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Secondhand smoke exposure and risk following the Irish smoking ban: an assessment of salivary cotinine concentrations in hotel workers and air nicotine levels in bars

M Mulcahy, D S Evans, S K Hammond, J L Repace, M Byrne

Objective: To investigate whether the Irish smoking ban has had an impact on secondhand smoke (SHS) exposures for hospitality workers.

Design, setting, and participants: Before and after the smoking ban a cohort of workers (n = 35) from a sample of city hotels (n = 15) were tested for saliva cotinine concentrations and completed questionnaires. Additionally, a random sample (n = 20) of city centre bars stratified by size (range 400–5000 square feet), were tested for air nicotine concentrations using passive samplers before and after the ban.

Main outcome measures: Salivary cotinine concentrations (ng/ml), duration of self reported exposures to secondhand smoke, air nicotine (µg/cubic metre).

Results: Cotinine concentrations reduced by 69%, from 1.6 ng/ml to 0.5 ng/ml median (SD 1.29; p < 0.005). Overall 74% of subjects experienced decreases (range 16–99%), with 60% showing a halving of exposure levels at follow up. Self reported exposure to SHS at work showed a significant reduction from a median 30 hours a week to zero (p < 0.001). There was an 83% reduction in air nicotine concentrations from median 35.5 µg/m³ to 5.95 µg/m³ (p < 0.001). At baseline, three bars (16%) were below the 6.8 µg/m³ air nicotine significant risk level for lung cancer alone; at follow up this increased to 10 (53%).

Conclusions: Passive smoking and associated risks were significantly reduced but not totally eliminated. Exposure to SHS is still possible for those working where smoking is still allowed and those working where smoke may migrate from outdoor areas. Further research is required to assess the true extent and magnitude of these exposures.

On the 29 March 2004, Ireland became the first country to introduce a smoking ban in indoor workplaces. Smoking has been banned in all bars, restaurants, cafes, and hotels (excluding bedrooms, outdoor areas, and properly designed smoking shelters). The ban aims to protect workers, particularly those in the hospitality industry, from the health consequences of secondhand smoking (SHS) such as increased risks for lung cancer, coronary heart disease, and asthma. Although several studies have assessed exposure levels relative to workplace smoking policy, relatively few studies have illustrated the levels of exposure of those most highly exposed such as hospitality workers and fewer still have assessed the exposure reductions yielded by smoking bans.

As more jurisdictions consider legislating for smoke-free workplaces, these may become the most sought after elements of the evidence base. The purpose of this research was to investigate whether the Irish smoking ban has had an impact on SHS exposures for hotel and bar workers.

METHODS

Cotinine monitoring of hotel workers

Cotinine is a biomarker of recent passive smoking exposure, as it is produced when the body breaks down nicotine absorbed over the previous 2–3 days. Saliva samples were obtained from a sample of hospitality staff 2–3 weeks before (baseline) and 4–6 weeks after (follow up) the smoking ban. All 40 hotels in Galway city in the Health Service Executives' Environmental Health database were visited. Each manager was asked for permission to approach staff and request their participation in the study. Hotel staff were then recruited; screening questions ensured participants were non-smokers from non-smoking or restricted smoking households (less than 45 minutes exposure per day to one smoker). All participants gave informed consent. The saliva samples were obtained during a working shift using salivettes (Sarstedt Ltd, County Waterford, Ireland; part no. 15/1534) and the time worked before the test recorded. The samples were coded and stored in cool boxes before frozen storage within three hours. Samples were shipped in dry ice to ABS Laboratories, London for analysis by gas chromatography.

Baseline demographic data were obtained with a questionnaire completed after the cotinine test.

Airborne nicotine monitoring of bars

Passive samplers were located behind the bar counters of 20 city centre bars for 7–10 hours, on the Friday preceding the ban (baseline) and six weeks later (follow up). An even number of bars were selected randomly from a database categorised into small, medium, large, and super pubs. Of the 20 bars approached, none refused to participate. Four follow up samples had to be repeated because of lost data. As one venue had closed, 19 of the 20 venues were sampled at follow up.

Abbreviations: CHD, coronary heart disease; LDL-C, low density lipoprotein cholesterol; OHSA, US Occupational Safety and Health Administration; SHS, secondhand smoke.
The baseline profile of subjects is given in Table 1. All samplers were clipped to spirit drink dispensers in the bars before lunchtime on Fridays and collected during the evening. In order to be able to interpret whether recorded levels were associated with high/low smoking rates, and to check for tampering with the samplers between times, a series of observations were made. At the time of placing and retrieving the samplers, the number of occupants and smokers in the bar was recorded. In addition, two further counts were covertly undertaken, also checking the positioning of the sampler for signs of tampering. Samples were sent to the Chemist Service Laboratory, Public Health Agency, Barcelona, Spain, for analysis by gas chromatography/mass spectrometry. Weather conditions at both baseline and follow up were mild/fine with no rain.

RESULTS

Profile

At baseline 52 workers from 15 hotels gave saliva samples and completed a questionnaire. Eight workers (15%) refused to participate in the study. At follow up, 11 participating workers were unavailable to give samples as they no longer worked at the venues and so they were excluded (baseline cotinine values: 0.2, 0.5, 1.3, 2.2, 3.9, 4.1, 5.6, 10.5, 14.1, 31.4, and 105.1 ng/ml). In addition, to control for misreported smoking status, we excluded subjects with saliva concentrations exceeding 15 ng/ml (baseline cotinine values: 21.8, 64.9, 107.9, 161.3, 322.9, and 349.7 ng/ml, respectively). Baseline and follow up comparisons were therefore undertaken on 35 of the 55 workers at 15 of the hospitality venues. The baseline profile of subjects is given in Table 1.

Bar venues ranged in size from 384 square feet to 5200 square feet and represented a cross section of venue types including traditional bars, music bars, hotel bars, and super pubs. At follow up eight bars had provided newly designated outdoor smoking areas. These uncovered areas were adjacent to main entrance doorways and consisted of areas cordoned off by waist level demountable screens.

Cotinine concentration

At baseline, 20% of workers had cotinine concentrations above 5 ng/ml, with 6% having concentrations greater than 10 ng/ml. At follow up, no cotinine concentrations greater than 5 ng/ml were recorded (Fig 1). At baseline 80% exceeded 0.57 ng/ml with this reducing to 49% at follow up. Overall 74% experienced decreases of 16–99%, with over a third of all hotel staff (34%) experiencing reductions greater than 70%. For 60%, exposure levels were halved at follow up. Of the nine that experienced increases, all were less than 2 ng/ml. Workers were tested 6.6 hours into their shift on average at baseline and 4.95 hours into their shift at follow up.

Table 1 shows that median cotinine concentrations declined from 1.6 ng/ml at baseline to 0.5 ng/ml at follow up (p < 0.005: 95% confidence interval (CI)). This represents a 69% reduction in median cotinine concentrations. Median concentrations reduced for all types of work except management, which experienced a small, insignificant increase from already low values (p > 0.05). Reductions were significant for waiters and “other” duties (p < 0.05: 95% CI) undertaken by 31% of workers—namely, front desk (3), portering (2), leisure club (1), accommodation (2), and kitchen (1) bar staff (2) (p < 0.05: 95% CI). The highest median cotinine concentrations at baseline and at follow up were experienced by bar staff (5.4 and 2.6 ng/ml) and waiters (3.9 and 1.6 ng/ml).

A reduction in median cotinine concentrations from 1.6 to 1.4 ng/ml (p < 0.05) was experienced by subjects living in households where no one smoked (n = 23). Those living in restricted smoking households (n = 12) experienced a reduction in median cotinine concentrations from 1.45 ng/ml to 0.4 ng/ml (p < 0.05).

Self-reported exposure to SHS at work showed a significant reduction from a median of 30 hours a week at baseline, to a median of zero at follow up (p < 0.001: 95% CI). At baseline, 68% reported that they had over 21 hours of exposure to SHS with 23% having up to 20 hours of exposure and 10% having no exposure. At follow up, 70% reported no exposure and 30% reported 1–2 hours of exposure at work in the last seven days.

Air nicotine

The median air nicotine concentrations reduced significantly from 35.52 µg/m³ (mean = 35.81 µg/m³ SD = 25.74) to 5.95 µg/m³ (mean = 10.23 µg/m³ SD = 9.66) representing an 83% reduction in air nicotine levels (p < 0.01: 95% CI). Reductions were experienced in 16 of the 19 bars at follow up (Fig 2). The level of reduction ranged from 38–90%. At baseline higher concentrations were generally observed in small and medium sized bars than in large bars and super pubs. Only small and medium venues exceeded 60 µg/m³ at baseline and only small bars exceeded 20 µg/m³ at follow up. The three venues that experienced increases in air nicotine concentrations had low concentrations at baseline (< 6 µg/m³) and at follow up remained below the average (< 10 µg/m³). At baseline, three bars (16%) were below 6.8 µg/m³, at

| Table 1 Baseline profile of subjects (n = 35) in 15 venues |
|-----------------|----------------|----------------|
|                | Male n (%)     | Female n (%)   | Total n (%)  |
| Age            |                |                |              |
| 18-21          | 2 13           | 10 53          | 2 6          |
| 22-30          | 5 31           | 8 42           | 15 43        |
| 31-40          | 7 38           | 1 5            | 14 40        |
| 41-50          | 3 19           | 4 11           | 4 11         |
| People allowed to smoke in home | 11 69         | 7 37           | 18 51        |
| Length of time current job | 5 31         | 0 0            | 5 15         |
| < 6 months     |                |                |              |
| 6 months to 1 year | 1 6           | 9 50           | 10 30        |
| 1-2 years      | 5 31           | 3 17           | 8 24         |
| 3+ years       | 5 31           | 6 33           | 11 32        |
| Work duties    |                |                |              |
| Waiting        | 4 25           | 7 37           | 11 31        |
| Management     | 5 31           | 5 26           | 10 29        |
| Other          | 5 31           | 6 32           | 11 32        |
| Mixture        | 2 13           | 1 5            | 3 9          |

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follow up this increased to 10 (53%). Bars with designated outside smoking areas (n = 8) had higher concentrations of airborne nicotine at follow up than those that did not (n = 11) (mean 13 μg/m³, median 8.2 μg/m³, median 5.6 μg/m³, p = 0.31).

The mean number of smokers observed actively smoking indoors during four separate baseline visits to the venues ranged from 2.5–11.75; almost two thirds of bars (63%) had 4–8 active smokers. At follow up no smokers were observed smoking indoors.

**DISCUSSION**

The study found that since the introduction of the national workplace smoking ban in Ireland, significant reductions in both air nicotine (80%) and saliva cotinine concentrations (70%) have been experienced. This corresponds with the reported compliance rates with the smoking ban exceeding 90%23 and should result in substantially lowered health impacts.

When compared to data available for England, median cotinine concentrations of Galway hotel workers fell to less than a sixth (0.5 v 3.2 ng/ml) of those found in London bar workers15; however, much like other studies, exposure has not been totally eliminated11 12 with Galway hotel workers’ cotinine concentrations 25% higher than those found in all English non-smokers15 (0.5 v 0.4 ng/ml median) and more than twice those found in Galway school children from non-smoking homes24 (0.6 ng/ml v 0.267 ng/ml geometric mean: SD 0.969 ng/ml). This continuing exposure seems particularly the case for those undertaking bar work or waiting duties, whose work could involve them circulating in smoking areas or in areas in close proximity to where the migration of SHS into non-smoking areas from elsewhere may be a factor.

Biological markers are useful in quantifying exposure and an important step in linking exposure to relative risk of adverse outcomes.7 Whincup et al25 related the 20 year cardiovascular mortality of 4729 randomly selected men in the UK to their serum cotinine at study entry. In the first four years of follow up, those with more cotinine had 3.7 times greater risk of coronary heart disease (CHD) compared to those in the lowest quartile (< 0.7 ng/ml), and the proportion with major CHD was comparable to that of light smokers. The proportion of Galway hotel workers exposed above this cut off point (saliva cotinine equivalent 0.56 ng/ml) declined from 80% (28) before the ban to 49% (17) after the ban. Although the Galway bar workers were selected to have little or no home exposure (usually a major contributor to cotinine), those in the upper half of exposure had higher cotinine concentrations, with a median 12% higher and 75th centile 50% higher, respectively, than those in the Whincup et al25 study, which reported a significant dose–response trend.

Much of the recent data on the relationship of SHS and heart disease indicate the effect is acute, as well as chronic. Thus, healthy young non-smokers exposed to SHS for 30 minutes had compromised endothelium function comparable to that of smokers, activated platelets comparable to levels found in smokers, and increased presence of endothelial cell mortality in the blood27; for six hours after exposure, adverse impact on serum lipids were observed, including a significant decrease in the resistance of low density lipoprotein cholesterol (LDL-C) to oxidation and a significant increase in macrophage uptake of LDL-C.28 The recent experience of Helena, Montana, which had a 40% decrease in acute myocardial infarction after smoking was banned in workplaces and public places,29 demonstrates that short SHS exposures effects are not limited to cellular effects, but that SHS also affects the system as whole.
In addition, although any level of preventable exposure is unacceptable, it can be useful to interpret measured air nicotine concentrations in terms of the population risk thresholds they may exceed. Repace et al. estimated the point at which population risk decision thresholds would be exceeded for US regulatory authorities. They calculated that workplace environmental nicotine exposure of 2.0 mg/m³ for 45 years presented a lung cancer risk of three in 10,000, a de manifestis risk—that is, "a risk of obvious or evident concern" or risks so high that the US regulatory agencies have almost always acted to reduce them. All the bars in the Galway study exceeded this level both before and after the ban. However, the US Occupational Safety and Health Administration (OSHA), in evaluating urgently actionable risk to workers, uses a risk rate of one death per 1000 workers per WLT, called the "significant risk of material impairment of health" level. It has been estimated that "significant risk" would occur at or above 6.8 mg/m³ for lung cancer and at 0.68 mg/m³ for heart disease. In Galway, median air nicotine concentrations fell from 35.52 mg/m³ to 5.95 mg/m³ and whereas three bars met the OSHA risk threshold at baseline, an additional seven bars complied at follow up resulting in 53% (10) of bars falling below the "significant risk level for lung cancer only".

The implementation of the workplace smoking ban has been accompanied by a growth in café society, pavement cafes, and outdoor smoking shelters. The resultant new trend in smoking and its potential impact on outdoor and indoor exposures has received little consideration in the legislation. While a requirement for roofed outdoor smoking shelters is that not more than half of their perimeter is surrounded by one or more walls or similar structures, there is no legal requirement as to their minimum proximity to buildings nor is there a requirement as to the minimum proximity of smokers to open doors or windows. So that despite the dilution effect of outdoor air there is the potential for infiltration of tobacco smoke into indoor areas. Although there was an 83% reduction in air nicotine concentrations and median concentrations fell below 6.8 mg/m³, in those bars with designated outside smoking areas airborne nicotine concentrations were higher than in those bars without such an arrangement, suggesting that SHS may be migrating indoors. The extent to which outdoor smoke infiltration may be a contributor to indoor pollution and exposure levels in workers requires further investigation. Nevertheless, in the early weeks of the smoking ban significant reductions in exposure and associated risks have been observed. Based on an equation by Benowitz that relates airborne nicotine exposure to blood cotinine concentrations, we calculate that a 2–3 hour visit to a typical Galway pub would have exposed patrons to enough SHS with no other exposure to raise their cotinine above the referent level in Whincup et al. Before the smoking ban, three quarters of the pubs had measured levels of nicotine sufficient to produce this elevation, while after the ban fewer than 20% had such high levels of nicotine.

In recruiting hotel workers for the study, all staff who were on duty at the time the hotel was visited by a member of the study team were asked to participate. It must be acknowledged that this convenience sampling approach combined with the small sample size cannot ensure that the results are statistically representative of hotel workers in Galway city. This would have required a random sample of all hotel workers employed which was outside the time and cost constraints of the current study. In addition, by selecting only hotels for inclusion in the study, the sample is not representative of all hospitality establishments. This places limitations on the study in terms of its ability to make generalisations about the findings for the hospitality industry in Galway and indeed for Ireland as a whole. Because visits to obtain samples were unannounced and the shift patterns of the volunteers varied it was not possible to standardise the time into a shift that samples were obtained. At follow up our subjects were tested an hour and a half earlier into their shift than at baseline (mean 4.95 hours v 6.6 hours) equating to...
REFERENCES