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# Impairment of Endothelial Function by Little Cigar Secondhand Smoke

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**Objectives:** Little cigars and cigarillos are gaining in popularity as cigarette use wanes, mainly due to relaxed regulatory standards that make them cheaper, easier to buy individually, and available in a variety of flavors not allowed in cigarettes. To address whether they should be regulated as strictly as cigarettes, we investigated whether little cigar secondhand smoke (SHS) decreases vascular endothelial function like that of cigarettes. **Methods:** We exposed rats to SHS from little cigars, cigarettes, or chamber air, for 10 minutes and measured the resulting acute impairment of arterial flow-mediated dilation (FMD). **Results:** SHS from both little cigars and cigarettes impaired FMD. Impairment was greater after exposure to little cigar SHS than by cigarette SHS relative to pre-exposure values, although the post-exposure FMD values were not significantly different from each other. **Conclusions:** Exposure to little cigar SHS leads to impairment of FMD that is at least equal to that resulting from similar levels of cigarette SHS. Our findings support the need to prevent even brief exposure to little cigar SHS, and support tobacco control policies that regulate little cigars as strictly as cigarettes.

**Key words:** little cigars; filtered cigars; cigarillos; secondhand smoke; vascular endothelial function; flow-mediated dilation  
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Little cigars (ie, filtered cigars) and cigarillos have been increasing in popularity for at least 2 decades,<sup>1</sup> counteracting the success of tobacco control efforts on cigarette smoking. As of October 2015, little cigars and cigarillos are not subject to the same product regulations as cigarettes, including warning labels on packages, minimum pack size, and prohibition of marketing using characterizing flavors other than menthol, and they are often taxed at a lower rate.<sup>1,2</sup> Little cigars are also perceived as being less harmful than cigarettes.<sup>3</sup> Filtered little cigars in particular are practically identical to cigarettes in size, shape, and filter style<sup>4,5</sup> (Figure 1). To avoid being regulated as

cigarettes, the small cigar products consist of tobacco wrapped in a tobacco leaf or in paper containing tobacco (cigarettes do not contain tobacco in their wrapper), and cigarillos are heavier than the weight range that defines cigarettes. In addition, the tobacco is of different pH and blend than that in cigarettes.<sup>6,7</sup> Nonetheless, unlike conventional cigars, the smoke from little cigars and cigarillos is often inhaled as in cigarette smoking, and secondhand smoke (SHS) poses hazards to bystanders regardless of smoking technique.<sup>8</sup> Regulating little cigars as strictly as cigarettes would arguably prevent them from simply replacing cigarettes,<sup>9-12</sup> but the relative lack of knowledge about their smoke

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**Figure 1**  
**Cigarettes and Little Cigars Used for this Study**



**Note.**

The UPC code and additional ID numbers on one carton of Swisher Sweets filtered little cigars are shown at right, along with the UPC code on individual packs. Little cigars were 100 mm long and needed to be trimmed to fit in our machine optimized for the ~80 mm standard cigarettes. All photographs by M.L. Springer.

composition and their health effects<sup>1</sup> makes such regulatory expansion difficult to achieve.

Like cigarettes, little cigar smoke contains nicotine and the thousands of chemicals that result from tobacco combustion, and is particularly rich in carbon monoxide, nitrosamines, nitrogen oxides, and ammonia.<sup>5-8</sup> Cigar smoke is associated with elevated risk of oral, lung, and esophageal cancers.<sup>8</sup> The risk of coronary heart disease is 30% higher for cigar smokers than non-smokers, and doubles for those who inhale the smoke.<sup>13</sup> These effects demonstrate the risks associated with long-term use of these products, but the case for regulating them as equivalents of cigarettes would be strengthened by evidence that their immediate health consequences are comparable to those of cigarettes.

One of the most acute health consequences of exposure to cigarette smoke is the immediate impairment of vascular endothelial function, measured as arterial flow-mediated dilation (FMD). FMD is a well-validated marker of cardiovascular risk that is chronically impaired in humans by both active smoking of cigarettes and conventional cigars and by cigarette SHS exposure.<sup>14-16</sup> FMD is temporarily impaired in humans by 30 minutes of exposure to SHS or aged sidestream smoke at real-world SHS levels.<sup>17-19</sup> Sidestream smoke is smoke from the smoldering tip that comprises ~85% of SHS

with the rest being exhaled mainstream smoke.<sup>20</sup> Because the sidestream smoke ages in the exposure chamber prior to exposure, like real SHS does in real exposure scenarios, we refer to it here as SHS. Our micro-ultrasound-based approach to measure FMD in living rats yields results whose pharmacological and biophysical effects are similar to those observed in humans.<sup>21,22</sup> This rat model showed that impairment of FMD occurred with one minute of exposure to cigarette SHS.<sup>23</sup> We report here that brief exposure to little cigar SHS impairs vascular function in rats as least as much as exposure to cigarette SHS.

## METHODS

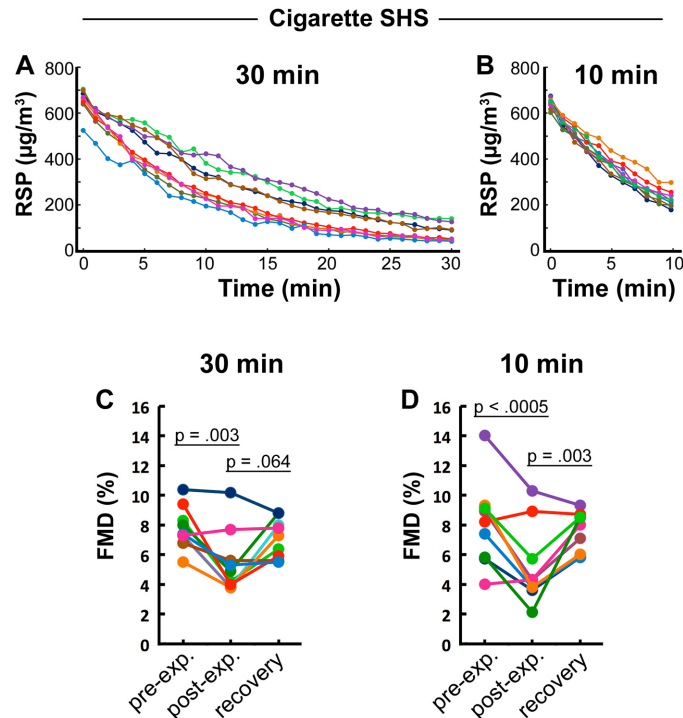
### Animals

We used male Sprague-Dawley rats, 10 weeks old, N = 8 or 9 rats/group. Rats remained anesthetized (ketamine 100 mg/kg, xylazine 5 mg/kg) throughout the experiment and were euthanized immediately afterward. All procedures were approved by the UCSF Institutional Animal Care and Use Committee.

### Measurement of Endothelial Function

Flow-mediated dilation was measured in anesthetized (ketamine 100 mg/kg, xylazine 5 mg/kg) rats

**Figure 2**  
**Comparable Impairment of FMD from 30 and 10 Minutes of Exposure to Declining Levels of Cigarette SHS**



**Note.** Decline of particle concentrations during (a) 30-minute and (b) 10-minute exposure are shown for each rat. (c) FMD response to 30-minute exposure. (d) FMD response to 10 minute exposure. Lines correspond to individual rats; colors track individual animals through smoke and FMD graphs. Thin lines underneath p values denote the data pair being compared.

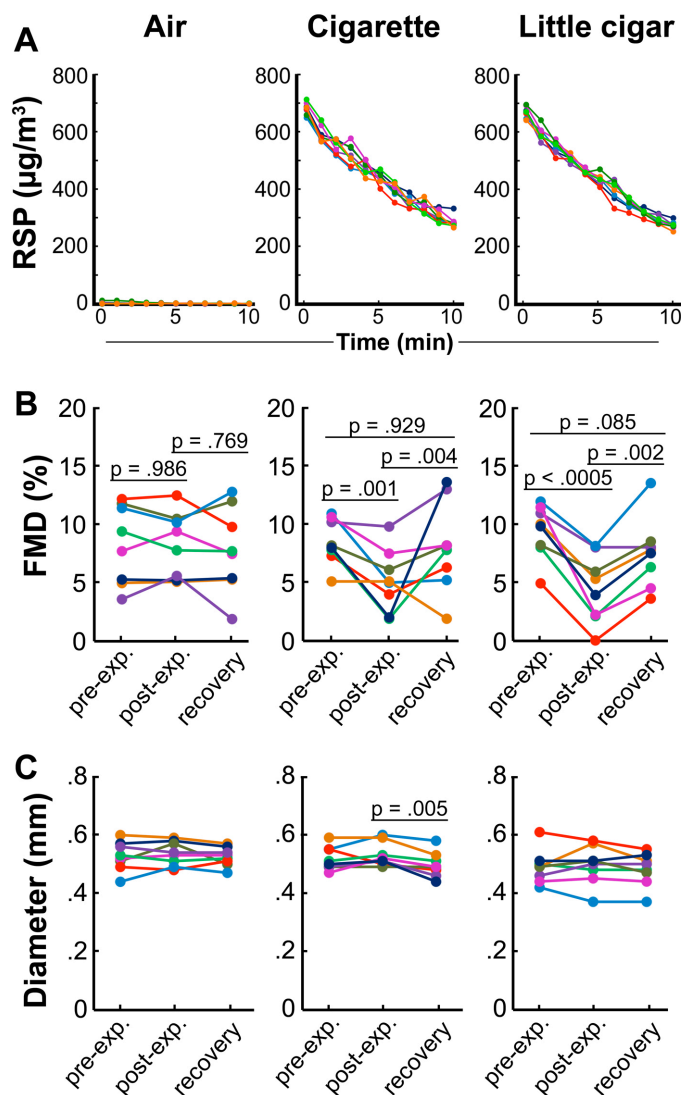
as we have previously described.<sup>22,23</sup> At first, an incision was made to expose the right common iliac artery. Then a suture snare was placed loosely around the common iliac artery to keep the ends of the snare externalized. A baseline ultrasound measurement of femoral artery diameter was taken at diastole with a 35 MHz ultrasound transducer (Vevo660, VisualSonics) system. The artery was occluded for 5 minutes, during which the femoral artery was prevented from moving by a supportive piece of tubing. The snare was released to re-establish perfusion with a rush of blood flow (hyperemia), and ultrasound measurements of femoral artery diameter were performed every 30 seconds for 3 minutes with additional measurements at 4 and 5 minutes. FMD was calculated as % change:  $(\text{peak diameter} - \text{diameter}_{\text{baseline}}) / \text{diameter}_{\text{baseline}} \times 100$ .

### Exposure to Smoke

Little cigars were Swisher Sweets brand (20/pack) and cigarettes were Marlboro Red brand, neither of which was flavored (Figure 1). We were only able to obtain the little cigars at 100 mm in length (slightly longer than the length of the Marlboro cigarettes typically used in our system, ~80 mm), so their non-filter ends were trimmed to enable them to be lit by our system's fixed-position automatic lighter coil. The Swisher Sweets product was 1.22 g/stick, or 2.69 lb per thousand sticks, falling within the US Federal Government's tax definition of a little cigar (maximum of 3 lb per 1000 count).<sup>24</sup>

Our modified cigarette smoking machine<sup>23</sup> uses a single chamber to collect sidestream smoke; the anesthetized rat is exposed by placing its head

**Figure 3**  
**Impairment of FMD from Exposure to Little Cigar SHS**



**Note.**

(a) Particle kinetics and quantitation for comparison of cigarette and little cigar sidestream smoke. Each line represents particle measurements over 10 minutes for each rat; errors are SD. (b) Impairment and recovery of FMD after 10 minutes of exposure to air, cigarette SHS, and little cigar SHS. (c) Lack of effects on baseline arterial diameter (pre-occlusion).  $p > .44$  for all sequential diameter measurements unless otherwise noted. Lines correspond to individual rats; colors track individual animals through panels a, b, and c. Thin lines underneath p values denote the data pair being compared.

through a gasket into the chamber. For each experiment, a cigarette or little cigar was pre-humidified overnight by placement over 16% glycerol in distilled water and then was lit and smoked for 3 minutes under well-established research conditions (ISO Standard 3308:2012, one 35 ml puff of

2 sec duration once per minute) and extinguished. Respirable suspended particles  $<2.5 \mu\text{m}$  (RSP) were measured with a TSI Sidepak AM510 monitor sampling and returning air from the chamber once per minute. The AM510 was factory calibrated and was then specifically calibrated for tobacco



smoke particles in our laboratory by gravimetric sampling of smoke from the exposure chamber, resulting in a calibration factor of 0.3 applied to the raw data. Due to particle adsorption and deposition, particle concentration falls over time (Figures 2 and 3). Particle concentration in the exposure chamber was adjusted by venting until the desired starting concentration was reached. Our target starting concentration was  $600 \mu\text{g}/\text{m}^3$  RSP, representative of smoke levels found in restaurants where smoking is allowed,<sup>25,26</sup> and similar to the conditions of our earlier study of cigarette SHS.<sup>23</sup> The cigarette or little cigar was extinguished, and an individual anesthetized rat, after baseline FMD measurement (denoted as “pre”), was exposed for the specified duration (which was determined in a preliminary experiment; see beginning of Results section) and was then returned to the ultrasound system for post-smoke FMD measurement. Due to technical limitations, the initial post-smoke FMD measurement (“post”) took place roughly 10 minutes after the end of exposure. FMD was measured a third time 30 minutes later to assess recovery (ie, 40 minutes after end of exposure; “recovery”). Exposure to air in the cleaned exposure chamber without tobacco product provided a negative control.

For each experiment, rats from each group were exposed in a random order, and arterial diameter measurements were obtained by an investigator unaware of the experimental condition.

### Statistics

To evaluate differences in FMD or baseline diameter versus times or exposure conditions, we fit a 2-factor (exposure condition and time) repeated measures ANOVA to all data at once using a linear mixed model estimated with restricted maximum likelihood estimation, then tested for differences over time and across exposure conditions using contrasts and pairwise comparisons, adjusted for multiple comparisons using the Šidák method using Stata 13.1. Time was modeled as a repeated effect and the residual covariance structure was independent. Variability in the data are reported as standard error of the mean (SEM) for paired analyses of FMD and diameter results, and standard deviation (SD) for smoke particle concentrations.

## RESULTS

### Exposure to Cigarette SHS for 30 minutes or 10 minutes Impairs FMD to Comparable Extents

To determine an appropriate exposure time for our primary purpose of studying brief exposure to little cigar SHS, we first validated a hypothesis that a 10-minute exposure would be sufficient in the context of cigarette SHS, with which we had extensive experience. We showed previously that one minute of exposure to cigarette SHS impairs FMD significantly but modestly, and that exposure for 30 minutes substantially impairs FMD, but most of the exposure occurs during the first 10 minutes due to the decline in particle concentration over time in our smoke system.<sup>23</sup> Therefore, we directly compared FMD impairment after 30 minutes versus 10 minutes of exposure to cigarette SHS (Figure 2). For 30 minutes: mean starting concentration was  $655 \pm 53.4 \mu\text{g}/\text{m}^3$ , mean concentration over time was  $253 \pm 59.4 \mu\text{g}/\text{m}^3$ , total exposure (area-under-curve) was  $7493 \pm 1804 \mu\text{g}/\text{m}^3 \cdot \text{min}$ ; for 10 minutes: mean starting concentration was  $644 \pm 23.5 \mu\text{g}/\text{m}^3$ , mean concentration over time was  $396 \pm 27.2 \mu\text{g}/\text{m}^3$ , total exposure was  $3921 \pm 279 \mu\text{g}/\text{m}^3 \cdot \text{min}$ .

Flow-mediated dilation declined after a 10-minute exposure from a mean of  $8.0 \pm 1.0\%$  to  $5.2 \pm 0.9\%$  ( $p < .0005$ ) and after a 30-minute exposure declined from  $7.8 \pm 0.5\%$  to  $5.5 \pm 0.7\%$  ( $p = .003$ ). Recovery of FMD after an additional 30 minutes of exposure (to clean air) in each group reached  $7.5 \pm 0.5\%$  and  $7.1 \pm 0.4\%$  in the 10 and 30 min groups ( $p = .003$  and  $p = .064$ , respectively, relative to the initial post-exposure values). There was no significant difference between the initial mean post-exposure impaired FMD values between the 2 exposure time conditions ( $p = .759$ ), nor between the subsequent mean recovery values ( $p = .682$ ). This result indicates that exposure to the low levels of smoke remaining during the last 20 minutes of a 30-minute exposure did not decrease FMD further after the higher level exposure during the first 10 minutes. Therefore, a 10-minute exposure to declining SHS levels was used as the standard exposure time for the subsequent experiment comparing SHS from cigarettes and little cigars.

### Comparable Impairment of FMD by SHS from Cigarettes and Little Cigars

We performed a direct comparison between little

cigar and cigarette SHS, both at starting smoke RSP concentrations of  $\sim 670 \mu\text{g}/\text{m}^3$  declining over 10 minutes, with chamber air as a negative control (Figure 3). For air, mean starting concentration was  $3 \pm 4 \mu\text{g}/\text{m}^3$ , mean concentration over time was  $1.7 \pm 1.3 \mu\text{g}/\text{m}^3$ , total exposure was  $17 \pm 13 \mu\text{g}/\text{m}^3 \cdot \text{min}$ ; for cigarettes, mean starting concentration was  $682 \pm 20 \mu\text{g}/\text{m}^3$ , mean concentration over time was  $449 \pm 14 \mu\text{g}/\text{m}^3$ , total exposure was  $4461 \pm 138 \mu\text{g}/\text{m}^3 \cdot \text{min}$ ; for little cigars, mean starting concentration was  $670 \pm 19 \mu\text{g}/\text{m}^3$ , mean concentration over time was  $441 \pm 10 \mu\text{g}/\text{m}^3$ , total exposure was  $4380 \pm 104 \mu\text{g}/\text{m}^3 \cdot \text{min}$ .

Flow-mediated dilation in the group exposed to little cigar SHS declined from  $9.4 \pm 0.8\%$  before exposure to  $4.4 \pm 1.0\%$  and recovered to  $7.5 \pm 1.1\%$  by 30 minutes later. FMD in the group exposed to cigarette SHS declined from  $8.5 \pm 0.7\%$  before exposure to  $5.2 \pm 0.9\%$  initially after exposure and recovered to  $8.0 \pm 1.4\%$  by 30 minutes later. FMD in the air group did not change significantly. Baseline (pre-occlusion) diameters were not affected by smoke exposure and were comparable for all groups from pre- to post-exposure ( $p > .5$ ), although for cigarettes only, there was a slight but significant ( $p = .005$ ) reduction in diameter between the initial post-exposure and recovery values, for reasons that remain unknown. The difference between absolute values of post-exposure FMD measurements for the cigarette and little cigar groups did not approach significance ( $p = .948$ ). We conclude that exposure to little cigar SHS leads to impairment of FMD that is at least comparable to that resulting from the same level of cigarette SHS.

## DISCUSSION

A limitation is that the extent of acute endothelial functional impairment was based on response to equivalent smoke particle concentrations from the 2 products, whereas the amount of SHS liberated from each product during real-world use may vary. However, when Swisher Sweets little cigars (100 mm) and Marlboro cigarettes (85 mm) were smoked to completion using laboratory smoking conditions standardized for each product, the little cigars produced roughly 150% as much tar, 250% as much CO, and 350% as much nicotine as the cigarettes,<sup>7</sup> indicating that toxicity of little cigar SHS may be even greater relative to that of ciga-

rettes than we have reported here.

We used Marlboro Red cigarettes and Swisher Sweet little cigars because they are representative of commonly-used brands of each product, because we have used Marlboro Red cigarettes in our previous studies,<sup>23</sup> and because of the earlier report mentioned above<sup>7</sup> that compared the smoke composition from these brands. Given the similarity of effects from exposure to SHS from these 2 products, it is unlikely that other cigarette or little cigar brands would be fundamentally different in terms of effects of SHS on vascular endothelial function.

We conclude that differences between tobacco and rolling paper composition in cigarettes and little cigars do not translate into differences in acute endothelial toxicity. Exposure to little cigar SHS leads to impairment of vascular function that is at least comparable to that resulting from similar levels of cigarette SHS. Our findings support the need to prevent even brief exposure to little cigar secondhand smoke.

## IMPLICATIONS FOR TOBACCO REGULATION

Cardiovascular toxicity is a major consequence of active and passive smoking alike,<sup>27</sup> and tobacco use causes over 140,000 cardiovascular deaths annually in the United States.<sup>28</sup> Cardiovascular toxic effects are rapid and result in increased risk of myocardial infarction and stroke.<sup>26,29</sup> Repeated exposure to SHS causes lasting reduction in FMD<sup>14</sup> and exposure to SHS during childhood correlates with lower FMD during adulthood.<sup>30</sup> For these reasons, acute vascular toxicity is a relevant and important consequence by which to evaluate harmful effects of SHS from little cigars. In our direct comparison of vascular response to similar levels of SHS from little cigars and cigarettes, we observed greater decrease in FMD after exposure to little cigar SHS.

Santo-Tomas et al<sup>16</sup> reported that FMD is impaired by active smoking of conventional cigars. Our results extend those findings to passive exposure to SHS from little cigars. The potential public health impact of demonstrating comparable acute harm to vascular function from little cigar and cigarette SHS demonstrates the need not only for regulation but also education about the health effects of little cigars, which are viewed by some smokers as less dangerous than cigarettes.<sup>3</sup> These results

contribute to tobacco regulation objectives both by countering this impression, and by strengthening the scientific basis for regulatory decisions regarding little cigars. Policies that protect people from even brief exposure to little cigar SHS are well-justified. Little cigars should be regulated like cigarettes and, in the absence of specific evidence to the contrary, regulators should apply knowledge of the cardiovascular effects of cigarettes to little cigars.

### Human Subjects Statement

Not applicable.

### Conflict of Interest Statement

The authors have no competing interests to declare.

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