Menthol is the only cigarette additive that is advertised and promoted. It is a flavouring agent that is sprayed on tobacco and offsets the heat sensation during smoking by stimulating cold receptors. Menthol is the chief constituent of peppermint oil and is found in many oral hygiene products, medications, skin lotions, and candy. It is regulated as a food and drug additive but not as a cigarette additive. Menthol increases salivary flow and acts as a bronchodilator. Although mentholated cigarettes were first sold to the public in the 1920s, menthol market share remained low until the 1960s and now accounts for about 25% of all cigarettes sold domestically in the USA. The increase in the popularity of menthol is likely due to advertising that promotes its association with good health. It has become the most commonly smoked brand of cigarette among young African Americans (blacks), but is also smoked by other racial and ethnic groups.13 However the health effects from smoking mentholated cigarettes have not been systematically studied. Smokers of mentholated cigarettes take fewer puffs but expire higher levels of carbon monoxide.6 It has been hypothesised that the higher mortality rate of smoking related cancers among blacks is due to menthol although several studies found little or no risk.10–12

A fundamental question concerning mentholated cigarettes and health is whether the number of cigarettes smoked per day (cpd) and the rate of quitting vary by menthol flavour. There are little data on whether the lower quit rate and cpd in blacks13–15 versus whites is due to their preference for mentholated cigarettes. In one study, the average number of cigarettes smoked per day was 16.2 cpd among smokers of mentholated cigarettes and 17.1 cpd for smokers of non-mentholated cigarettes, although race specific figures were not determined.1 This difference could be due to the higher nicotine content in mentholated cigarettes.1 The preference for high nicotine menthol cigarettes among blacks could reflect their lower amounts of smoking. In active smokers, higher concentrations of blood cotinine have been found in blacks than in whites after controlling for cpd.17–24

Menthol might directly produce its own addicting effect or increase the reinforcing effects of nicotine. Menthol could potentially facilitate addiction by its sensory effects or its perceived qualities as a healthy substance. In one study of 36 smokers, menthol smokers had increased craving relative to non-menthol smokers.25 The possibility that menthol could be a factor associated with the lower quit rate among black smokers remains unexplored. Successful quitters smoke fewer cigarettes per day, at least in some studies,19,26 yet blacks are less likely to quit than whites despite smoking fewer cigarettes per day.

The smoking habits of almost 20 000 white and black smokers were analysed to determine whether the daily consumption of cigarettes and the quit rate were associated with cigarette menthol content.

METHODS
The subjects for this cross sectional analysis were current and former smokers who participated in a case–control study of tobacco related cancers. The original study was conducted to determine the relation between the dose of cigarette exposure and lung, head and neck, kidney, and pancreas cancers. In particular, the risk was studied in relation to cigarette formulation, occupation, social class, and diet.11 For example, it was reported that the histologic specific risk of lung cancer depended on the cigarette “tar” yield.27 Cigarette additives were also hypothesised to affect the risk of aerodigestive cancers, although no association was observed with the use of mentholated cigarettes (compared to non-mentholated cigarettes).28–30

The study was conducted in several hospitals in New York, Washington, DC and Pennsylvania between 1981 and 1999. Newly diagnosed cancer patients were identified from thoracic and other surgery schedules. Non-surgical patients were sought out in oncology wards. Pathology reports were obtained to
confirm the diagnoses and medical reports were reviewed to ensure that the patient had no previous history of lung cancer. Control patients with medical conditions unrelated to cigarette smoking were frequency matched to the cases by age (within five years), sex, race, hospital, and month of interview. Controls were selected randomly from general hospital admitting rosters. Annual refusal rates were always below 15%. All subjects signed a consent form that was approved by the institutional review board.

The current analysis was limited to current and former smokers. Using $\chi^2$ analyses, significant differences were found in the use of mentholated cigarettes by sex, age, smoking status, and cigarettes per day (table 1). Smokers of mentholated cigarettes were significantly more likely to have been women (blacks: 36.6% v 31.3%; whites: 35.1% v 28.8%), less than 55 years of age (blacks: 44.1% v 30.7%; whites: 36.1% v 27.8%), former smokers (blacks), and smoked fewer cigarettes per day (blacks: 18.2 cpd v 20.9; whites: 28.1 v 28.9). There were no differences in the percentage of cases and controls who smoked menthol. For smokers of non-mentholated cigarettes, the mean number of years of smoking was 34.8 for blacks and 32.4 for whites. For smokers of mentholated cigarettes, the mean number of years of smoking was 31.7 for blacks and 33.0 for whites.

Comparing smoking habits between blacks and whites, blacks preferred mentholated cigarettes (34.4% v 13.3%, p < 0.01) and were more likely to have been current smokers (66.4% v 48.3%, p < 0.01). Whites smoked more cigarettes per day than blacks (men: 30.6 cpd v 20.2 cpd, p < 0.01; women: 24.0 cpd v 16.6 cpd, p < 0.01).

The POR of smoking $\geq$ 21 cpd associated with mentholated cigarettes was 0.7 (95% CI 0.5 to 0.9) in blacks who currently smoked and 0.9 (95% CI 0.8 to 1.0) in whites who currently smoked (table 2). Very similar findings were observed in former smokers. Blacks were less likely than whites to smoke more than one pack per day, after adjustment for menthol content, smoking status, and other covariates (POR 0.30, 95% CI 0.27 to 0.34). In an analysis limited to control subjects, the findings were nearly identical (data not shown). There were little differences in the POR by time interval, except for a stronger relation among blacks in the most recent time period. For former smokers, the POR was 0.8 (95% CI 0.4 to 1.2) for the early time period, 0.3 (95% CI 0.5 to 1.2) for the middle period, and 0.3 (95% CI 0.2 to 0.7) for the late time period. For former smokers the POR was 0.8 (95% CI 0.3 to 2.0), 0.9 (95% CI 0.5 to 1.5), and 0.2 (95% CI 0.1 to 0.7), respectively.

### Table 1 Differences in characteristics between 16540 smokers of non-mentholated cigarettes and 3005 smokers of mentholated cigarettes

<table>
<thead>
<tr>
<th></th>
<th>Black</th>
<th>White</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Non-menthol (%) (n=1251)</td>
<td>Menthol (%) (n=655)</td>
</tr>
<tr>
<td>Sex</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>860 (68.8)</td>
<td>415 (63.4)</td>
</tr>
<tr>
<td>Female</td>
<td>391 (31.3)</td>
<td>240 (36.6)</td>
</tr>
<tr>
<td>Age (years)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;45</td>
<td>88 (7.0)</td>
<td>109 (16.6)</td>
</tr>
<tr>
<td>45–54</td>
<td>296 (23.7)</td>
<td>180 (27.5)</td>
</tr>
<tr>
<td>55–64</td>
<td>532 (42.5)</td>
<td>238 (36.3)</td>
</tr>
<tr>
<td>≥65</td>
<td>333 (26.8)</td>
<td>128 (19.5)</td>
</tr>
<tr>
<td>Subjects</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Controls</td>
<td>453 (36.2)</td>
<td>257 (39.2)</td>
</tr>
<tr>
<td>Cases</td>
<td>793 (63.8)</td>
<td>398 (60.8)</td>
</tr>
<tr>
<td>Smoking</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Current</td>
<td>804 (64.3)</td>
<td>461 (70.4)</td>
</tr>
<tr>
<td>Former</td>
<td>447 (35.7)</td>
<td>194 (29.6)</td>
</tr>
<tr>
<td>Cigarettes per day</td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;10</td>
<td>374 (29.9)</td>
<td>226 (34.5)</td>
</tr>
<tr>
<td>11–20</td>
<td>519 (41.5)</td>
<td>294 (44.9)</td>
</tr>
<tr>
<td>21–39</td>
<td>167 (13.4)</td>
<td>77 (11.8)</td>
</tr>
<tr>
<td>≥40</td>
<td>191 (15.3)</td>
<td>58 (8.9)</td>
</tr>
</tbody>
</table>

*p Value* on cigarettes per day missing for 31 white subjects.

NS, not significant.

*Data on cigarettes per day missing for 31 white subjects.
Cigarette mentholation was not associated with continued smoking. The POR was 1.1 (95% CI 0.8 to 1.4) in blacks and 1.1 (95% CI 1.0 to 1.2) in whites (table 3). Blacks were more likely to have been current smokers than whites for both men (POR 1.4, 95% CI 1.2 to 1.7) and women (POR 1.3, 95% CI 1.1 to 1.7) (table 4). The findings from analyses limited to the control subjects were nearly identical to analyses based on cases and controls combined. When stratified by time period, there were few differences in the POR.

The independent predictors of currently smoking and smoking more than one pack per day are shown in table 4. In a model that simultaneously adjusted for sex, race, age, and other covariates, the association with menthol was consistent with previous analyses. Menthol was not associated with continued smoking, and was inversely associated with smoking more than one pack per day.

DISCUSSION
The findings from this study are consistent with another survey showing that smokers of mentholated cigarettes consume fewer cigarettes per day. Our data show that this relation is more pronounced in blacks, which could reflect a higher nicotine content among the menthol brands chosen by blacks than by whites. However, cigarettes with relatively lower nicotine yields, as determined by the Federal Trade Commission (FTC), are smoked more intensely. Consequently, the dose of nicotine delivered is similar among cigarette brands that vary by FTC nicotine content. Blacks smoked fewer cigarettes per day than whites after statistical adjustment for cigarette menthol content and other factors.

Cigarette mentholation was not associated with continued smoking in blacks and in whites. Blacks were less likely than whites to have quit smoking regardless of the brand of cigarette. These findings are similar to newly published data from the Community Intervention Trial for Smoking Cessation (COMMIT). In this cohort of 13 268 smokers, cigarette menthol content was not predictive of quitting in both blacks and whites after five years of follow up. Based on these two studies, it appears that menthol does not modify the addictive properties of cigarettes, although more population based data and physiological studies of menthol and quitting are needed to confirm this. The reason for the lower quit rate in blacks is poorly understood but could be due to lack of perceived benefits, medical advice, and social support. Other factors that have been suggested include targeted cigarette advertising, social stress, greater nicotine dependence, and other psychological reasons.

The limitations of the current study include its cross sectional design and some potential biases. It used a convenience sample and not a random population based sample. Because most subjects were older adults, it is not possible to generalise the findings to younger persons. There might have been a selection bias in the smoking habits among blacks because the participating institutions were large academic medical centres that treat predominantly white populations. In more recent years, blacks tend to seek cancer treatment at municipal hospitals located in minority catchment areas. Furthermore, the study was conducted for almost 20 years and the results might have been influenced by temporal changes in cigarette smoking patterns or in the formulation of mentholated cigarettes. However, when the results were stratified by three time periods, there were little differences in the PORs, with the exception of a higher inverse association between menthol and cigarette amount in blacks in the most recent years of the study. This might simply reflect variability in subgroup analyses, or possibly a trend related to social, demographic or other factors. The percentage of both black and white subjects who smoked mentholated cigarettes is lower than has been reported in national surveys that were conducted during this time period, but this difference could be due to the older ages of the current study subjects or to...
geographic preferences for menthol. The current data are consistent with other reports that showed blacks were about twice as likely to smoke mentholated cigarettes as whites.\(^3\)

It has been reported that the burning of menthol does not produce carcinogens,\(^3\) although one study found that burned menthol produced polycyclic aromatic hydrocarbons.\(^4\)

In experimental animals that were treated with tobacco carcinogens, menthol supplementation in their drinking water did not increase the tumour burden.\(^2\) Despite these findings, it is unknown whether tobacco sprayed with menthol might burn differently from untreated tobacco, or whether the tobacco blend of mentholated cigarettes is different. Smokers of mentholated cigarettes take fewer numbers of puffs per cigarette, but have higher levels of expired carbon monoxide. Although cigarette mentholation was not associated with an increased cancer risk in several studies, there are no studies of mentholated cigarettes and risk of cardiovascular disease.

It is well established that tobacco advertisements of specific menthol brands are targeted to young black consumers, although it remains uncertain if this is a consequence or a preference for menthol among blacks. One study\(^9\) of the content of popular magazines read primarily by blacks found that over an eight year period there were nearly 1500 tobacco advertisements. In contrast, only six of 84 articles about cancer specifically addressed lung cancer. There is a need to better understand the reasons for choosing mentholated cigarettes and how these reasons might be related to quitting success. It is unknown whether black youths would choose different brands or choose not to smoke at all if menthol cigarettes were unavailable. The cooler taste of mentholated cigarette might contribute to a false psychological perception of safety compared to non-mentholated cigarettes.

ACKNOWLEDGEMENTS

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REFERENCES

17. (US Government Printing Office Publication No/S:017/001/005274.)