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Effect of Overnight Smoking Abstinence on a Marker for Microglial Activation: A [¹¹C]DAA1106 Positron Emission Tomography Study

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Abstract

Rationale—Microglia are the main immune cells in the central nervous system and participate in neuroinflammation. When activated, microglia express increased levels of the translocator protein 18 kDa (TSPO), thereby making TSPO availability a marker for neuroinflammation. Using positron emission tomography (PET) scanning, our group recently demonstrated that smokers in the satiated state had 16.8% less binding of the radiotracer [¹¹C]DAA1106 (a radioligand for TSPO) in the brain than non-smokers.

Objectives—We sought to determine the effect of overnight smoking abstinence on [¹¹C]DAA1106 binding in the brain.

Methods—Forty participants (22 smokers and 18 non-smokers) completed the study (at one of two sites) and had usable data, which included images from a dynamic [¹¹C]DAA1106 PET scanning session (with smokers having been abstinent for 17.9±2.3 h) and a blood sample for TSPO genotyping. Whole brain standardized uptake values (SUVs) were determined, and analysis of variance was performed, with group (overnight abstinent smoker vs. non-smoker), site, and TSPO genotype as factors, thereby controlling for site and genotype.

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Results—Overnight abstinent smokers had lower whole brain SUVs (by 15.5% and 17.0% for the two study sites) than non-smokers (ANCOVA, $P=0.004$). The groups did not significantly differ in injected radiotracer dose or body weight, which were used to calculate SUV.

Conclusions—These results in overnight abstinent smokers are similar to those in satiated smokers, indicating that chronic cigarette smoking leads to global impairment of microglial activation which persists into early abstinence. Other explanations for study results, such as smoking leading to reduced numbers of microglia or smokers having more rapid metabolism of the radiotracer than non-smokers, are also possible.

Keywords

Tobacco Dependence; Cigarette Withdrawal; Positron Emission Tomography; Magnetic Resonance Imaging; [^{11}C]DAA1106; Microglial Activation; Neuroinflammation

Introduction

Microglia are the main innate immune cells in the central nervous system (CNS), (Waisman et al. 2015). Under homeostatic conditions, they continuously monitor the surrounding environment for signs of infection or homeostasis-perturbing events (Gonzalez et al. 2014; Song et al. 2018). Microglia react to counteract such perturbations in order to protect neurons, which have a limited capacity to regenerate, leading to elevated levels of activated microglia in neurodegenerative diseases (Low and Ginhoux 2018; Schettters et al. 2017). In this context, activated microglia participate in various functions, such as clearance of apoptotic cells and extracellular pathogens, removal of degenerating neurons and extracellular proteins, and cytokine/chemokine production (Anthony and Pitossi 2012). When activated, microglial cellular morphology changes and expression of the translocator protein (TSPO) 18 kDa is increased, thereby making expression of TSPO a marker for neuroinflammation.

Epidemiological studies have shown that cigarette smokers have a lower risk of neurodegenerative diseases than the general population (Foucault-Fruchard and Antier 2017; Quik 2004), and much research has examined the mechanism by which smoking could diminish neuroinflammation, which contributes to neurodegenerative damage. Extensive pre-clinical work has shown that nicotine and/or other agonists at nicotinic acetylcholine receptors (nAChRs) protect against neuronal cell damage (Quik et al. 2015) via binding to α_7 nAChRs and inhibition of microglial activation (Foucault-Fruchard and Antier 2017; Li et al. 2016; Noda and Kobayashi 2017; Shytle et al. 2004).

The radioligand N-(2,5-dimethoxybenzyl)-N-(5-fluoro-2-phenoxyphenyl) acetamide labeled with carbon-11 (abbreviated as [^{11}C]DAA1106) has emerged as a reliable second generation radiotracer for labeling TSPO (Maeda et al. 2004; Okubo et al. 2004; Zhang et al. 2003) with high affinity (Chaki et al. 1999; Chauveau et al. 2008; Venneti et al. 2007; Venneti et al. 2008) for positron emission tomography (PET) scanning *in vivo*. Specific binding of DAA1106 is reported to be greater than previously-used ligands (e.g., PK11195) but less than at least one recently-developed ligand (PBR28) (Owen et al. 2011). Specific binding of DAA1106 correlates with the presence of activated microglia identified by

immunohistochemistry *in situ* (Venneti et al. 2008) and immunohistochemistry combined with autoradiography in brain tissue (Venneti et al. 2007). [¹¹C]DAA1106 was chosen for use here because of these favorable properties and previous experience by our group with this radiotracer (Brody et al. 2017; Brody et al. 2014).

Our group recently used [¹¹C]DAA1106 with PET scanning to compare smokers who had smoked to satiety (~15 min prior to scanning) with non-smoking controls (Brody et al. 2017). The groups differed in whole brain standardized uptake values (SUVs) for the radiotracer, with smokers having 16.8% lower values than non-smokers (and lower mean SUVs in menthol than non-menthol cigarette smokers). Smokers also had lower SUVs in a range of smaller brain volumes of interest (14.6–19.7%). In addition, whole-brain SUV was negatively correlated with participant-reported cigarettes per day. These study findings were consistent with much prior research demonstrating that smokers have less inflammatory functioning than non-smokers.

For the study presented here, we sought to determine if the reduction in [¹¹C]DAA1106 SUV (the marker for neuroinflammation) found in satiated smokers was still present in early (overnight) smoking abstinence. In order to study smokers in a state with potentially significant differences from non-smoking controls, overnight (>12 h) abstinence from smoking was chosen as the time point of interest because nicotine would be expected to be recently eliminated from the body at that time (plasma half-life of ~2 h (Benowitz et al. 2009)) and withdrawal symptoms (e.g., urge to smoke and anxiety/irritability) would be expected to be elevated (Schuh and Stitzer 1995; Ward et al. 2001). As in our prior study (Brody et al. 2017), we hypothesized that smoker versus non-smoker effects would occur globally, based on prior research demonstrating that cigarette smoke is rapidly absorbed (Hukkanen et al. 2005) and results in saturation (or near saturation) of nicotinic acetylcholine receptors throughout the brain (Brody et al. 2009a; Brody et al. 2011; Brody et al. 2006a; Esterlis et al. 2010). We also sought to examine the effect of menthol, because menthol smoking is common (~1/3 of U.S. smokers) (SAMHSA 2009), menthol smokers have more trouble quitting in standard treatment programs than non-menthol cigarette smokers (Gandhi et al. 2009; Okuyemi et al. 2007; Pletcher et al. 2006), menthol smoking has been found to lead to elevated serum nicotine/cotinine and exhaled carbon monoxide levels (Williams et al. 2007) (though not all studies agree on this point (Heck 2009; Werley et al. 2007)), and menthol smokers had lower mean [¹¹C]DAA1106 SUVs than non-menthol smokers in our prior study of smokers in satiety (Brody et al. 2017).

Method

Forty participants (22 cigarette smokers and 18 non-smokers) completed the study and had usable data. These participants underwent telephone and in-person screening, overnight smoking abstinence prior to the PET session (for the smoker group), a bolus [¹¹C]DAA1106 PET scanning session, a blood draw to determine TSPO affinity genotype, and a structural magnetic resonance imaging (MRI) scan, as described below. Four additional participants were enrolled, but were excluded because they were homozygous for the low affinity TSPO genotype (Figure 1). All participants provided written informed consent on forms approved by the Institutional Review Boards of either the VA San Diego Healthcare System or the VA

Greater Los Angeles Healthcare System. A subset of the non-smoker group in the analysis presented here consisted of participants who were included in our previous report (n=13) (Brody et al. 2017).

Participants were recruited through posted flyers and the Internet (e.g., Craigslist). Inclusion criteria were: healthy adult (18 to 65 years old) daily cigarette smokers (range 2–25 cigarettes per day and evidence of tobacco dependence on the Fagerström Test for Nicotine Dependence (FTND)) or non-smokers (< 100 cigarettes lifetime and none within the past year), smoking primarily (> 80%) either menthol or non-menthol cigarettes (for the smoker group), ability to give voluntary informed consent, and an exhaled CO or < 8 ppm (and urine cotinine or < 200 ng/mL) during the study screening visit to support smoking or non-smoking status, respectively. Exclusion criteria were: any psychiatric diagnosis (including mood, anxiety, psychotic, and substance abuse disorders other than Tobacco Use Disorder) within the past six months; any current medication or history of a medical condition that might affect the central nervous system at the time of scanning (e.g., current treatment with a psychotropic medication, or history of severe head trauma with loss of consciousness, epilepsy, or other neurological diseases); regular use (> 1x/week) of anti-inflammatory medication, such as steroidal or non-steroidal anti-inflammatory medications (e.g., corticosteroids, ibuprofen, naproxen, aspirin, or celecoxib [Celebrex[®]]); unstable cardiovascular disease, severe liver disease, or renal insufficiency, which might make tolerating study procedures difficult; or pregnancy. Occasional drug/alcohol use not meeting criteria for abuse or dependence was not exclusionary, but participants were instructed to abstain from drug/alcohol use for at least 48 h prior to PET scanning.

For the telephone screening, a thorough smoking history, including age of first cigarette, maximum smoking habit, menthol or non-menthol cigarette use, length and dates of abstinence periods, previous treatments used, and current smoking habit, was obtained. A brief medical, psychiatric, and substance use history was also obtained. During a subsequent in-person visit, eligibility criteria were confirmed and general demographics, smoking history, and symptom ratings were obtained with the Fagerström Test for Nicotine Dependence (FTND) (Fagerstrom 1978; Heatherton et al. 1991) (to assess severity of Nicotine Dependence), Spielberger State Trait Anxiety Index (STAI) (Spielberger 1983), and Beck Depression Inventory (BDI) (Beck et al. 1996) (to confirm the absence of potentially confounding psychiatric symptoms) for all participants, and Smoker's Profile Form (Brody et al. 2006a; Brody et al. 2013) and Shiffman-Jarvik Withdrawal Scale (SJWS) (Shiffman and Jarvik 1976) (to measure craving and withdrawal) for participants scanned at UCSD. A brief medical review of systems and chart review were also performed by a study physician, along with an exhaled carbon monoxide (CO) measurement (Micro+ Smokerlyzer Breath CO Monitor; Bedfont Scientific, Ltd, UK), urine cotinine screen (The Accutest[®] NicAlert[™]; Jant Pharmacal Corp., Encino, CA), breathalyzer (AlcoMatePro), urine toxicology screen (Test Country I-Cup Urine Toxicology Kit), and urine pregnancy test (Test Country Cassette Urine Pregnancy Test), in order to determine if participants met inclusion/exclusion criteria.

Participants who met inclusion/exclusion criteria and wished to participate underwent a [¹¹C]DAA1106 PET scanning session one week later at either the University of California at

San Diego (UCSD) Center for Molecular Imaging (n=15; 10 overnight abstinent smokers and 5 non-smokers) or VA Greater Los Angeles Healthcare System (VAGLAHS) PET Center (n=25; 12 overnight abstinent smokers and 13 non-smokers), using a procedure similar to the one developed in previous studies (Ikoma et al. 2007; Takano et al. 2010; Yasuno et al. 2012; Yasuno et al. 2008). Smokers were instructed to abstain from cigarettes and other nicotine-containing products from prior to midnight on the night before PET scanning. The PET session was initiated in the afternoon with participants undergoing a brief clinical interview, breathalyzer, exhaled CO level, and urine toxicology and pregnancy screens, in order to verify continued meeting of inclusion/exclusion criteria (including drug abstinence at the time of scanning). An exhaled CO of ≤ 6 ppm was considered consistent with overnight abstinence. Following these procedures, participants were positioned on the PET scanning bed and a venous line was placed. They then received a bolus injection of 350 (± 53) MBq of [^{11}C]DAA1106, and underwent dynamic PET scanning of the brain for the next 90 min. PET scans were obtained using either an ECAT HR+ PET scanner (CTI PET systems, Knoxville, TN) (UCSD) or a Philips Gemini TruFlight PET Scanner (Koninklijke Philips Electronics N.V., Eindhoven, the Netherlands) (VAGLAHS). [^{11}C]DAA1106 was prepared by an established method (Wang et al. 2012). All scans consisted of eighteen 5-min frames. For the UCSD scanner, a 20-min transmission scan from a Ge-68 rod source was performed for attenuation correction following the dynamic scan. The reconstruction used a manufacturer's OSEM algorithm with 4 iterations and 16 subsets. For the VAGLAHS scanner, CT scanning was performed for attenuation correction following the dynamic scan, and the manufacturer's RAMLA algorithm was used for image reconstruction. Investigational new drug (IND) approvals from the Food and Drug Administration (INDs 133984 [UCSD] and 122041 [VAGLAHS]) were obtained to use the radiotracer [^{11}C]DAA1106 for the studies described here.

A 5-ml blood sample was drawn prior to the initiation of PET scanning for genotyping of each individual's TSPO affinity subtype (high [C/C], medium [C/T], or low [T/T]), because these affinity subtypes affect radiotracer binding of all currently used radiotracers determining TSPO availability (Owen et al. 2011; Owen et al. 2012; Yoder et al. 2013). Genomic DNA was extracted from whole blood using QiaAmp DNA Blood Mini Kits (Qiagen, Valencia, CA) (by E.N. and L.S.) and TSPO single-nucleotide polymorphism (rs6971) genotyping using the TaqMan Allelic Discrimination (Thermo Fisher Scientific, Canoga Park, CA) platform was performed in duplicate, according to the manufacturer's specified protocol. Quality control was ensured by perfect concordance of replicate samples, expected minor allele frequencies, and adherence to Hardy-Weinberg equilibrium. Only scans from participants with the high- or medium- affinity genotypes, present in $>90\%$ of North Americans (Mizrahi et al. 2012), were included in study analyses, in order to avoid a potential confound. The exclusion of homozygous low-affinity binders from data analysis is standard practice in recent research in this field (Hafizi et al. 2016; Hannestad et al. 2013; Koshimori et al. 2015; Zurcher et al. 2015).

Within 1–2 weeks of PET scanning, an MRI scan of the brain was obtained, in order to facilitate localization of regions on the PET scans. At UCSD, high-resolution T1-weighted 3D MRIs were obtained on a Siemens 3T Skyra scanner (Erlangen, Germany), with the following specifications: TR=2300ms, TE=3ms, 9-degree angle, 2 acquisitions, $160 \times 256 \times$

256 matrix. Participants who received PET scans at VAGLAHS had MRI scans with the following specifications: 3T GE Medical Systems Signa scanner (Milwaukee, WI) three-dimensional Fourier-transform (3DFT) spoiled-gradient-recalled acquisition with TR=30ms, TE=7ms, 30-degree angle, 2 acquisitions, 256 × 192 view matrix.

As in previous research by our group (Brody et al. 2009a; Brody et al. 2002; Brody et al. 2006a; Brody et al. 2009b; Brody et al. 2006b; Brody et al. 2004), MRI/PET co-registration was performed using Statistical Parametric Mapping software (FIL Methods Group, UK), and automated volumes of interest (VOIs) were determined on MRI using FSL tools for structural MRI. These automated VOIs were transferred from each participant's MRI to his/her co-registered PET scan and visually inspected using PMOD (PMOD Technologies Ltd, Zurich, Switzerland). The primary VOI was whole brain (including gray and white matter) for reasons noted above. However, since automated volumes are easily attained and regional differences are possible, VOIs were also determined for the amygdala, caudate, hippocampus, nucleus accumbens, putamen, and thalamus, similar to VOIs obtained in prior research (Takano et al. 2010; Yasuno et al. 2012).

In order to obtain semi-quantitative measurements of radiotracer binding to TSPO in brain, standardized uptake values (SUVs) were calculated using the standard definition of SUV = mean tissue activity concentration (Bq/mL)/(injected dose [Bq]/body weight [g]). Mean tissue activity from 20–40 min post-injection was used, based on time-activity curves from our previous study (Brody et al. 2017) demonstrating stable activity during this time period. SUV was used as the primary outcome measure because it avoids invasive arterial blood sampling.

For the statistical analysis of data, demographic and rating scale variables were compared between groups using Student t tests for continuous variables and Chi Square tests for categorical variables, in order to determine if groups were similar on these variables. For the central study analysis, an analysis of covariance (ANCOVA) was performed, with whole-brain SUV as the measure of interest and both group (overnight abstinent smoker versus non-smoker) and TSPO genotype (heterozygous or homozygous for the high affinity allele) as between-subject factors (as in (Setiawan et al. 2015; Varrone et al. 2015)). Research site (UCSD versus VAGLAHS) was included in the model as a nuisance covariate, in order to control for possible systematic differences in SUVs related to the different PET scanners or other differences in methodology at the two sites (which were somewhat, but not significantly, different). To determine if group differences were due to regional effects, a multivariate ANCOVA (MANCOVA), using the smaller automated VOIs, was performed with the same structure as the preceding ANCOVA, followed by univariate ANCOVAs for the individual VOIs. The use of an overall MANCOVA for determination of main effect of group (non-smoker versus overnight abstinent smoker) controls for multiple comparisons. For the smaller VOIs, means of left and right SUVs were used. For descriptive purposes, percent difference between study groups for both sites was calculated as:

$100 * (SUV_{\text{non-smokers}} - SUV_{\text{smokers}}) / SUV_{\text{non-smokers}}$. Based on prior research reporting greater brain exposure to cigarette smoke in menthol- than non-menthol- cigarette smokers, we also performed an ANCOVA for whole brain SUV with the same structure as the above test, using non-smoker versus menthol- versus non-menthol- cigarette preference as a

between-subject factor, followed by the same analyses for smaller automated VOIs as for the preceding analysis. For the exploration of associations between PET data and smoking-related behavioral symptoms, correlations were determined for the smoker group between whole brain SUV and smoking-related ratings (for smokers with completed rating scales), while controlling for genotype. For exploratory associations between SUVs and demographic variables, correlations or tests for associations with the same structure as in the preceding analyses were performed for age, sex, and race/ethnicity for the whole study sample. Statistical tests were performed using the statistical software program SPSS version 24 (SPSS Inc., Chicago, IL).

Results

The non-smoker and overnight abstinent smoker groups did not differ significantly in age ($P=.24$), sex ($P=.71$), race/ethnicity ($P=.50$), height ($P=.18$), weight ($P=.25$), depression ($P=.16$) or anxiety ($P=.94$) levels, or caffeine ($P=.23$), alcohol ($P=.27$), or marijuana ($P=.29$) use (Table 1). On average, participants were middle-aged, mostly male, and had generally low levels of depression/anxiety and drug/alcohol use. On the scan day, participants had low levels of exhaled CO (1.9 ± 0.8 ppm for the non-smoker group and 3.3 ± 1.4 ppm for smokers), and smokers had a mean of 17.9 ± 2.3 h abstinence at the time of radiotracer injection (due to overnight abstinence and availability of PET scanning for research at our institutions in the mid-to-late afternoon).

PET data analysis comparing overnight abstinent smokers and non-smokers revealed a significant effect of group for whole brain SUVs (ANCOVA, $P=0.004$), due to the overnight abstinent smokers having lower values than non-smokers. Mean whole brain SUVs were 0.87 for non-smokers and 0.73 for overnight abstinent smokers at UCSD and 1.00 for non-smokers and 0.83 for overnight abstinent smokers at VAGLAHS (15.5% lower at UCSD and 17.0% lower at VAGLAHS). Consistent with these global findings, a significant multivariate effect of group was found (MANCOVA, $P=0.002$) for the smaller VOIs, with all VOIs having a significant between-group effect on univariate analysis (range of P values <0.0005 to 0.026), due to overnight-abstinent smokers having lower SUV values than nonsmokers (range 6.8 to 29.5%) in all VOIs studied.

For the three-group comparison (non-smokers versus non-menthol cigarette smokers versus menthol cigarette smokers), the whole brain SUV comparison was significant (ANCOVA, $P=0.001$), due to non-smokers having the highest values, followed by non-menthol cigarette smokers, and then menthol cigarette smokers. In the multivariate analysis of smaller VOIs, a significant effect of group was found (MANCOVA, $P<0.0005$), with all VOIs having a significant between-group effect, due to the range (from high to low) of SUV values from smokers to non-menthol smokers to menthol smokers. In comparing only the non-menthol with the menthol cigarette smokers, the whole brain SUV comparison was significant (ANCOVA, $P=0.02$), and similar results were found for the smaller VOIs (ANOVAs; P 's=0.02 to 0.07), possibly less highly significant than the overall analysis due to the smaller samples used for comparing the non-menthol- with the menthol- cigarette smoker subgroups.

In the exploratory analysis of smoking-related behavioral variables, a significant inverse relationship was found between depth of inhalation of cigarettes (as rated by participants on the Smoker's Profile Form) and whole brain SUV (correlation=-0.74, P=0.02), suggesting that higher levels of cigarette smoke exposure were associated with lower levels of TSPO availability. No significant associations were found for cigarettes per day, FTND, SJWS scores, age, sex, or race/ethnicity.

Discussion

Cigarette smokers who underwent overnight abstinence had less [¹¹C]DAA1106 binding than nonsmokers throughout the brain, indicating less TSPO availability (and less neuroinflammatory function). Furthermore, menthol cigarette smokers had lower levels of [¹¹C]DAA1106 binding than non-menthol cigarette smokers. These findings are remarkably similar to those of our previous study of smokers in the satiated state (Brody et al. 2017), and indicate that chronic cigarette smoking leads to reduced TSPO availability found in satiety and persisting into early (overnight) abstinence. In addition, the global (rather than localized) effect found here is consistent with prior research demonstrating global effects of smoking on other molecules (such as nAChRs) that are located throughout the brain (Brody et al. 2009a; Brody et al. 2011; Brody et al. 2006a; Brody et al. 2013; Cosgrove et al. 2009; Staley et al. 2006).

A straightforward explanation for the central study finding is that chronic cigarette smoking leads to global impairment of microglial activation in early abstinence. This explanation is consistent with much prior research demonstrating that smokers have impaired inflammatory functioning in other parts of the body, an effect that leads to compromised wound healing (Goncalves et al. 2011; Towler 2000) and lasts for weeks (Pluvy et al. 2015; Rinker 2013). It is also consistent with basic science research demonstrating that microglial cells express nAChRs (Shytle et al. 2004) and that pre-treatment with nicotine (or acetylcholine) inhibits microglial activation (De Simone et al. 2005; Park et al. 2007; Shytle et al. 2004). Similarly, in an animal study of autoimmune encephalomyelitis, nicotine administration was found to decrease microglial activation (Gao et al. 2014), although other condensate of cigarette smoke was found to increase it. In addition, the inverse correlation in this study between [¹¹C]DAA1106 binding and participant self-reports of depth of inhalation indicates that severity of impaired microglial activation may be related to the amount of exposure to cigarette smoke (since depth of inhalation is known to affect absorbed constituents of tobacco smoke (Benowitz et al. 2009)).

Other explanations for study findings are possible. For example, cigarette smoking has been shown to reduce numbers of resident microglial cells (Shi et al. 2009), which would presumably lead to less radiotracer binding. Cigarette smokers are also known to have higher metabolism of some medications than non-smokers (Li and Shi 2015), which (if applicable to [¹¹C]DAA1106) would be expected to lessen radiotracer uptake and binding. In addition, cigarette smoking may result in inflammation in parts of the body other than brain (Rom et al. 2013), which could lead to sequestration of the radiotracer and less binding in brain.

In addition to the overall difference between smokers and non-smokers, the menthol cigarette smoker subgroup had less [^{11}C]DAA1106 binding than the non-menthol cigarette smoker subgroup. This finding is consistent with prior research by our group (Brody et al. 2013) showing greater up-regulation of nicotinic acetylcholine receptors throughout almost all brain regions in menthol- than non-menthol- cigarette smokers, along with a similar finding for TSPO availability in our prior study with [^{11}C]DAA1106 PET and smokers in satiety (Brody et al. 2017). Also, research by others demonstrates that menthol cigarette smoking is associated with more severe biological abnormalities in some (Williams et al. 2007), but not all (Abobo et al. 2012; Muscat et al. 2009), studies that have examined this issue. Therefore, as in prior research, the present finding may be due to greater brain exposure to cigarette smoke (leading to greater impairment of microglial activation) in menthol-cigarette smokers, a direct effect of menthol flavoring, or some other mechanism.

The primary limitation of this study is the absence of arterial blood sampling, precluding the determination of total distribution volume (V_t), which is the gold standard outcome measure for this type of research. V_t may control for the potential confounds of between-subject differences in radiotracer metabolism and binding to vascular endothelium and plasma protein (Koshimori et al. 2015; Rizzo et al. 2014; Turkheimer et al. 2015). While V_t is a common outcome measure in PET studies examining TSPO in conditions other than tobacco dependence (Colasanti et al. 2016; Haarman et al. 2016; Narendran et al. 2014), the SUV measure was used because it is less invasive than methods that include arterial blood sampling and arterial sampling was not feasible. Other similar studies have used pseudo-reference regions for PET data analysis (Colasanti et al. 2016; Coughlin et al. 2014; Hamelin et al. 2016; Kreisl et al. 2016; Lyoo et al. 2015; Zurcher et al. 2015) to minimize potential confounds, but this method would not have been appropriate here due to the hypothesized and confirmed effect of smoking throughout the brain. Additional limitations included scanning at two sites with different model scanners (though results were similar at the two sites), a modest sample size, the absence of measurement of brain nAChR levels, and the fact that smokers were scanned in early abstinence, such that we did not determine whether TSPO availability normalizes with prolonged abstinence. Associations between TSPO availability and brain nAChR availability as determined with PET or single photon emission computed tomography imaging (as we and others have done in prior research (Brody et al. 2016; Brody et al. 2013; Cosgrove et al. 2009; Staley et al. 2006)) or evaluation of normalization of TSPO availability with prolonged abstinence could be addressed in future research.

In summary, cigarette smokers in the overnight abstinent state have lower TSPO availability than non-smoking controls, similar to our recent finding in satiated smokers. The effect was global, and greater for menthol- than non-menthol- cigarette smokers. Future research could build upon this and our previous (Brody et al. 2017) initial studies and directly compare TSPO availability in smokers in satiety with those undergoing short-term and prolonged abstinence. This approach could provide direct information about the time course of normalization of available TSPO levels in smokers who maintain abstinence. Additional future studies could focus on the interplay between smoking, neuroinflammation, and the progression of diseases thought to be mediated by neuroinflammation.

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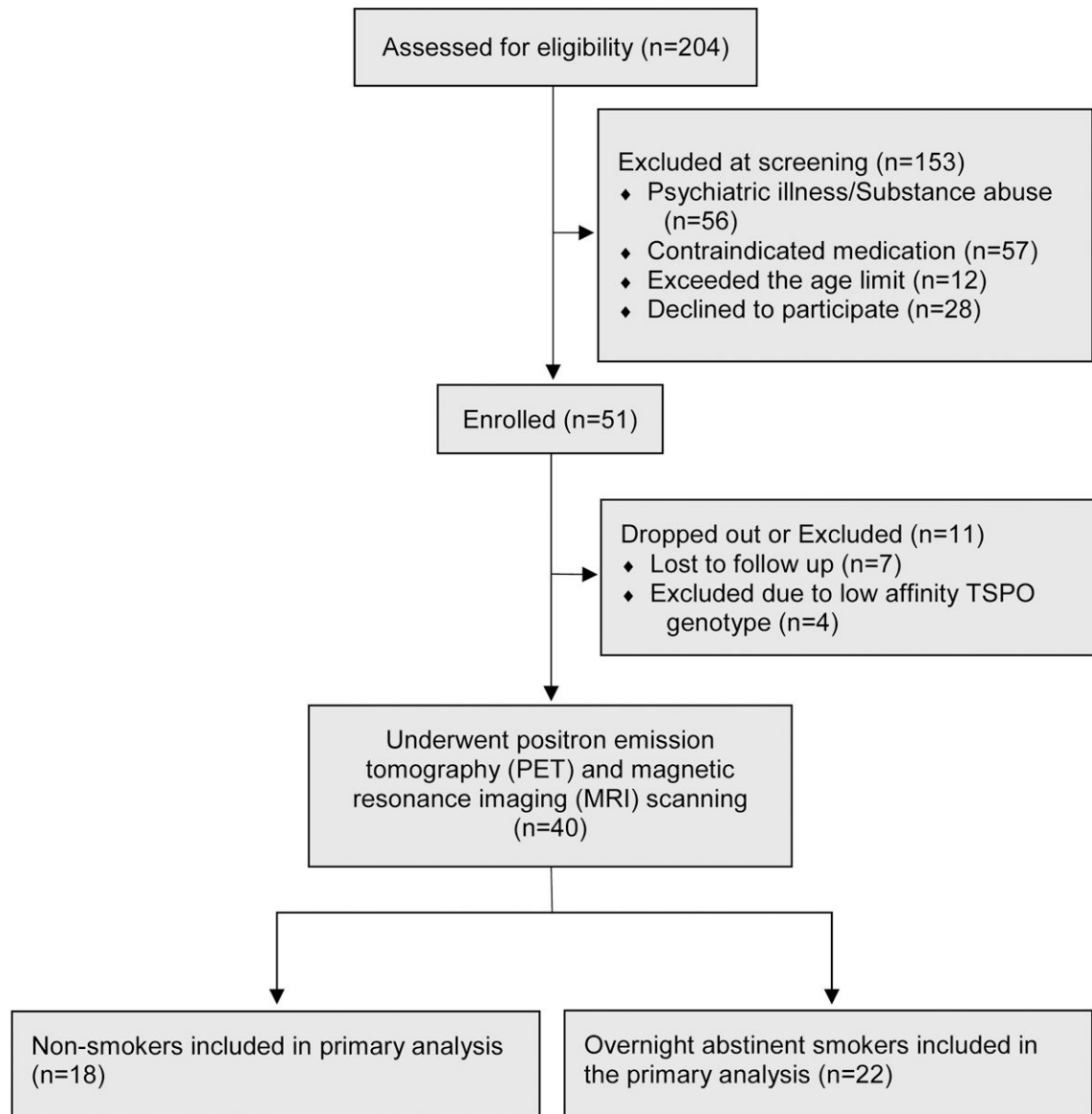


Figure 1: Flow diagram showing the number of potential and actual participants at each step of the study, including reasons for potential participants being screened out of participation.

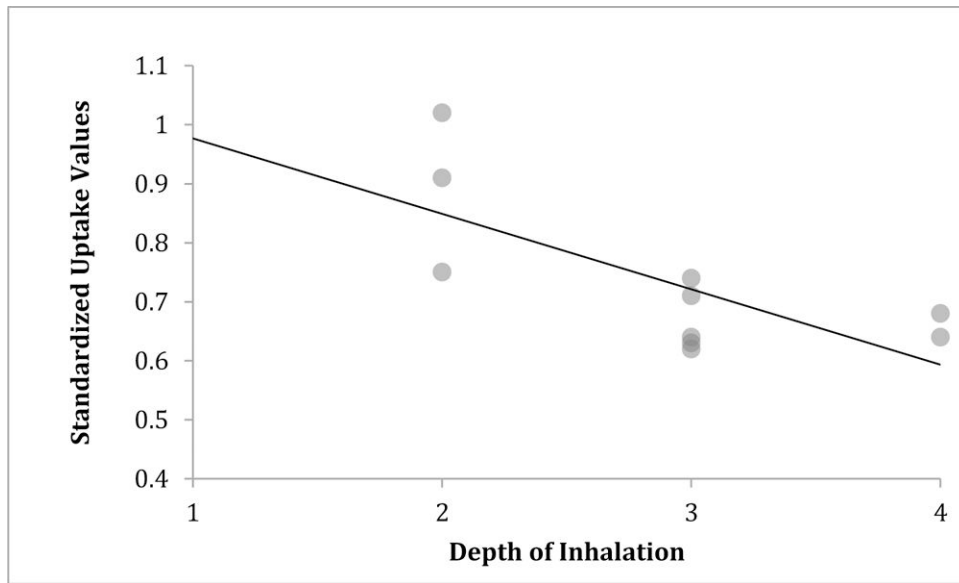


Figure 2: Scatterplot showing inverse correlation (-0.74 , $P=0.02$) between whole brain standardized uptake values and smokers' ratings of depth of inhalation (for study smokers who filled out this questionnaire). The correlation value controlled for translocator protein genotype (high versus medium affinity).

Table 1:

Baseline Demographics and Rating Scale Scores for the Non-Smoker and Overnight Abstinent Smoker Groups

Variable	Non-Smoker Group (n=18)	Whole Smoker Group (n=22)	Non-Menthol Smoker Subgroup (n=16)	Menthol Smoker Subgroup (n=6)
Age	46.9 (\pm 13.4)	41.9 (\pm 13.2)	41.5 (\pm 12.6)	43.0 (\pm 16.7)
Sex (% Female)	22.2	27.3	25.0	33.3
Race/Ethnicity (%)				
African American	22.2	36.4	37.5	33.3
Asian	22.2	9.1	12.5	0
Hispanic	22.2	13.6	12.5	16.7
White	33.3	40.1	37.5	50.0
TSPO genotype (% high affinity)	83.3	77.2	87.5	50.0
Height (Inches)	69.7 (\pm 4.2)	68.0 (\pm 3.4)	68.5 (\pm 3.5)	66.6 (\pm 2.6)
Weight (Kilograms)	89.8 (\pm 23.8)	82.3 (\pm 16.5)	86.8 (\pm 15.8)	70.1 (\pm 12.1)
Cigarettes per day	0 (\pm 0)	11.2 (\pm 6.7)	11.1 (\pm 6.7)	11.7 (\pm 7.3)
Exhaled Carbon Monoxide (ppm)	1.7 (\pm 0.7)	12.7 (\pm 9.0)	11.6 (\pm 8.7)	15.7 (\pm 9.6)
Fagerström Test for Nicotine Dependence (FTND)	0 (\pm 0)	4.3 (\pm 2.9)	4.6 (\pm 3.0)	3.5 (\pm 2.6)
Beck Depression Inventory	1.1 (\pm 1.4)	2.1 (\pm 2.9)	2.3 (\pm 2.8)	1.8 (\pm 3.5)
Trait Anxiety Inventory	30.9 (\pm 9.1)	31.2 (\pm 11.6)	31.4 (\pm 12.0)	30.4 (\pm 11.8)
Caffeine Use (Coffee Cup Equivalents/Day)	0.9 (\pm 1.0)	1.3 (\pm 1.0)	1.2 (\pm 1.0)	1.6 (\pm 1.0)
Alcohol Drinks Per Week	0.2 (\pm 0.5)	1.0 (\pm 1.7)	1.2 (\pm 1.9)	0.6 (\pm 0.8)
Marijuana Cigarettes Per Week	0.1 (\pm 0.5)	0.5 (\pm 1.6)	0.7 (\pm 1.8)	0.2 (\pm 0.6)

All values are presented as means (\pm standard deviation) or percentages. Using χ^2 tests for categorical variables and Student t-tests for continuous variables, no between-group (non-smoker versus smoker) tests were significant, other than differences in measures of smoking (cigarettes per day, exhaled carbon monoxide, and FTND scores; all P values < 0.0005). TSPO = translocator protein.

Table 2:

Ratios of Mean Standardized Uptake Values (SUVs) between Overnight Abstinent Smokers and Non-Smokers for the Two Study Sites

Region	Ratio of Mean SUVs between All Overnight Abstinent Smokers and Non-Smokers by site		Ratio of Mean SUVs between Overnight Abstinent Non-Menthol Smokers and Non-Smokers by site		Ratio of Mean SUVs between Overnight Abstinent Menthol Smokers and Non-Smokers by site	
	UCSD	VAGLAHS	UCSD	VAGLAHS	UCSD	VAGLAHS
Whole Brain	0.84	0.83	0.92	0.85	0.73	0.71
Accumbens	0.73	0.80	0.80	0.83	0.63	0.69
Amygdala	0.87	0.80	0.95	0.82	0.75	0.69
Caudate	0.85	0.85	0.93	0.88	0.73	0.71
Hippocampus	0.89	0.79	0.97	0.81	0.77	0.70
Putamen	0.73	0.78	0.80	0.80	0.63	0.66
Thalamus	0.70	0.79	0.76	0.80	0.60	0.71

Ratios of mean standardized uptake values (SUVs) for whole brain and smaller volumes of interest (VOIs) between overnight abstinent smokers and non-smokers for the two study sites, demonstrating that results from the two sites were consistent with one another. All values were less than 1, indicating lower [^{11}C]DAA1106 binding on positron emission tomography scans for overnight abstinent smokers (and subgroups of smokers based on cigarette type) than non-smokers for all VOIs and both study sites. UCSD = University of California at San Diego. VAGLAHS = VA Greater Los Angeles Healthcare System.