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Heavy smoking and lung cancer: Are women at higher risk? Result of the ICARE study

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Background: Whether women are more or equally susceptible to the carcinogenic effects of cigarette smoke on the lungs compared with men is a matter of controversy. Using a large French population-based case–control study, we compared the lung cancer risk associated with cigarette smoking by gender.

Methods: The study included 2276 male and 650 female cases and 2780 male and 775 female controls. Lifetime smoking exposure was represented by the comprehensive smoking index (CSI), which combines the duration, intensity and time since cessation of smoking habits. The analysis was conducted among the ever smokers. All of the models were adjusted for age, department (a regional administrative unit), education and occupational exposures.

Results: Overall, we found that the lung cancer risk was similar among men and women. However, we found that women had a two-fold greater risk associated with a one-unit increase in CSI than men of developing either small cell carcinoma (OR = 15.9, 95% confidence interval (95% CI) 7.6, 33.3 and 6.6, 95% CI 5.1, 8.5, respectively; $P < 0.05$) or squamous cell carcinoma (OR = 13.1, 95% CI 6.3, 27.3 and 6.1, 95% CI 5.0, 7.3, respectively; $P < 0.05$). The association was similar between men and women for adenocarcinoma.

Conclusion: Our findings suggest that heavy smoking might confer to women a higher risk of lung cancer as compared with men.

Whether women are more or equally susceptible to the carcinogenic effects of cigarette smoke on the lungs compared with men is a matter of controversy. Several case–control studies have indicated that compared with non-smokers, the lung cancer odds ratio associated with a given increase in the amount and duration of cigarette smoking may actually be higher among women than among men (Brownson *et al*, 1992; Harris *et al*, 1993; Risch *et al*, 1993; Zang and Wynder, 1996). However, other case–control studies have not shown higher smoking-related risks among females than among males, including a recent large Italian-based case–control study (Schoenberg *et al*, 1989; Osann *et al*, 1993; De Matteis *et al*, 2013). In contrast, the results of four cohort studies published to date have been homogeneous and have found that the incidence of lung cancer among female smokers was about the same as that in male smokers after standardising for the

amount smoked (Bach *et al*, 2003; Bain *et al*, 2004; Haiman *et al*, 2006; Freedman *et al*, 2008). These results are not confirmed by a recent work based on a large UK medical research database which found, using the non-smokers as the reference group, that moderate/heavy smoker women are at higher risk for lung cancer than men, for the same level of smoking (Powell *et al*, 2013). The debate does not seem to be limited to lung cancer, as exemplified by the recent systematic review showing that women who smoke had a 25% greater risk of coronary heart disease than male smokers (Huxley and Woodward, 2011).

Most studies have used pack-years of cigarette smoking to investigate the potential modifying effect of gender in smoking-related lung cancer risk. However, this straightforward index of cumulative exposure makes strong assumptions on the equivalence of the roles of intensity and duration, an assumption that has been

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questioned for example in the context of smoking and lung cancer (Peto, 2012). Other metrics summarising smoking history have been used such as $\ln(1 + \text{pack-years}/5)$ (Risch *et al*, 1993; De Matteis *et al*, 2013) or $\ln(\text{cigarettes} + 1) \times \text{years of smoking}$ (Thurston *et al*, 2005). However, neither of these other metrics takes into account the time since cessation (TSC), an important parameter of lung cancer risk (Peto *et al*, 2000; Pirie *et al*, 2013). The comprehensive smoking index (CSI) is a single aggregate measure of smoking exposure that incorporates intensity, duration and TSC (Leffondre *et al*, 2006). As the CSI aggregates the three most important components of smoking history into a single variable, it is a practical tool for investigating the interactions between smoking and other factors.

Occupational exposure constitutes the other major determinant of lung cancer and has been regarded as a potential confounder in the comparison of the lung cancer risk related to smoking consumption between genders in some studies (Kreuzer *et al*, 2000; De Matteis *et al*, 2013).

The objective of this work was to estimate the overall effect of smoking on lung cancer risk, using the CSI separately for men and women and to investigate whether a modification effect could be observed. We considered all lung cancer together and each main histological type.

MATERIALS AND METHODS

Study population. The ICARE study, conducted from 2001 to 2007, is a large multicentre population-based case-control study of respiratory cancers. Details of the study design have been described previously (Luce and Stucker, 2011). The present analysis focused on women and men with lung cancer and their population controls. The eligible cases were aged <76 years at diagnosis. All histological types were included (ICDO codes C33–34). Among the 3360 eligible cases identified and contacted, 434 refused to participate (response rate = 87%). The present analysis included 650 female cases and 2276 male cases.

Population controls with no history of previous respiratory cancer were randomly selected through incidence density sampling from residential telephone directories (Rothman and Greenland, 2008). The controls were frequency matched to the cases by age (± 3 years), gender and department. Additional stratification was used to achieve a distribution by socioeconomic status among the controls comparable to that of the general population in each department. A total of 775 women out of the 1005 female controls eligible that were contacted agreed to participate (77%), and 2780 out of the 3406 male controls agreed to participate (81%).

The ICARE study was approved by the Institutional Review Board of the French National Institute of Health and Medical Research (IRB-Inserm, n° 01-036).

Data collection. We used a detailed questionnaire during face-to-face interviews. A detailed smoking history was collected for different time periods over the entire smoking lifespan. Periods of non-consumption of at least 1 year were also recorded and were considered when determining the duration of consumption.

Ever smokers were defined as subjects who had smoked at least 100 cigarettes in their lifetime and former smokers were defined as subjects who had stopped smoking for at least 2 years.

Lifetime occupational history was collected and coded according to the *International Standard Classification of Occupations* (ISCO) and the *French Nomenclature of Activities* (NAF). In 1982, a list (known as list A) of industries and occupations known to be associated with lung cancer was proposed and is a useful tool when considering occupational exposures. The latest update of this list (Consonni *et al*, 2010) was applied to our data to adjust for the burden of occupational exposures in this population.

Comprehensive smoking index. Lifetime smoking history was estimated by the CSI proposed for lung cancer (Leffondre *et al*, 2006), which incorporates duration of smoking (dur), time since cessation (tsc) and smoking intensity (int):

$$\text{CSI} = (1 - 0.5^{\text{dur}^*/\tau}) \times (0.5^{\text{tsc}^*/\tau}) \times \ln(\text{int} + 1),$$

where τ is the half-life parameter, δ is a lag time parameter that modulate the effects of TSC and total duration as follows: $\text{tsc}^* = \max(\text{tsc} - \delta, 0)$ and $\text{dur}^* = \max(\text{dur} + \text{tsc} - \delta, 0) - \text{tsc}^*$.

The lag time parameter δ accounts for the increase in lung cancer risk that has been observed in the first years after cessation (Leffondre *et al*, 2002). Tau (τ) gives information on the form of the dose-response curve between duration (or TSC) and the risk of lung cancer (Leffondre *et al*, 2006). A shorter τ implies a faster levelling-off of the impact of increasing both duration and TSC (see right panels of figures 1 and 2 in Leffondre *et al*, 2006). To estimate τ and δ , we estimated a series of logistic regression models, considering men and women together. Each model included the classical adjustment factors (see below) and the CSI as a continuous covariate, with the CSI calculated using a (τ , δ) pair selected from a dense two-dimensional grid. We searched a range of 11–50 years in 1-year increments for τ and a range of 0.1–2.7 in 0.1-year increment for δ . These ranges correspond to the limits of the 95% confidence intervals (CIs) for τ and δ found in two other case-control studies on lung cancer (Leffondre *et al*, 2006). The goodness of fit of the models was compared using Akaike's Information Criterion (AIC). The model that yielded the minimum AIC was identified and the corresponding values of τ (26 years) and δ (0.7 year) were used in all the analyses. A similar analysis developed in men and women separately indicated that these values of τ (26 years) and δ (0.7 years) led to the best goodness of fit.

Other dose metrics of smoking. For comparison purposes, we attempted to replicate our main results using the metrics $\ln(1 + \text{pack-years}/5)$ (Risch *et al*, 1993; De Matteis *et al*, 2013) or $\ln(\text{intensity} + 1) \times \text{duration}$ (Thurston *et al*, 2005) instead of the CSI.

Statistical analysis. The analysis was conducted only among ever smokers (i) to avoid the statistical instability due to the very small number of lung cancer cases among never smokers, especially among men and (ii) due to the strong association between lung cancer and gender by histological type among never smokers (i.e., OR = 7.0 (95% CI 4.6–10.6) and OR = 2.8 (95% CI 1.3–5.9)) for adenocarcinoma and squamous cell carcinoma, respectively, for women vs men).

The cutoffs were defined according to the quartiles of the CSI distribution among all the subjects (i.e., cases and controls, male and female). The reference class was the smallest CSI class.

We tested the linearity of the effect of CSI on the logit of lung cancer risk using a smoothing spline function of CSI with 4 degrees of freedom and the linearity was not rejected. We considered therefore the CSI as a continuous variable and estimated the ORs of lung cancer per unit of CSI and formally tested the interactions between gender and CSI by using the likelihood ratio test.

All of the models were adjusted for age, department, education, and occupational exposure (list A).

RESULTS

The overall socio-demographic characteristics of the cases and controls and the histological subtypes of the lung cancer cases are presented in Table 1. Half of the female cases had adenocarcinomas (53%), whereas we observed similar proportions of squamous cell and small cell carcinomas (17% and 15%, respectively). For male cases, the percentages were 33% adenocarcinomas, 35% squamous cell carcinomas and 14% small cell carcinomas.

Table 1. Distribution of cases and controls among men and women by socio-demographic characteristics and histological type among cases

	Male							Female						
	Cases		Controls		OR ^a	95% CI	P-value	Cases		Controls		OR ^a	95% CI	P-value
	N	%	N	%				N	%	N	%			
Total	2275		2780					648		775				
Age at interview, years														
<50	310	14	663	24	1.0	Ref.		160	25	167	21	1.0	Ref.	
50–60	773	34	855	31	1.9	1.6–2.3		224	35	160	21	1.5	1.1–2.1	
60–70	824	36	923	33	1.9	1.6–2.2		171	26	263	34	0.7	0.5–1.0	
≥70	368	16	339	12	2.3	1.9–2.9	<i>P</i> <0.001	93	14	185	24	0.6	0.4–0.8	<i>P</i> <0.001
Educational level^b														
Elementary school or less	675	34	521	19	1.0	Ref.		204	34	242	32	1.0	Ref.	
Middle school	869	43	1081	41	0.6	0.5–0.7		208	35	270	36	0.6	0.5–0.8	
High school	185	9	310	12	0.4	0.4–0.6		72	12	90	12	0.7	0.5–1.0	
University	274	14	752	28	0.3	0.2–0.3	<i>P</i> <0.001	110	19	149	20	0.5	0.4–0.8	<i>P</i> <0.001
Occupational exposure (list A)^c														
Non-exposed	1826	81	2450	88	1.0	Ref.		623	97	767	99	1.0	Ref.	
Exposed	438	19	329	12	1.8	1.5–2.1	<i>P</i> <0.01	21	3	8	1	3.0	1.3–6.9	<i>P</i> <0.001
Histological types by smoking status														
Never smokers (CSI = 0)	59		813					188		513				
Squamous cell carcinoma	12	20	—					23	12	—				
Small cell carcinoma	2	4	—					6	3	—				
Adenocarcinoma	32	54	—					133	70	—				
Other types	13	22	—					27	14	—				
Multiple phenotype ^d	—		—					1		—				
Ever smokers (CSI > 0)	2191		1958					457		260				
Squamous cell carcinoma	786	36	—					82	18	—				
Small cell carcinoma	330	15	—					89	19	—				
Adenocarcinoma	759	35	—					221	48	—				
Other types	336	15	—					70	15	—				
Multiple phenotype ^d	20	3	—					5		—				

Abbreviations: CI = confidence interval; CSI = comprehensive smoking index; OR = odds ratio.
^aOdds ratio of lung cancer adjusted for age at interview (<50, 50–60, ≥60) and department.
^bEducational level was not available in the summarized questionnaire.
^cData missing for 12 men (11 cases and 1 control) and 4 women cases.
^dMultiple phenotype corresponds to tumours with several histological entities. Subjects with multiple phenotypes must be subtracted from the total to not be counted more than once.

The range of the values of the CSI was 0–2.71 among controls and 0–3.08 among cases ever smokers. Mean pack-years, intensity, duration and TSC across CSI classes in men and women controls are presented in Table 2. As can be seen, women classified in the first two classes of CSI had a smoking history lighter than men, mainly because of a lower average intensity. We found that the effect of smoking by CSI class on lung cancer was quite similar for men and women. The OR associated with a one-unit increase in CSI was also close and not significantly different (*P* = 0.5).

Table 3 presents the OR for the three histological types of lung cancer associated with the CSI. For squamous cell lung cancer, the OR was two-fold greater in women than in men (*P* = 0.04). In contrast, for adenocarcinomas the OR was higher in men than in women but the difference was not significant. Finally, the analysis for small cell lung cancer showed an OR two-fold greater in women than in men (*P* < 0.05).

Table 4 presents the ORs of lung cancer associated with the CSI (model 1), with ln (1 + PY/5) (model 2) or ln (intensity + 1) × duration (model 3) in our population of men and women ever smokers, each metric of smoking consumption being considered as

a continuous variable. Each model included an interaction term between the metric of smoking consumption and sex. Models 2 and 3 additionally included the TSC, plus an interaction term with sex. When all lung cancers were pooled together, the fits to data of models 1 and 3 were equal and both better than model 2. The ORs of lung cancer were systematically higher for men smokers than for women smokers although not significant. Model 1 in squamous and small cells lung cancer yielded a better AIC than models 2 or 3. The ORs were higher in women than in men whatever the metric used, but interaction with sex was significant only in model 1. On the other hand for adenocarcinoma the fits to data of models 1 and 3 were similar and better than model 2. The OR for adenocarcinoma was stronger in men smokers than in women smokers.

DISCUSSION

In this work, we have shown that when comparing lung cancer risks associated with smoking between genders, results vary according to histological type.

Table 2. Means of pack-years, intensity, duration, time since cessation among controls, by categories of CSI and estimated odds ratio for the association between lung cancer and CSI among the ever smokers

CSI	Pack-years ^a	Intensity ^a (cig/day)	Duration ^a (years)	TSC ^a (years)	Cases (%)	Controls (%)	Ever smokers	
							OR ^b	95% CI
Male					N = 2191	N = 1958		
≤0.73	8.4	11.3	13.3	27.6	202 (9)	859 (44)	1.0	Ref.
0.73–1.40	22.8	16.3	27.2	14.1	447 (20)	582 (30)	3.3	2.6–4.1
1.40–1.90	34.2	18.8	35.1	7.6	643 (29)	347 (18)	8.0	6.4–10.1
> 1.90	58.1	25.0	43.7	5.0	899 (41)	170 (9)	20.3	15.8–26.0
Dose–response ^c							5.5	4.9–6.3
Female					N = 457	N = 260		
≤0.73	3.7	6.9	11.5	25.0	48 (10)	116 (45)	1.0	Ref.
0.73–1.40	16.7	12.2	27.0	12.0	94 (21)	84 (32)	2.7	1.6–4.5
1.40–1.90	31.0	18.8	33.8	5.8	183 (40)	45 (17)	9.1	5.4–15.2
> 1.90	54.9	28.7	41.9	4.2	132 (29)	15 (6)	20.0	10.0–40.0
Dose–response ^c							4.9	3.6–6.7
P-value for interaction gender × CSI							P = 0.5	

Abbreviations: CI = confidence interval; CSI = comprehensive smoking index; OR = odds ratio; TSC = time since cessation.
^aStatistics calculated among controls.
^bOdds ratio of lung cancer associated with CSI, adjusted for age at interview (<50, 50–60, ≥60), départements, education and occupational exposure (list A).
^cThe CSI is considered continuously.

Table 3. Estimated odds ratio for the association between CSI and each histological type of lung cancer

	CSI ^a	Ever smokers			
	mean ± s.d.	Cases (%)	Controls N (%)	OR ^b	95% CI
Squamous cell carcinoma					
Male	1.71 ± 0.60	786 (98)	1958 (71)	6.1	5.0–7.3
Female	1.68 ± 0.48	82 (78)	260 (34)	13.1	6.3–27.3
P-value for sex × CSI interaction					P = 0.04
Adenocarcinoma					
Men (N = 759)	1.61 ± 0.60	759 (96)	1958 (71)	4.9	4.2–5.9
Women (N = 221)	1.43 ± 0.59	221 (62)	260 (34)	3.6	2.5–5.2
P-value for sex × CSI interaction					P = 0.13
Small cell carcinoma					
Male (N = 330)	1.72 ± 0.55	330 (99)	1958 (71)	6.6	5.1–8.5
Female (N = 89)	1.76 ± 0.51	89 (94)	260 (34)	15.9	7.6–33.3
P-value for sex × CSI interaction					P = 0.03

Abbreviations: CI = confidence interval; CSI = comprehensive smoking index; OR = odds ratio.
^aMean of CSI calculated among cases ever smokers, according to gender and histology.
^bOdds ratio of lung cancer associated with CSI (considered continuously), adjusted for age at interview (<50, 50–60, ≥60 years), départements, education and occupational exposure (list A).

Synthesizing the various dimensions of smoking information to better capture the smoking history represents a challenge, and the most common approach has been to use pack-years. However, this index does not include the delay since smoking cessation, which is also an important component of lung cancer risk (Peto *et al*, 2000). Cohort studies have compared the incidence rate of lung cancer between gender stratified by smoking categories including a category for ex-smokers, even though this complicates the global assessment of the gender/smoking interaction (Freedman *et al*, 2008).

We estimated the two parameters (δ and τ) of the CSI considering men and women together to avoid introducing

variability between genders in the shape of the dose–response curves that are determined by τ . Furthermore, we obtained similar values for δ and τ when we checked that these estimates also corresponded to the best goodness of fit when considering men and women separately. We treated the CSI as a continuous variable in the logistic models and restricted the population to the ever smokers as previous studies had done (Brownson *et al*, 1992; Harris *et al*, 1993; Zang and Wynder, 1996; De Matteis *et al*, 2012, 2013).

Several case–control studies (Schoenberg *et al*, 1989; Osann *et al*, 1993; Kreuzer *et al*, 2000; De Matteis *et al*, 2013) and

Table 4. Estimated odds ratio of lung cancer associated with smoking, according to smoking metric, for all lung cancer and by histological type among ever smokers; comparison of the fit of models 1, 2 and 3

Smoking metric (model ^a)	All lung cancer				Squamous cell carcinoma				Small cell carcinoma				Adenocarcinoma			
	AIC ^b	Males OR ^c 95% CI	Females OR ^c 95% CI	P ^d	AIC ^b	Males OR ^c 95% CI	Females OR ^c 95% CI	P ^d	AIC ^b	Males OR ^c 95% CI	Females OR ^c 95% CI	P ^d	AIC ^b	Males OR ^c 95% CI	Females OR ^c 95% CI	P ^d
Model 1	4578	5.5 (4.9–6.3)	4.9 (3.6–6.7)	0.40	2337	6.1 (5.0–7.3)	13.1 (6.3–27.3)	0.04	1514	6.6 (5.1–8.5)	15.9 (7.6–33.3)	0.03	2791	4.9 (4.2–5.9)	3.6 (2.5–5.2)	0.10
Model 2	4587	4.0 (3.4–4.7)	3.1 (2.2–4.3)	0.10	2348	4.1 (3.3–5.2)	5.1 (2.7–9.7)	0.50	1537	4.0 (3.0–5.3)	5.6 (3.0–10.5)	0.32	2802	3.8 (3.1–4.7)	2.6 (1.8–3.9)	0.09
Model 3	4580	1.3 (1.3–1.4)	1.3 (1.2–1.3)	0.10	2350	1.3 (1.3–1.4)	1.3 (1.2–1.5)	0.80	1538	1.29 (1.2–1.4)	1.4 (1.2–1.5)	0.20	2790	1.32 (1.3–1.4)	1.2 (1.1–1.3)	0.07

Abbreviations: CI = confidence interval; CSI = comprehensive smoking index; OR = odds ratio; PY = pack-years; TSC = time since cessation.
^aModel 1: CSI, sex, CSI × sex; Model 2: ln(1 + PY/5), sex, ln(1 + PY/5) × sex, TSC, TSC × sex; Model 3: ln(intensity + 1) × duration, sex, ln(intensity + 1) × duration × sex, TSC, TSC × sex (CSI, PY, intensity, duration and TSC are considered continuously).
^bAIC, Akaike's information criterion with lower AIC indicating the best fit to data.
^cOdds ratio of lung cancer associated with smoking metric adjusted for age at interview (<50, 50–60, ≥60), départements, education and occupational exposure (list A).
^dP-value for interaction sex × smoking metrics (using the likelihood ratio test with 1 degree of freedom for model 1, and 2 degrees of freedom for models 2 and 3).

prospective studies (Bach *et al*, 2003; Bain *et al*, 2004; Haiman *et al*, 2006; Freedman *et al*, 2008) have found no overall differences in risk between genders. Although we also found no overall difference in risks between genders, the risk of squamous and small cell lung cancer was higher in women as compared with that in men. The high proportion of adenocarcinoma (38%) explained why we did not observe any overall difference. Squamous and small cells lung cancer are the histological types the most strongly related to smoking (Khuder and Mutgi, 2001). In our population, women with squamous cells and especially small cells lung cancer were heavy smokers. Their average CSI was close to that of male cases for squamous cells (1.71 vs 1.68 in women and men, respectively) and even higher for small cells (1.76 vs 1.72 in women and men, respectively). The same mean estimated on the whole lung cancer group is lower in women than in men (1.55 and 1.66, respectively). It is noteworthy that the higher associations among women than among men were also found for these two histological types in the previous studies (Brownson *et al*, 1992; Harris *et al*, 1993; Risch *et al*, 1993; Zang and Wynder, 1996), a result confirmed by the meta-analysis from Khuder and Mutgi (2001).

It could therefore be likely that heavy smoking carries a higher risk for women than for men. Lung cancer in men and women differs in many aspects, survival, histology, genetic alterations in tumours, which suggests that carcinogenesis might be different in men and women (Novello and Baldini, 2006; O'keeffe and Patel, 2008; Gasperino, 2010; Kiyohara and Ohno, 2010). A recent work based on a UK medical research database, comprising 12 121 incident cases of lung cancer and 48 216 controls, also found a higher risk of lung cancer in women than in men, that increased with the level of smoking assessed by the highest intensity ever smoked (Powell *et al*, 2013). Various reasons could support a deleterious effect of smoking higher in women than in men including possibly differential enzymatic activity of genes involved in tobacco smoke metabolism (Mollerup *et al*, 2006). The role of hormonal and reproductive factors on female lung carcinogenesis is currently heavily debated, and a possible interaction between smoking and hormonal factors should also be considered. Such an interaction between smoking and estrogens replacement therapy has been observed once, but to our knowledge never reproduced (Taioli and Wynder, 1994). In addition, EGFR and KRAS, two driver mutations of lung carcinogenesis, are now well known for their particular distribution with gender and smoking history (Cooper *et al*, 2013). A large recent study specific for adenocarcinoma showed a specificity of the type of K-RAS mutations with sex and also, interestingly, that women with this specific mutation had a significantly lower pack-year consumption

as compared with men (Dogan *et al*, 2012). A similar result had been previously found with P53, a tumor suppressor gene frequently mutated in lung cancer (Kure *et al*, 1996; Toyooka *et al*, 2003). Altogether, these findings may also support a possible higher susceptibility of tobacco carcinogenesis in women.

Finally, this higher susceptibility might not be specific of lung cancer, since a large review of cigarette smoking and coronary heart disease also shows that women present a risk ratio relative to men of 1.25 (1.12–1.39) (Huxley and Woodward, 2011).

A comparison of our findings with those of similar studies is not easy because we are the first to have compared the effects of smoking between genders using the CSI to estimate the lifelong smoking history. The most recent publication on this topic, using a different metric (i.e., ln(1 + PY/5) to evaluate the lifelong smoking history, did not find any difference in risk between genders (De Matteis *et al*, 2013). Their population and ours are similar in many respects: they are two European populations, with similar gender distribution, displaying a similar proportion of highly educated men and women (high school or more). The proportion of never smokers among cases in men as in women is very close, the mean pack-years of cigarette smoking among women ever smokers were almost identical to our numbers (i.e., 33 and 15 in women cases and controls, respectively) and the average delay since smoking cessation almost identical in cases and controls, among men as well as among women. All these comparisons demonstrate a level of smoking intoxication similar in the two populations. However, although the De Matteis *et al* study found slightly higher odds ratios in women than in men for squamous and small cells lung cancer associated with smoking, they did not find any statistical interactions. However, their number of women with squamous and small cells lung cancer (38 and 35, respectively) was smaller than in this study. To compare our results with already published studies, including De Matteis *et al*, we examined the smoking-gender interaction with a logistic regression that included the cumulative exposure index estimated with ln(1 + PY/5), in addition to an interaction term between sex and TSC to make models 2 (i.e., ln(1 + PY/5)) and 3 (i.e., ln(intensity + 1) × duration) more comparable to the CSI. Our results also became non-significant, although the ORs associated with smoking were still higher in women than in men. These last results demonstrate that the manner in which smoking history is synthesized is clearly critical. Because the range of the values of each metric is clearly different, it is not possible to compare the values of the ORs between the three models. Time since cessation modulates the risk of lung cancer (Pirie *et al*, 2013). Comprehensive smoking index integrates this important parameter in the assessment of lifelong smoking history

and it is interesting to note that the best AIC is achieved with the model that includes CSI while models 2 and 3 treat TSC as a confounder (including an interaction term with sex).

Among the criticism also often levelled at studies indicating higher risks for women is that baseline risks are likely to be different in never-smoking women than in never-smoking men (i.e., less work-related exposure to lung carcinogens for women), although this was not observed in Freedman's cohort (Freedman *et al*, 2008). To consider this possible confounder, we adjusted our results for occupational exposure through the typical variable List A that we already used in a previous work focused specifically on occupational exposures (Guida *et al*, 2011). We also made an additional adjustment for the type of tobacco smoked (light tobacco, dark tobacco or mixed). This did not alter any result (data not shown).

Recall bias in case-control design is always a matter of concern. However, in the topic we are interested in, the problem is less in the validity of lifelong smoking history reported than in a potential differential recall bias between men and women. Indeed, if women tend to underreport their true consumption more than men or to a lesser extent, the consequence would tend to overestimate the risk in women as compared with men. This hypothesis should also be considered although this bias is not specific to the case-control design. Cohort studies also rely on self-questionnaire or face-to-face interviews at baseline and during follow-up. It could be interesting to examine this hypothesis in the near future, using both cotinine biomarker and self-report to assess smoking and then comparing results in men and women.

Our study also has several strengths: we included incident cases, population controls were randomly selected in the same *départements* as cases through incidence density sampling, the participation rate was over 80% in cases and controls and subjects were interviewed face to face by trained interviewers using a structured questionnaire.

In summary, to resolve the question of whether women have a greater susceptibility to smoking-related lung cancer than men, it is important to consider the results both for all lung cancer cases and also by histological type. Incorporating time since smoking cessation in the smoking metric seems to better summarize the lifelong history. Further studies using the CSI instead of pack-years of cigarette smoking to compare the smoking effects on lung cancer in men and women are needed to move forward in this debate. The results we obtained for small cells and squamous cells lung cancer might also suggest that women heavy smokers are more susceptible to tobacco smoking. This hypothesis should be considered like any others, especially as the aetiology of women lung cancer is known only by the knowledge obtained in men.

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CONFLICT OF INTEREST

The authors declare no conflict of interest.

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