

Original Investigation

Comparison of Urine Cotinine and the Tobacco-Specific Nitrosamine Metabolite 4-(Methylnitrosamino)-1-(3-Pyridyl)-1-Butanol (NNAL) and Their Ratio to Discriminate Active From Passive Smoking

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Abstract

Objectives: Cotinine is the most widely used biomarker to distinguish active versus passive smoking. However, there is an overlap in cotinine levels when comparing light or occasional smokers versus heavily exposed passive smokers. 4-(methylnitrosamino)-1-(3-pyridyl)-1-butanol (NNAL) is a tobacco-specific nitrosamine measurable in urine with a much longer half-life than cotinine. The aim of the study was to determine optimal cutoff points to discriminate active versus passive smokers and to compare sensitivity and specificity for the use of cotinine, NNAL, and the ratio of the NNAL/cotinine in urine.

Methods: Cotinine and NNAL were measured in urine of 373 active smokers and 228 passive smokers.

Results: Geometric mean cotinine levels were 2.03 ng/ml (interquartile interval: 0.43–8.60) and 1,043 ng/ml (658–2,251) and NNAL levels were 5.80 pg/ml (2.28–15.4) and 165 pg/ml (90.8–360) pg/ml in passive and active smokers, respectively. NNAL/cotinine ratio in urine was significantly higher for passive smokers when compared with active smokers (2.85 vs. 0.16, $p < .01$). The receiver operating characteristics analysis determined optimal cutoff points to discriminate passive versus active smokers: 31.5 ng/ml for cotinine (sensitivity: 97.1% and specificity: 93.9%), 47.3 pg/ml for NNAL (87.4% and 96.5%), and 0.74×10^{-3} for NNAL/cotinine ratio (97.3% and 87.3%).

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Conclusions: Both urine cotinine and NNAL are sensitive and specific biomarkers for discriminating the source of tobacco smoke exposure. Cotinine is the best overall discriminator when biomarkers are measured while a person has ongoing exposure to tobacco smoke. NNAL because of its long half-life would be particularly useful when there is a delay between exposure and biomarker measurement. The NNAL/cotinine ratio provides similar sensitivity but poorer specificity at discriminating passive versus active smokers when compared with NNAL alone.

Introduction

Biomarkers are useful in assessing both active and passive smokers' exposure to tobacco smoke. Self-reported exposure such as cigarettes smoked per day by active smokers and hours per day exposed to secondhand smoke (SHS) in passive smokers is imprecise. Exposure patterns differ significantly from person to person due to differences in how cigarettes are smoked, differences in tobacco products, room ventilation, proximity of smokers to nonsmokers, and many other environmental factors (N. L. Benowitz, 1996).

Biomarkers are used to discriminate active from passive smokers in epidemiological studies, population surveillance of tobacco use, and research on the health risks of active and passive smoking. Determination of optimal biomarker cutoff

points to distinguish active versus passive smokers for use in epidemiological studies is important to minimize misclassification (N. L. Benowitz, Bernert, Caraballo, Holiday, & Wang, 2009). Two biomarkers of tobacco smoke exposure are specific to tobacco and can be measured with adequate sensitivity to assess SHS exposure. One is cotinine and the other is 4-(methylnitrosamino)-1-(3-pyridyl)-1-butanol (NNAL).

Measurement of cotinine, the major proximate metabolite of nicotine, in blood, saliva, or urine has supported epidemiological findings of a causal relationship between SHS and lung cancer, cardiovascular disease, and aggravation of chronic obstructive pulmonary disease (COPD) in adults and asthma in children (Boffetta et al., 2006; Strachan, Jarvis, & Feyerabend, 1990; Whincup et al., 2004). Moreover, cotinine is widely used to distinguish smokers from nonsmokers in epidemiology studies and smoking cessation trials. The optimal cutoff point for cotinine to discriminate active smokers from nonsmokers varies from study to study. There are substantial differences between proposed urine cotinine cutoff points in various studies ranging from 20 to 550 ng/ml (Zielinska-Danch, Wardas, Sobczak, & Szoltysek-Boldys, 2007). The most widely cited cotinine cutoff points of 14 ng/ml for serum and 50 ng/ml for urine were provided by Jarvis, Tunstall-Pedoe, Feyerabend, Vesey, and Saloojee (1987). Recently, N. L. Benowitz et al. (2009) proposed optimal serum cotinine cutoff points of 3.08 and 2.99 ng/ml for adults and adolescents, respectively, and estimated an optimal urine cotinine cutoff point of 15 ng/ml. Although plasma and saliva cotinine are considered to be the best biomarkers for both active and passive smoking, the concentration in individuals exposed to SHS is very low. Urine concentrations of cotinine are much higher than in blood or saliva, and in many studies, urine is easier to collect than plasma or saliva.

Tobacco-specific nitrosamines are formed from nicotine and other tobacco alkaloids during tobacco curing and burning. These compounds are potent animal and human carcinogens (Hecht et al., 1999). 4-(methylnitrosamino)-1-(3-pyridyl)-1-butanone (NNK) induces adenocarcinoma of the lung in rodents (Hecht, 1998). NNK is metabolized in human body to NNAL, and this metabolite is further conjugated with glucuronic acid. The level of NNAL in urine of smokers is associated with future risk of lung cancer. NNAL can be measured in urine, whole blood, plasma, and toenails (Bernert et al., 2005; Carmella, Han, Villalta, & Hecht, 2005; Hecht et al., 1999; 2002; Jacob et al., 2008). The half-life of NNAL (10–16 days) is much longer than that of cotinine (16 hr), suggesting that NNAL might be a better measure over tobacco exposure over time and that NNAL might be a better biomarker when sampling cannot be done in temporal proximity to tobacco smoke exposure (such as in hospitalized patients; Goniewicz et al., 2009). We are unaware of any published analysis of the optimal cutoff point to distinguish active versus passive smoking using urine NNAL.

As SHS ages, concentrations of nicotine decline faster than many other gaseous and particulate components of smoke due to absorption of nicotine on surfaces such as walls and carpets (Singer, Hodgson, Guevarra, Hawley, & Nazaroff, 2002). In addition, NNK levels increase as SHS ages, presumably related to the reaction of tobacco-specific alkaloid nicotine with nitric oxide in SHS (Schick & Glantz, 2007; Sleiman et al., 2010). Consistent with these changes in the composition of SHS over time, we recently reported that cotinine measurement leads to an under-

estimation of exposure to NNK from SHS compared with active smoking (N. Benowitz et al., 2010). Thus, we hypothesized that the ratio of NNAL/cotinine would be a better discriminator of active versus passive smoking compared with either NNAL or cotinine alone.

The aims of our study were to determine optimal cutoff points to discriminate active versus passive smokers and to compare sensitivity and specificity for the use of cotinine, NNAL, and the ratio of the NNAL/cotinine in urine.

Materials and Methods

Study Design and Subjects

Subjects from the United States, Poland, and Mexico included 376 smokers and 261 nonsmokers with evidence of exposure to SHS (passive smokers). Characteristics of study groups are presented in Table 1. Subjects were classified as active or passive smokers according to their self-declared smoking pattern or exposure to tobacco smoke. Each subject provided urine for measurement of the concentration of NNAL and cotinine. For the analysis of optimal cutoff points, subjects who did not have an NNAL level higher than the level of quantitation were excluded as we wanted people who definitely had some exposure to SHS. Subjects with a cotinine level below the limit of quantitation were excluded so that we could compute an NNAL/cotinine ratio. After these exclusions, data from 373 smokers and 228 passive smokers were used for cutoff point analysis.

Adult daily smokers were recruited from three different studies (Studies A–C). Study A was a study of tobacco smoke biomarkers comparing Black and White smokers in San Francisco, USA ($N = 128$). Study B assessed the relationship between smoking topography and tobacco biomarkers among daily smokers recruited in Silesia region, Poland ($N = 187$, BK and AS). Study C compared urine biomarkers in daily ($N = 36$) compared with intermittent (nondaily; $N = 22$) smokers in Pittsburgh, USA (Saul Shiffman). Both daily and nondaily smokers were qualified as active smokers.

Nonsmokers exposed to SHS had participated in two previously published studies (Studies D and F) and one unpublished study (Study E) in which urine was collected for the assessment of SHS exposure. The first study (Study D) was a U.S. cohort study of nonsmoking adults with chronic obstructive lung disease (COPD), who collected samples in their homes and mailed samples to the investigators ($N = 72$, MDE; Eisner, Jacob, Benowitz, Balmes, & Blanc, 2009; Eisner et al., 2006). Among these subjects, none were current smokers, but almost half of them had history of smoking. Study E was a Polish cohort study of nonsmoking adults to assess SHS exposure in home and work environments. Subjects provided morning spot samples during screening medical tests ($N = 108$, WZ-D). Study F included volunteers from central Mexico who provided a urine sample before and after visiting a discotheque in which smoking was ongoing for at least 1 hr ($N = 81$, EL-P; Lazcano-Ponce et al., 2007).

Biomarkers Analysis

Cotinine was analyzed by liquid chromatography–atmospheric pressure chemical ionization tandem mass spectrometry (LC-MS/MS) as described by Bernert et al. (2005). Total

Table 1. Characteristics of Study Groups

Study	All subjects					
	Active smokers (<i>n</i> = 376)			Passive smokers (<i>n</i> = 261)		
	A	B	C	D	E	F
Demographic data						
Sample size	130	187	59	72	108	81
Subjects with detectable cotinine levels (%)	129 (99)	187 (100)	58 (98)	54 (75)	108 (100)	80 (99)
Subjects with detectable NNAL levels (%)	128 (98)	187 (100)	59 (100)	48 (67)	106 (98)	77 (95)
Valid subjects	128	187	58	45	106	77
Sex (male), %	74 (58)	83 (44)	26 (45)	19 (42)	50 (47)	27 (35)
Race–ethnicity (White, non-Hispanic), %	67 (52)	187 (100)	33 (57)	42 (93)	106 (100)	NA
Age (mean ± SD)	38.2 ± 10.9	36.3 ± 13.8	NA	64.6 ± 5.8	34.6 ± 16.6	25.7 ± 7.4
Nationality	USA	Poland	USA	USA	Poland	Mexico
Cigarettes/day (mean ± SD)	18.4 ± 8.2	15.0 ± 8.4	6.9 ± 7.1	–	–	–
Subjects included in final analysis						
Analytic data	Active smokers (<i>n</i> = 373)			Passive smokers (<i>n</i> = 228)		
	Cotinine (ng/ml)			Cotinine (ng/ml)		
	Geometric mean (95% CI): 1,043 (919–1,183)			Geometric mean (95% CI): 2.03 (1.59–2.61)		
	IQI: 658–2,251			IQI: 0.43–8.60		
	Range: 4.05–10,788			Range: 0.06–470		
	NNAL (pg/ml)			NNAL (pg/ml)		
	Geometric mean (95% CI): 165 (146–187)			Geometric mean (95% CI): 5.80 (4.90–6.87)		
	IQI: 90.8–360			IQI: 2.28–15.4		
	Range: 3.36–2,319			Range: 0.31–153		
	NNAL/cotinine ratio (×10 ⁻³)			NNAL/cotinine ratio (×10 ⁻³)		
	Geometric mean (95% CI): 0.16 (0.15–0.17)			Geometric mean (95% CI): 2.85 (2.44–3.33)		
	IQI: 0.09–0.26			IQI: 1.32–6.14		
	Range: 0.01–18.2			Range: 0.16–361		

Note. IQI = interquartile interval; NNAL = 4-(methylnitrosamino)-1-(3-pyridyl)-1-butanol.

NNAL (NNAL and NNAL-gluc) was analyzed by LC-MS/MS as described by Jacob et al. (2008). The limits of quantification were 0.05 ng/ml and 0.25 pg/ml for cotinine and NNAL, respectively.

Data Analysis

Nonparametric Kruskal–Wallis analysis of variance was used to compare biomarker levels and their ratio among study groups. Receiver operating characteristics (ROC) curves were generated for three subsets of data (two biomarkers separately and NNAL/cotinine ratio). For generating ROC curves and estimating optimal cutoff points, we assumed 21% population smoking prevalence and that both sensitivity and selectivity had the same importance. To compare diagnostic effectiveness of cotinine, NNAL, and their ratio, we compared their areas under the curve (AUC_{ROC}) using statistical test provided by software system in ROC module. All analyses were performed using STATISTICA data analysis software system v. 8 (StatSoft, Inc., Tulsa, OK).

Results

Characteristics of subjects and the geometric mean data on urine cotinine, NNAL, and NNAL/cotinine ratio are presented in Table 1. As noted previously, after excluding subjects whose cotinine or NNAL were below the limit of quantitation, we had a population of 373 active smokers and 228 passive smokers for ROC analysis.

ROC curves are presented in Supplementary Figure 1, and their detailed characteristics are given in Supplementary Table 1. The ROC analyses showed that optimal cutoff points to discriminate passive versus active smokers were 31.5 ng/ml for cotinine (with a sensitivity of 97.1% and a specificity of 93.9%), 47.3 pg/ml for NNAL (87.4% and 96.5%), and 0.74 × 10⁻³ (NNAL [picograms per milliliter]/cotinine [nanograms per milliliter]) for NNAL/cotinine ratio (97.3% and 87.3%). The frequency distributions of cotinine, NNAL, and NNAL/cotinine ratio among active and passive smokers, with dotted lines showing the optimal ROC cutoff points, are presented in Figures 1–3, respectively.

Results of AUC_{ROC} comparison are presented in Supplementary Table 2. For discriminating active versus passive smokers, there were significant differences between the AUC_{ROC} of cotinine and the AUC_{ROC} of both NNAL and NNAL/cotinine ratio (*p* ≤ .0001), indicating that cotinine had the superior overall discriminating performance. There was no significant difference between AUC_{ROC} of NNAL and NNAL/cotinine ratio (*p* = .26).

Discussion

Our study is novel in determining the optimal urine NNAL cut point to distinguish active versus passive smoking and comparing the performance of urine cotinine and NNAL as discriminators of active versus passive smoking. We defined

Urine Cotinine

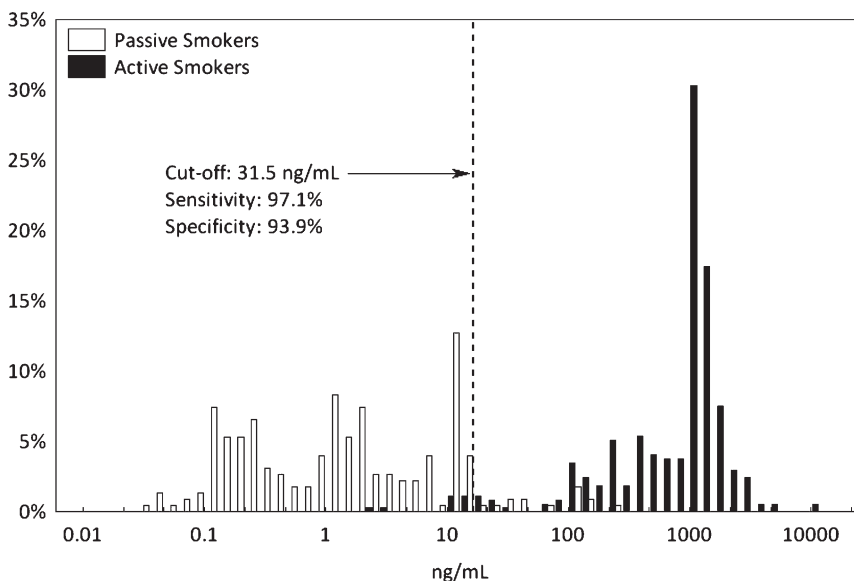


Figure 1. Distribution of urine cotinine concentrations with receiver operating characteristics of the optimal cutoff point among passive and active smokers. The dashed line is drawn at the optimal ROC cutoff point.

passive smoke exposure as having NNAL in the urine at levels above the level of quantitation based on the fact that NNK, the precursor to NNAL, is entirely specific to tobacco.

The utility of urine cotinine to separate passive smokers from active smokers was studied by Zielinska-Danch et al. (2007), who estimated an optimal cutoff point of 550 ng/ml to distinguish the two groups in healthy Polish population. Other studies of Holl, Grabert, Heinze, and Debatin (1998), Hobbs, Wilminck, Adam, and Bradbury (2005), and Savitz, Dole, Terry,

Zhou, and Thorp (2001) have used a similar cutoff point of 500 ng/ml to distinguish active versus combined group of light, nondaily, or passive smokers but did not perform an analysis to specifically determine an optimal cutoff point. The cutpoint of 500 or 550 ng/ml used in these five studies is much higher than the optimal cutoff point of 31.5 ng/ml that we have determined in the present paper.

A urine cotinine cutoff of 15 ng/ml was recently estimated by N. L. Benowitz et al. (2009) based on extrapolation from

Urine NNAL

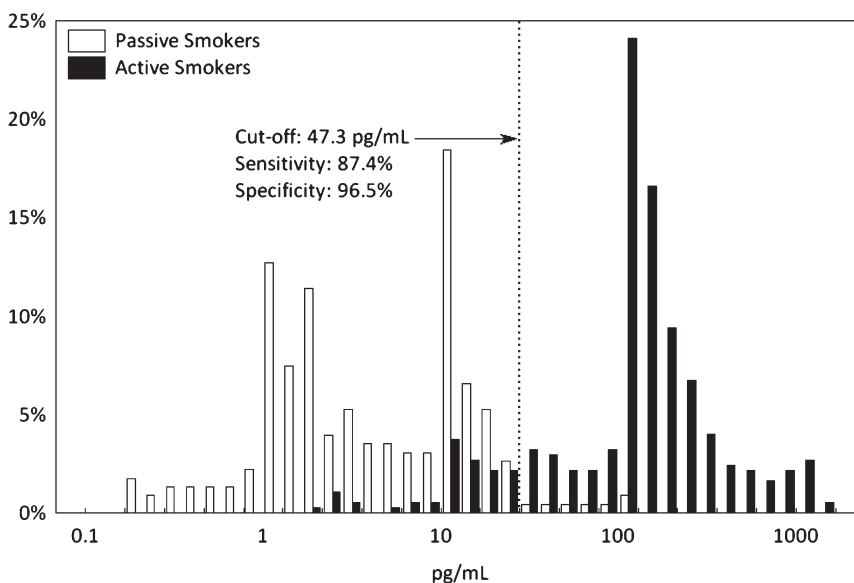


Figure 2. Distribution of urine 4-(methylnitrosamino)-1-(3-pyridyl)-1-butanol (NNAL) concentrations with receiver operating characteristics (ROC) of the optimal cutoff point among passive and active smokers. The dashed line is drawn at the optimal ROC cutoff point.

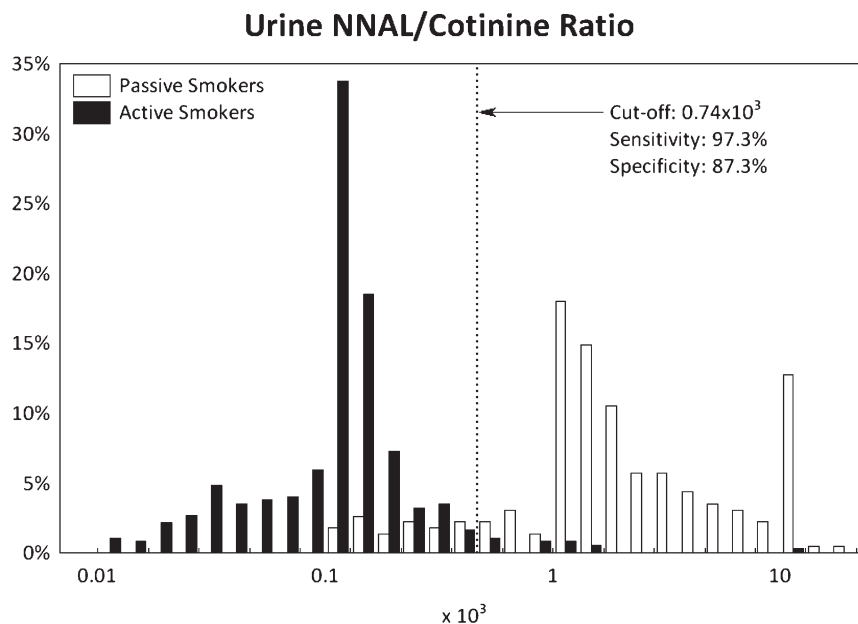


Figure 3. Distribution of the urine 4-(methylnitrosamino)-1-(3-pyridyl)-1-butanol (NNAL)/cotinine ratio with receiver operating characteristics (ROC) of the optimal cutoff point among passive and active smokers. The dashed line is drawn at the optimal ROC cutoff point.

optimal serum levels for distinguishing cigarette smokers and nonsmokers within different racial/ethnic groups in the United States between 1999 and 2004, using data from average ratios of cotinine in urine versus plasma. The differences between the cutoff points for urine cotinine observed in the present study and the *N. L. Benowitz et al. (2009)* study compared with those reported by WZ-D and other in the literature are likely due to different levels of secondhand exposure patterns among examined populations (higher passive exposure results in the higher value of cutoff point) and/or to misclassification of intermittent or light smokers as nonsmokers (such misclassification also leads to higher cutoff points). We believe that our estimate is generalizable to populations of SHS-exposed people broadly since it was estimated based on three subpopulations of active and passive smokers with various patterns of tobacco smoke exposure and smoking behaviors.

Our study presents the first formal analysis of the cutoff point for urine NNAL to distinguish passive versus active smokers. Our optimal cutoff point of 47.3 pg/ml allows discrimination of active versus passive smokers with very high specificity of 96.5% but only moderate sensitivity of 87.4%. To compare the results of our analysis to data from other published studies on active and passive smokers when urine NNAL data were provided, we summarized finding from other studies in Supplementary Table 3. All studies on passive smokers found urine NNAL levels to be lower than our proposed cutoff point value. We identified only one study of active smokers showing that some smokers in Shanghai have lower urine NNAL levels than our cutoff point of 47.3 pg/ml (*Yuan et al., 2009*). The other studies of active smokers showed NNAL much higher than our proposed cutoff point value. It should be noted that the proposed cutoff point for NNAL is based on discriminating known active versus known passive smokers, the latter defined by the presence of NNAL in the urine. If one uses urine NNAL to discriminate active smokers from all nonsmokers (both those ex-

posed and not exposed to SHS, many of whom would have undetectable levels of NNAL in urine), the cutoff point would be lower. We recommend the following two-step approach to using urine NNAL to classify smoke exposure. First, the presence of NNAL at levels above the limit of quantitation indicates tobacco exposure—either active or passive smoke exposure (or smokeless tobacco exposure, which is not considered in the present analysis). Second, considering only individuals with detectable NNAL, the cutoff of 47.3 pg/ml is used to determine if a person is an active or a passive smoker.

Our analysis finds that overall urine cotinine performs better than NNAL or the ratio of NNAL/cotinine in discriminating active versus passive tobacco smoke exposure. Cotinine provides better sensitivity (97.1% vs. 87.4%) but slightly worse specificity (93.9% vs. 96.5%) than NNAL. This finding suggests that when cotinine is used in the epidemiological studies, fewer active smokers are misclassified as passive, whereas using NNAL leads to fewer passive smokers misclassified as active.

However, since NNAL has a long elimination half-life, urine NNAL represents a cumulative measure of exposure over time. The half-life of NNAL averages 10–16 days, so NNAL can be measured in the urine for a month or longer after the last exposure (*Goniewicz et al., 2009*). Cotinine has an average half-life of 16 hr, so cotinine is eliminated from the body within 3–4 days after the last exposure (*N. L. Benowitz, 1996*). Thus, for epidemiological studies of long-term intermittent exposure to SHS or in studies in which biomarker measurements cannot be made in temporal proximity to exposure, such as in hospitalized patients, the use of NNAL is expected to be of greater utility than cotinine.

Recently, we published a paper showing that cotinine measurements lead to an underestimation of exposure to the carcinogen NNK from SHS when compared with active smoking

(N. Benowitz et al., 2010). We showed that passive smoking is associated with a much higher ratio of NNAL/cotinine in the urine compared with active smoking. Thus, in the present analysis, we hypothesized that the ratio of NNAL/cotinine would be a better discriminator than either biomarker alone. This was not the case. The ratio had comparable sensitivity but less specificity compared with cotinine. Overall, the ratio of NNAL/cotinine performed no better than NNAL itself.

In conclusion, we provide novel information on the use of urine biomarkers to discriminate active versus passive smokers. Overall, urine cotinine performs best with a combination of high sensitivity and specificity. However, because cotinine has a short half-life, it is useful only when biomarker samples are taken in close temporal proximity to tobacco smoke exposure. Urine NNAL has high specificity and moderate sensitivity but because of its long half-life has the advantage that it can be used when subjects have not been exposed to tobacco smoke for some period of time, such as in studying tobacco-related disease in hospitalized patients.

Supplementary Material

Supplementary Figure 1 and Tables 1–3 can be found online at <http://www.ntr.oxfordjournals.org>

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Declaration of Interests

Dr. NLB is a consultant to several pharmaceutical companies that market medications to aid smoking cessation and has served as a paid expert witness in litigation against tobacco companies. The other authors have no conflicts to declare.

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