. All functional volumes were smoothed with an overall Gaussian kernel with a full width at half maximum (FWHM) of 7 mm.

Genotyping procedures

Genomic DNA was obtained from saliva samples using Oragene collection system and extracted using the protocol provided (Genotek, Ontario, Canada). STin2 and 5-HTTLPR/rs25531 polymorphisms were genotyped using polymerase chain reaction and oligonucleotide primers. 5-HTTLPR was genotyped using primers from Yonan et al. (2006). PCR products were analyzed on 1.5% TBE agarose gels (expected band sizes: S—415 bp, L—459 bp). To additionally genotype the A–G SNP (rs25531), product was digested withMspI. Digest products were resolved on 3% TBE agarose gels (expected band sizes: L_A, L_B, L_C—157 bp, 174 bp, 66 bp, and 62 bp; S—287 bp, 66 bp, and 62 bp). STin2 was genotyped using primers from Kaiser et al. (2001). PCR products were resolved for size on 1.5% agarose gels. Gels were visualized using ethidium bromide under UV light and reviewed by two independent people, with 100% concordance. Six of the samples were verified by Sanger sequencing, with 100% concordance.

Statistical analyses

After pre-processing, the functional data were analyzed using a modified General Linear Model (GLM) and a blocked design. Regressors of interests (tailored smoking-cessation messages, untailored smoking-cessation messages, and neutral messages) were convolved with a canonical hemodynamic response function (HRF) with a time derivative to account for between-subject and between-voxel variability in the response peak. Movement parameters were included as covariates. Statistical analyses were conducted in a series of steps using a random-effects model in SPM5. First, anatomically defined ROI masks of right and left amygdalae were constructed using WFU PickAtlas (Maldjian et al., 2003) (right amygdala: 87 voxels; left amygdala: 75 voxels). For each individual subject, we extracted the average parameter estimates (betas) for the tailored smoking-cessation messages–neutral messages contrast and the untailored smoking-cessation messages–neutral messages contrast for right and left amygdalae ROIs. These individual parameter estimates, together with genotyping and outcome data, were then entered into second-level group analyses in SPSS 17.0. Race (Caucasian or not), gender, pre-intervention number of cigarettes smoked per day in Non-Quitters compared to Quitters (p = 0.053) (Supplementary Table 1).

Amygdala response to smoking-cessation messages predicts quitting

To test for emotional enhancement memory effect for smoking-cessation messages, and to verify that the subjects paid attention during the Messages Task, we gave them a surprise memory test after they left the scanner (see Supplementary Results). Both tailored and untailored smoking-cessation messages were remembered more accurately than neutral messages (p < 0.01), supporting an emotional enhancement effect for smoking-cessation messages.

We assessed the amygdala response to tailored and untailored smoking-cessation messages compared to neutral messages using blood-oxygenation-level dependent (BOLD) signal and the smoking-cessation messages > neutral messages contrast values extracted from anatomically defined right and left amygdala masks (Maldjian et al., 2003). To control for other potential sources of variability in susceptibility to relapse following a quit attempt, we included race, gender, pre-intervention cigarettes smoked per day, and post-intervention use of nicotine patches as nuisance covariates in all analyses.

Consistent with our main hypothesis, the mean amygdala response to smoking-cessation messages (averaged across right and left amygdalae) was a significant predictor of subsequent post-intervention quitting outcome (B = 1.17, SE = 0.40, Wald χ² = 8.32, p = 0.004, odds ratio = 3.21) (see Supplementary Table 2). A greater magnitude of the mean amygdala response to smoking-cessation messages predicted better odds of quitting success. An omnibus test of model coefficients indicated that the model was significant (p = 0.001). The model explained 23% of the variance in quitting outcome (R square = 0.23) and correctly classified 33 out of 45 Quitters (73.3% accuracy). When tested with separate models, both the right and left amygdala responses to smoking-cessation messages also predicted subsequent quitting (right amygdala: B = 1.05, SE = 0.37, Wald χ² = 8.31, p = 0.004, odds ratio = 2.87, classification accuracy = 73.3% of Quitters, Supplementary Table 3; left amygdala: B = 0.73, SE = 0.30, Wald χ² = 5.84, p = 0.016, odds ratio = 2.08, classification accuracy = 75.6% of Quitters, Supplementary Table 4).

In addition, when we examined tailored and untailored smoking-cessation messages separately, the mean amygdala response to both types of messages was predictive of subsequent quitting (tailored messages: B = 0.96, SE = 0.33, Wald χ² = 8.34, p = 0.004, odds ratio = 2.62, classification accuracy = 73.3% of Quitters, Supplementary Table 5; untailored messages: B = 0.63, SE = 0.28, Wald χ² = 5.11, p = 0.024, odds ratio = 1.88, classification accuracy = 68.5% of Quitters, Supplementary Table 6). Both models were significant (p < 0.008) and explained 21% (tailored messages: R square = 0.21) and 18% (untailored messages: R square = 0.18) of the variance in quitting outcome, respectively.

Genotyping results

Next, we tested for modulation of amygdala response to smoking-cessation messages by genetic variation in the 5-HTT. Subjects provided saliva for DNA extraction and were genotyped for two common, functional polymorphisms in the regulatory regions of the 5-HTT gene (SLC6A4), 5-HTTLPR/rs25531 in the promoter and STin2 in intron 2.

We observed the following distribution of STin2 genotypes: 12 participants were 10/10 homozygotes, 38 were 10/12 heterozygotes, and 32 were 12/12 homozygotes. The observed 5-HTTLPR/rs22531 polymorphism was functionally grouped as follows: 24 participants were LG/LG homozygotes, 41 were LG/LS heterozygotes (i.e., LG/S or LG/LC), and 17 were LG/S, LS homozygotes (i.e., S/S, LC/S or LC/LC). No deviations from the Hardy–Weinberg equilibrium were observed. The genotyping results are given in Supplementary Table 7.
Because the STin2 and 5-HTTLPR/rs25531 polymorphisms both affect the rate of transcription and may have combined effects on transcription efficiency (Hranilovic et al., 2004), both genotypes were included in the initial linear regression models to test if either genotype predicted amygdala response to smoking-cessation messages. Consistent with previous reports (Gelernter et al., 1999; Kazantzseva et al., 2008), the STin2 and 5-HTTLPR/rs25531 loci were in linkage disequilibrium, i.e., the genotypes at the two loci were significantly associated with each other ($\chi^2 = 27.80, p < 0.0001$) (Table 1). Neither the STin2 genotype groups nor the 5-HTTLPR/rs22531 genotype groups differed in pre-intervention measures, except for a higher confidence in quitting in the LA/LA genotype group compared to the LA/LGS carriers ($t = 2.47, p = 0.015$) (Supplementary Tables 8 and 9).

**Table 1**

<table>
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<th>Linkage disequilibrium between the STin2 and 5-HTTLPR/rs25531 loci.</th>
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The STin2 genotype was a significant predictor of the mean amygdala response to smoking-cessation messages, when controlling for race, gender, pre-quit smoking severity, post-intervention use of nicotine patches, and for the 5-HTTLPR/rs25531 genotype ($B = 0.35, SE = 0.017, t = 2.06, p = 0.042$) (see Supplementary Table 10). The model including both STin2 and 5-HTTLPR/rs25531 was significant ($p = 0.002$) and accounted for 23% of the variance in the mean amygdala response to smoking-cessation messages ($R^2 = 0.23$). As hypothesized, a higher number of low-transcription alleles (STin2 10 alleles) predicted a greater magnitude of amygdala response. In contrast, the 5-HTTLPR/rs25531 genotype was not a significant predictor of the mean amygdala response when controlling for the STin2 genotype and the other covariates ($p = 0.188$). Because the genotypes at the two loci were highly associated with each other ($\chi^2 = 27.80, p < 0.0001$) (Table 1), we focused on the STin2 genotype and dropped the 5-HTTLPR/rs25531 from subsequent analyses.

Separate regression models for the right and left amygdalae showed that the STin2 genotype alone significantly predicted the response to smoking-cessation messages both in the right amygdala ($B = 0.50, SE = 0.15, t = 3.30, p = 0.001$) and in the left amygdala ($B = 0.41, SE = 0.18, t = 2.25, p = 0.027$) (see Supplementary Tables 11–12). Both models were significant ($p < 0.01$) and explained 24% of the variance in the right amygdala response ($R^2 = 0.24$) and 18% of the variance in the left amygdala response ($R^2 = 0.18$), respectively.

The STin2 genotype alone was also a significant predictor of the mean amygdala response to tailored smoking-cessation messages ($B = 0.38, SE = 0.15, t = 2.59, p = 0.012$). The model was significant ($p = 0.03$) and explained 15% of the variance in the mean amygdala response ($R^2 = 0.15$). The effects were observed both in the right amygdala ($B = 0.40, SE = 0.16, t = 2.43, p = 0.017$) and in the left amygdala ($B = 0.37, SE = 0.17, t = 2.18, p = 0.032$) when tested with separate models (Supplementary Tables 13–15). The STin2 genotype also significantly predicted the mean amygdala response to untailored messages ($B = 0.53, SE = 0.20, t = 2.60, p = 0.011$). The model was significant ($p = 0.002$) and explained 22% of the variance in the mean amygdala response ($R^2 = 0.22$). For untailored messages, the STin2 effect was significant in the right amygdala ($B = 0.60, SE = 0.19, t = 3.07, p = 0.003$) and trend significant in the left amygdala ($B = 0.46, SE = 0.24, t = 1.89, p = 0.063$) (Supplementary Tables 16–18).

Furthermore, the results for STin2 modulation of the mean amygdala response remained essentially unchanged when we restricted the analyses to Caucasians ($n = 63; 33$ females and $30$ males; STin2 genotypes: 10/10 = 12, 10/12 = 28, 12/12 = 23), controlling for gender, pre-intervention smoking severity, and post-intervention use of

**Fig. 1.** Mean amygdala response to tailored smoking-cessation messages mediates the impact of STin2 genotype on smoking cessation. A. Anatomically defined region-of-interest masks for right and left amygdalae shown against MNI 152 template. B. Mediation path diagram. Lines are labeled with unstandardized $B$ coefficients, with standard errors in parentheses. Path a denotes the association of the mean amygdala response with smoking cessation, controlling for the effects of STin2 genotype. Direct effect $c’$ denotes the effect of STin2 genotype on smoking cessation, controlling for the mediation effect. *$p < 0.05$. 
nicotine patches (all smoking-cessation messages: $B = 0.47$, $SE = 0.18$, $t = 2.60$, $p = 0.012$; tailored messages: $B = 0.44$, $SE = 0.17$, $t = 2.59$, $p = 0.012$; untailored messages: $B = 0.49$, $SE = 0.23$, $t = 2.13$, $p = 0.037$) (Supplementary Tables 19–21).

**Amygdala response to smoking-cessation messages mediates STin2 effects on quitting**

Because the mean amygdala response to smoking-cessation messages was both predictive of subsequent quitting outcome and modulated by the STin2 genotype in our data, we next tested for mediation effects. A mediation relationship is illustrated by a three-variable model with three causal pathways: Path $a$ represents the effect of the predictor on the mediator; Path $b$ represents the effect of the mediator on the outcome; and Path $c$ represents the total effect of the predictor on the outcome (Baron and Kenny, 1986). Path $c'$ denotes the direct effect of the predictor on the outcome, controlling for the effect of the mediator. Evidence for mediation is obtained if we can reject the null hypothesis of no difference between the total effect ($c$) and the direct effect ($c'$), that is, $c - c' \neq 0$, demonstrating that the predictor affects the outcome at least in part through the mediator. The benefit of conducting a mediation analysis is that it allows us to test a hypothesis about a relationship between three different variables: thus, a significant mediation effect informs us about the process through which the predictor affects the outcome.

Using the Sobel test of mediation and a bootstrapping approach (Preacher and Hayes, 2008), we found that the mean amygdala response to untailored smoking-cessation messages was associated with a greater amygdala response to untailored smoking-cessation messages, and this greater amygdala response was associated with better odds of quitting success and mediated the impact of the STin2 genotype on quitting.

Similarly, the mean amygdala response to untailored smoking-cessation messages was a significant mediator of STin2 genotype effects on quitting outcome (mean mediation effect value $= 0.35$, $SE = 0.25$, 95% CI: $0.02$–$0.95$, $p < 0.05$) (Path diagram in Fig. 2, see also Supplementary Results for a sample bootstrap output). Controlling for mediation effect in the amygdala reduced the association between the STin2 genotype and smoking cessation (total effect $c$: $B = 0.80$, $SE = 0.38$, $p = 0.037$; direct effect $c'$: $B = 0.61$, $SE = 0.40$, $p = 0.122$). As before, a greater number of the STin2 10 alleles was associated with a greater amygdala response to untailored smoking-cessation messages, and this enhanced amygdala response was associated with better odds of successfully quitting and mediated the impact of the STin2 genotype on quitting.

**Voxel-wise results: brain regions modulated by STin2 genotype**

We also performed an exploratory, voxel-wise search for brain regions whose response to untailored and tailored smoking-cessation messages was modulated by the STin2 genotype, again controlling for the effects of race, gender, pre-intervention smoking severity, and post-intervention use of nicotine patches ($p < 0.005$, 10 contiguous voxels, uncorrected). A right amygdala cluster extending to parahippocampal gyrus was a global maximum in both searches, showing an increase in response magnitude with an increased number of the low-transcription STin2 10 alleles for both tailored messages (xyz coordinates of peak voxel: 33, 0, –12; cluster extent $k = 16$ voxels; $T = 4.04$, $Z = 3.83$; Fig. 3A) and untailored messages (xyz coordinates of peak voxel: 33, 0, –12; cluster extent $k = 94$ voxels; $T = 4.27$, $Z = 4.03$; Fig. 3B), confirming the results obtained with an anatomically defined mask described above.

Apart from the right amygdala/parahippocampal cluster, the number of STin2 10 alleles was also positively associated with response magnitude to tailored messages in the bilateral superior frontal gyrus, right fusiform gyrus, and right uncus (Supplementary Table

[Fig. 2. Mean amygdala response to untailored smoking-cessation messages mediates the impact of STin2 genotype on smoking cessation. A. Anatomically defined region-of-interest masks for right and left amygdala shown against MNI 152 template. B. Mediation path diagram. Lines are labeled with unstandardized $B$ coefficients, with standard errors in parentheses. Path $b$ denotes the association of the mean amygdala response with smoking cessation, controlling for the effects of STin2 genotype. Direct effect $c'$ denotes the effect of STin2 genotype on smoking cessation, controlling for the mediation effect. $^*p \leq 0.05$.]
Our results suggest that smokers who show a greater amygdala response to smoking-cessation messages relative to neutral messages have less difficulty quitting. However, prior neuroimaging evidence also suggests that smokers who show a greater amygdala response to smoking-related cues relative to neutral cues are at a greater risk of slipping during their quit attempt (Janes et al., 2010). We propose that this apparent discrepancy can be explained as follows. The impact of amygdala response to emotionally salient stimuli on learning and behavior depends on the relation of these salient stimuli to current goals. If the emotionally salient stimuli are associated with behavioral responses consistent with the current goal, then amygdala activation to these stimuli should both enhance this goal representation and motivate (i.e., facilitate) the behaviors directed at attaining the goal. Thus, if a smoker has made a resolution to quit smoking, a greater amygdala response to smoking-cessation messages should increase his or her odds of quitting success. But if the emotionally salient stimuli are associated with behaviors that are contrary to the current goal, then we would expect that a greater amygdala activation to these salient stimuli may interfere with the attainment of the goal. For example, smoking-related cues are known to produce a subjective craving and trigger an urge to smoke in smokers (Ferguson and Shiffman, 2009; Rose, 2006), and amygdala activation to smoking-related cues as well as other drug-related cues is positively associated with such craving (Childress et al., 1999; Franklin et al., 2007; Kilts et al., 2001). Thus, a greater amygdala response to smoking-related cues should interfere with the goal to remain abstinent in smokers trying to quit.

Indeed, whereas exposure to drug-related cues is a key environmental trigger and a reliable predictor of relapse, including in smokers (Ferguson and Shiffman, 2009), the impact of this exposure is significantly moderated by individual differences in cue reactivity (Niaura et al., 1998). Our data and previous reports (Chua et al., 2011; Falk et al., 2010) suggest that individual differences in reactivity to smoking-cessation messages may similarly moderate the impact of such messages on quitting or reduction in smoking. Future studies could help elucidate the role of the amygdala by directly comparing the effectiveness of amygdala response to smoking-related cues vs. to smoking-cessation messages as a predictor of subsequent quitting outcome in smokers attempting to quit.

Of note, extensive evidence from both human and animal studies documents the role of the amygdala in detecting and responding to negative emotional stimuli and stressors (LeDoux, 2000; Phelps, 2006). And it is possible that it is specifically amygdala reactivity to negative emotional stimuli and stressors that determines the impact of smoking-cessation messages on subsequent quitting outcome in smokers. We certainly cannot rule out this possibility. However, because growing evidence also supports the role of the amygdala in detecting and responding to positive emotional stimuli (Baxter and Murray, 2002), and because we do not know if participants in our study perceived the smoking-cessation messages as negative or positive, we offer a more general interpretation focusing on amygdala reactivity to all emotionally salient stimuli, irrespective of perceived emotional valence.

Previous neuroimaging evidence from our group and others focused on the prefrontal cortical regions, and showed that the magnitude of response to smoking-cessation messages in the medial prefrontal cortex (MPFC) was a positive predictor of subsequent smoking cessation (Chua et al., 2011) or reduction in smoking (Falk et al., 2010). In the current report, we extend this evidence to the amygdala, a key subcortical region. Because the amygdala and the MPFC share bidirectional anatomical connections (Ghashghaei et al., 2007), and are thought to form a neural circuit of critical importance in addiction (Verdejo-Garcia and Bechara, 2009), future studies could examine whether variability in functional and structural connectivity in the amygdala-MPFC circuit also predicts subsequent quitting, above and beyond the responses of the two regions.

Discussion

The goal of the current study was twofold: first, to examine the role of amygdala response to smoking-cessation messages in quitting; and second, to examine the genetic sources of variability in amygdala response. With respect to the first goal, using a “brain-as-predictor” approach, we demonstrate that neural response to smoking-cessation messages in bilateral, anatomically defined amygdala is a significant predictor of subsequent quitting outcome following a tailored smoking-cessation intervention. Smokers who showed a greater amygdala response to smoking-cessation messages had better odds of quitting success. Importantly, these results were obtained controlling for other factors that are known to affect quitting outcome, including gender, pre-quit smoking severity, and post-intervention use of nicotine patches.

The amygdala response to both tailored and untailored smoking-cessation messages significantly predicted subsequent quitting success. Tailored messages were tailored to individual participants based on their baseline assessment and thus varied between participants, while untailored messages were relevant to all smokers and were identical for all participants. In our view, amygdala engagement reflects that both types of messages were interpreted as emotionally salient by smokers. But because the content of both tailored and untailored messages was related to smoking and was also self-relevant to some degree, we were unable to dissociate the impact of these two attributes on the observed effects. Future studies treating smoking-relatedness and self-relevance as orthogonal factors may be needed to accomplish that, although this may prove difficult in the context of smoking cessation (e.g., it is possible that smoking-related stimuli are interpreted as self-relevant by smokers).

Fig. 3. The right amygdala is a global maximum in a voxel-wise search for brain regions whose response to tailored smoking-cessation messages (A) and untailored smoking-cessation messages (B) is modulated by the STin2 genotype (thresholded at $p=0.005$, 10 contiguous voxels). The results are shown against the MNI template brain. The scale represents $t$ values.
The second goal of the current study was to investigate the genetic sources of variation in amygdala response to smoking-cessation messages in the context of quitting. Here, we employed an imaging genetics approach to examine whether genetic variation in the serotonin transporter (5-HTT) could account for some of the observed individual differences in amygdala reactivity to smoking-cessation messages in smokers. Previous imaging genetics studies (Dannlowski et al., 2010; Hariri et al., 2002, 2005) showed that the low-transcription S and Lc alleles of the promoter polymorphism in that gene (5-HTTLPR/rs25351) were associated with an increased amygdala reactivity to emotionally salient stimuli, such as signals of threat. However, the impact of 5-HTT gene variation on amygdala response to smoking-cessation messages in smokers has not been examined.

We tested both the 5-HTTLPR/rs25351 and the STin2 polymorphisms, and found that the STin2 genotype was a significant independent predictor of the anatomically defined amygdala response to smoking-cessation messages (controlling for 5-HTTLPR/rs25351). The direction of the modulation was broadly consistent with previous studies: namely, the low-transcription allele was associated with an increase in amygdala response relative to the high-transcription allele. However, we found significant effects only for the STin2 genotype, whereas the 5-HTTLPR/rs25351 genotype (controlling for STin2) did not have a significant effect on amygdala response to smoking-cessation messages in our data.

In consideration of our sample size, we limited the genetic analyses to two common, functional polymorphisms in the 5-HTT gene. However, other polymorphisms in the 5-HTT gene have also been described and may contribute to the observed modulation of amygdala response. It is also possible that one or more yet unmeasured polymorphisms in the 5-HTT gene are responsible for the observed effects. A simultaneous examination of the effect of all of these polymorphisms, including their possible interactions, would require sample sizes beyond the scope of a neuroimaging study. Other, alternative approaches may have to be used to systematically examine the impact of 5-HTT gene variation on brain and behavior, including on amygdala response to smoking-cessation messages and smoking-related cues in the context of smoking cessation. For example, future studies could assess the impact of all functional variation in the 5-HTT gene using expression assays as a combined measure of all functional polymorphisms acting together to affect gene transcription. Alternatively, the 5-HTT gene could be sequenced and all observed variants could be coded into one aggregate measure of predicted transcription efficiency. Such aggregate measures could then be integrated with neuroimaging and outcome data.

To conclude, in this report we demonstrate that individual differences in amygdala response to smoking-cessation messages have an impact on subsequent real-life smoking cessation in smokers, and that genetic variation in the central serotonin system accounts for some of these individual differences in amygdala function. Taken together, these results point to the gene–brain–behavior pathway relevant to smoking cessation, and add to our understanding of the role of the amygdala in nicotine addiction. In addition, knowledge of such pathway could impact two important domains of translational research and clinical practice on smoking cessation: prediction and intervention. First, identification of brain predictors of smoking cessation and of genetic variants that modulate these brain processes could enable better predictive models and aid in treatment selection. And second, this knowledge could lead to the development of more effective smoking-cessation interventions tailored to the genetic and neural–response profiles of individual smokers for optimal intervention efficacy.

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**Appendix A. Supplementary data**

Supplementary data to this article can be found online at doi:10.1016/j.neuroimage.2011.12.064.

**References**


