

Lung cancer risk of solid fuel smoke exposure: a systematic review and meta-analysis

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ABSTRACT: The aim of this systematic review was to quantify the impact of biomass fuel and coal use on lung cancer and to explore reasons for heterogeneity in the reported effect sizes.

A systematic review of primary studies reporting the relationship between solid fuel use and lung cancer was carried out, based on pre-defined criteria. Studies that dealt with confounding factors were used in the meta-analysis. Fuel types, smoking, country, cancer cell type and gender were considered in sub-group analyses. Publication bias and heterogeneity were estimated.

The pooled effect estimate for coal smoke as a lung carcinogen (OR=1.82, 95% CI 1.60, 2.06) was greater than that from biomass smoke (OR=1.50, 95% CI 1.17, 1.94). The risk of lung cancer for solid fuel use was greater in women (OR=1.81, 95% CI 1.54, 2.12) compared to men (OR=1.16, 95% CI 0.79, 1.69). The pooled effect estimates were 2.33 (95% CI 1.72, 3.17) for adenocarcinoma, 3.58 (1.58, 8.12) for squamous cell carcinoma, and 1.57 (1.38, 1.80) for tumours of unspecified cell type.

These findings suggest that in-home burning of both coal and biomass is consistently associated with an increased risk of lung cancer.

Keywords: Adenocarcinoma, biomass fuel, coal, indoor air pollution, squamous cell carcinoma

Introduction: Lung cancer is one of the leading mortality causes accounting for 1.3 million deaths annually worldwide [1]. While smoking is the major risk factor, 25% of cases are not attributable to tobacco use [2]. Epidemiological studies have shown that while globally lung cancer in never smokers is consistently more common in women than in men, geographical variations are substantial [2]. In East and South Asia, up to 83% of female lung cancer cases are never smokers, compared to 15% in the United States [2]. In developing countries, an estimated 2.4 billion (70%) people use biomass (wood, charcoal, crop residues, dung) or coal, collectively known as solid fuels, for cooking and heating [3]. Emissions from combustion of solid fuels have been shown to have high concentrations of polycyclic aromatic hydrocarbons (PAHs), benzo[a]pyrene (BAP) and particulate matter with diameter of 2.5 μm or less ($\text{PM}_{2.5}$), which in turn have been associated with high lung cancer rates [2].

Recently, indoor emissions from household combustion of coal and biomass (mostly wood) have been classified as carcinogenic (Group 1) and probably carcinogenic (Group 2A) respectively to humans [4]. However, data on the magnitude of lung cancer risk and the histological sub-type of lung cancer associated with solid fuel use are few. In the literature, four meta-analyses were identified, but three [5-7] were limited to studies conducted in China and one [8] focused only on coal use. A recent paper included a pooled estimate from several countries, but data were restricted only to studies from an international consortium [9].

In this meta-analysis we reviewed papers from all countries and calculated pooled estimates of the association of the use of solid fuels and lung cancer. We investigated whether these effects were influenced differently by other factors, notably the types of fuel used, smoking (including environmental tobacco smoke [ETS]), and study location. We also looked at whether there was a pattern of association between smoke exposure and lung cancer histological sub-type.

METHODS

Papers published from January 1980 to October 2010 were identified through a systematic literature search in Ovid MEDLINE, EMBASE and Google Scholar. Search terms used for the initial search on exposure were “biomass”, “biofuel”, “organic fuel”, “black smoke”, “wood”, “indoor air pollution”, “carbon monoxide”, “respirable dust”, “solid fuel”, “dung”, “charcoal”, “crop residue” and outcomes were “carcinogen”, “lung tumour”, “adenoma”, “adenocarcinoma”, “squamous carcinoma”, “carcinoma”, “lung cancer” and “cancer”. The articles obtained by using different exposure search terms were combined using “OR” and the same was done for outcomes. The combining term “AND” was used to combine the article obtained for exposure and outcome. References in each of the identified papers were screened for any articles that were not identified in the original search. There was no restriction on language in the original search but articles in English and Chinese were retained for inclusion in the meta-analysis. The search was carried out by two co-authors (PA and OK).

Study selection

All potentially relevant manuscripts were reviewed. Selection criteria were identified and defined by all co-authors. For studies to be a part of the review and meta-analysis they had to meet the criteria listed in Box 1. Most studies considered were those in which cases had cytological/histological findings alongside radiological confirmation. However, a minority of the studies where the assessment technique was not stated were still included in the review. No limitations were set for the age of participants in the studies or for the definition of exposure to solid fuels.

Box 1: Inclusion criteria for meta-analysis

1. Papers of primary studies written in English or Chinese
2. Case-control, cross sectional or cohort study design that controlled for smoking.
3. Solid fuel used primarily for household cooking and/or heating in the study population
4. Provided adjusted odds ratios (ORs) or relative risks (RRs) to measure the association between lung cancer and exposure to solid fuels with corresponding 95% confidence intervals (CIs) or p-values
5. Specify the technique by which exposure and lung cancer were assessed and ascertained, (although we specified no definitive criteria)

Data extraction

Selection of studies was undertaken at each stage by two co-authors (PA and OK) for studies written in English and KL for studies written in Chinese. Disagreements were settled by consensus. All data

were extracted by two co-authors (PA and KL) independently and uncertainties were discussed with all co-authors. We used the Newcastle-Ottawa Quality Assessment Scale to assess the quality of the studies [10]. A pre-defined form was then used to extract information from selected studies under the following headings - author, journal, year, country of study, organisation/funding body, type of fuel considered, study design, smoking (type, measure and assessment technique), sample size, indoor air pollution exposure assessment, primary outcome (type and assessment of outcome), effect size (relative risk [RR] or odds ratio [OR] and the associated 95% confidence intervals [CI] and p-values) and possible confounding factors considered.

Statistical analysis

Initially all studies were pooled and a sensitivity analysis was carried out to assess the impact of methodological concern by grouping them into different sub-groups, which include fuel types (biomass/mixed fuel/coal), gender (female only/male only/male and female), cancer histological sub-type (unspecified/adenocarcinoma/squamous carcinoma), adjustment for smoking (yes/non-smokers only), adjustment for ETS exposure (yes/no), study design (population-/hospital-based), sample size (>368 [median]/≤368), study location (China/Taiwan/India/other countries), year when study was conducted (2000 onwards/before 2000), year of publication (2000 onwards/before 2000), language of publication (Chinese/English), Newcastle-Ottawa score (>6 [median]/≤6), and the quality of exposure assessment based on the Newcastle-Ottawa criteria (1/2/3 stars). The natural logarithm of OR and the associated standard error (SE) were used to estimate the effect size of all studies and the sub-groups. Within-group heterogeneity was evaluated using Q-tests and/or I^2 statistics. Heterogeneity between different studies was visually explored using Galbraith plots, and sources of heterogeneity were systematically examined by meta-regression. We used random effects models as there was significant heterogeneity on Q-tests ($p < 0.05$) and/or I^2 statistic value >50%. Begg's funnel plot and Egger's test were used to assess publication bias [11]. All analyses were performed in STATA (version 11, STATA, College Station, Texas, USA).

RESULTS

The initial search revealed 11398 articles of which 2012 duplicates and 7908 irrelevant papers were removed by screening the titles. The abstracts of the remaining 1478 papers were reviewed and 203 were selected for full paper review, of which 51 papers were related to lung cancer and solid fuel use (Figure 1). Twenty eight studies (Table 1) were included in the meta-analysis, the rest (23 papers) being excluded (Supplementary Table S1) either because of failure to meet the inclusion criteria or because data were unusable, or both. The results presented are from 12419 cases and 34609 controls.

Table 1: Summary of studies included in the meta-analysis

References	Study location	Study period	Study design	Fuel type	Gender	Cancer type	Cases	Controls
Huang <i>et al</i> 1992 [12]	China	1990-1991	Hospital-based case-control	Coal	M & F	Unspecified	135	135
Lan and He 2004 [13]	China	1995-1996	Population-based case-control	Coal	M & F	Unspecified	122	122
Lan <i>et al</i> 2002 [14]	China	1976-1992	Population-based case-control	Coal	F	Unspecified	684	9380
				Coal	M	Unspecified	700	10648
Liu <i>et al</i> 1993 [15]	China	1983-1984	Hospital-based case-control	Coal	F	Unspecified	92	92
				Coal	M	Unspecified	224	224
Liu <i>et al</i> 1991 [16]	China	1985-1986	Population-based case-control	Biomass	F	Unspecified	54	202
				Biomass	M	Unspecified	56	224
Luo <i>et al</i> 1996 [17]	China	1990-1991	Population-based case-control	Coal	M & F	Squamous	39	306
Sun <i>et al</i> 1991 [#] [18]	China	1985-1987	Population-based case-control	Coal	F	Unspecified	418	398
Sun <i>et al</i> 2002 [#] [19]	China	1996-1999	Population-based case-control	Coal	M & F	Unspecified	206	618
Zhong <i>et al</i> 1999* [20]	China	1992-1994	Population-based case-control	Coal	F	Unspecified	504	601
Wu-Williams <i>et al</i> 1990 [21]	China	1985-1987	Population-based case-control	Coal	F	Unspecified	956	953
Xu <i>et al</i> 1989 [22]	China	1985-1987	Population-based case-control	Coal	F	Unspecified	520	557
				Coal	M	Unspecified	729	788
Galeone <i>et al</i> 2008 [23]	China	1987-1990	Hospital-based case-control	Coal	M & F	Unspecified	216	435
Lin <i>et al</i> 1996* [#] [24]	China	1985-1990	Population-based case-control	Coal	F	Adenocarcinoma	122	122
Hao 1998 [#] [25]	China	1981-1986	Population-based case-control	Coal	M & F	Unspecified	220	440
Lu <i>et al</i> 2003 [†] [26]	China	1998-2001	Population-based case-control	Coal	M & F	Unspecified	445	445
				Coal	M & F	Squamous	185	185
Liang <i>et al</i> 2004 [#] [27]	China	2001-2002	Hospital-based case-control	Coal	M & F	Adenocarcinoma	89	89
Huang <i>et al</i> 1999 [#] [28]	China	1993-1996	Hospital-based case-control	Coal	M & F	Unspecified	122	244
Ger <i>et al</i> 1993 [29]	Taiwan	1990-1991	Population-based case-control	Coal	M & F	Squamous	59	118
Ko <i>et al</i> 1997* [30]	Taiwan	1992-1993	Hospital-based case-control	Biomass	F	Unspecified	91	89
				Coal	F	Unspecified	52	66
Lee <i>et al</i> 2001 [31]	Taiwan	1993-1999	Hospital-based case-control	Biomass	F	Adenocarcinoma	162	273
				Coal	F	Adenocarcinoma	162	273
				Biomass	F	Squamous	84	134
				Coal	F	Squamous	84	134
Sapkota 2008 [32]	India	2001-2004	Hospital-based case-control	Biomass	M & F	Unspecified	381	237
				Coal	M & F	Unspecified	35	10
Gupta <i>et al</i> 2001 [33]	India	1995-1997	Hospital-based case-control	Mixed	F	Unspecified	30	90
				Mixed	M	Unspecified	232	431
Sobue 1990* [34]	Japan	1985	Hospital-based case-control	Biomass	F	Unspecified	144	731
Hernandez-Garduno <i>et al</i> 2004* [35]	Mexico	1986-1994	Hospital-based case-control	Biomass	F	Unspecified	113	273
Sasco <i>et al</i> 2002 [36]	Morocco	1996-1998	Hospital-based case-control	Coal	M & F	Unspecified	118	235
Wu <i>et al</i> 1985 [37]	USA	1981-1982	Population-based case-control	Coal	F	Adenocarcinoma	149	149
				Coal	F	Squamous	71	71
Ramanakumar 2007 [38]	Canada	1996-2001	Population-based case-control	Mixed	F	Unspecified	315	381
				Mixed	M	Unspecified	438	588
Lissowska <i>et al</i> 2005 [39]	Europe ^{††}	1998-2001	Hospital/population case-control	Biomass	M & F	Unspecified	2861	3118

*Studies with non-smoking participants only

[#]Papers published in Chinese^{††}Czech Republic, Hungary, Poland, Romania, Russia, Slovakia, and United Kingdom

Effect estimates

The pooled effect estimate size was obtained using the random effect model because of heterogeneity across studies (Q-statistic=107.30, degrees of freedom=40, $p<0.001$; $I^2=62.7\%$; tau-squared=0.081 and $Z=7.99$, $p<0.001$). The pooled OR with 95% CI was 1.70 (1.50, 1.94) for all studies.

Sub-group analyses were performed using random effect models. The OR values related to biomass, mixed fuel and coal were 1.50 (95% CI 1.17, 1.94), 1.13 (0.52, 2.46) and 1.82 (1.60, 2.06) respectively (Figure 2) (For other forest plots see Supplementary Figures S1-S8). Coal contributed 68.8% to the pooled effect sizes of lung cancer followed by biomass (19.8%) and mixed fuel (11.5%). The associated risk for women was greater compared to that for men ($p = 0.034$) (Table 2). The greater risk observed in the Chinese publications compared to those in English ($p=0.006$) remained after adjusting for potential confounders including types of fuel used, gender, smoking, and quality of the study assessed by Newcastle-Ottawa score. The same trend was found in both smoking and non-smoking participants.

Studies were then stratified according to the type of fuel used, then by various sub-groups (Table 3). No significant heterogeneity was observed in the different strata for studies related to the exposure to biomass smoke but heterogeneity among hospital based studies approached significance ($I^2 54.3\%$; $p=0.053$). On the other hand, there was significant heterogeneity among studies with coal smoke exposure in relation to squamous cell carcinoma ($I^2 61.2\%$, $p=0.035$), unspecified types of lung cancer ($I^2 38.1\%$; $p=0.047$), women only ($I^2 45.5\%$; $p=0.043$), population-based ($I^2 60.4\%$; $p=0.001$) and hospital-based studies ($I^2 43.4\%$; $p=0.008$), and those with sample size less than or equal to 368 ($I^2 49.3\%$; $p=0.019$) and for.

Of the 28 studies included in the meta-analysis, 14 collected data on ETS exposure and merely seven made adjustments for ETS. Even more surprising was the fact that only three out of seven of the female-only studies that measured ETS actually adjusted for it. Pooled effect estimates from studies that adjusted for ETS (1.28, 95% CI 0.91, 1.80) was significantly lower ($p=0.034$) compared to those that did not (1.91, 95% CI 1.65, 2.22).

The studies with poor quality, particularly in the exposure assessment, as measured by the Newcastle-Ottawa score, tend to report greater effect size (Tables 2, 3 and Supplementary Table S7).

Table 2: Sub-group analyses of the lung cancer risk associated with the use of solid fuels

Sub-group types	Studies (n)*	Heterogeneity (I ² (%); p-value)	OR (95% CI)	p-value
Types of solid fuel used				
<i>Biomass fuel (BM)</i>	7	41.2; 0.092	1.50 (1.17, 1.94)	0.180 (BM vs. C) 0.235 (MF vs. C)
<i>Mixed fuel (MF)</i>	2	89.4; <0.001	1.13 (0.52, 2.46)	
<i>Coal (C)</i>	22	43.4; 0.008	1.82 (1.60, 2.06)	
Gender				
<i>Female only (F)</i>	12	36.8; 0.051	1.81 (1.54, 2.12)	0.034 (F vs. M)
<i>Male only (M)</i>	6	80.1; <0.001	1.16 (0.79, 1.69)	
<i>Male and female[#]</i>	13	66.0; <0.001	1.93 (1.53, 2.44)	
Cancer histological sub-type				
<i>Unspecified</i>	22	64.1; <0.001	1.57 (1.38, 1.80)	0.335 (Adeno. vs. Squamous)
<i>Adenocarcinoma</i>	4	0.0; 0.553	2.33 (1.72, 3.17)	
<i>Squamous carcinoma</i>	5	51.8; 0.065	3.58 (1.58, 8.12)	
Adjustment for smoking				
<i>Yes</i>	24	64.6; <0.001	1.70 (1.47, 1.96)	0.710
<i>Non-smokers only</i>	7	69.2; 0.001	1.85 (1.21, 2.81)	
Adjustment for environmental tobacco smoke				
<i>Yes</i>	9	65.6; 0.003	2.27 (1.31, 3.96)	1.709
<i>No</i>	32	63.1; <0.001	1.67 (1.46, 1.91)	
Study design				
<i>Population-based</i>	15	71.3; 0.106	1.83 (1.51, 2.21)	0.402
<i>Hospital-based</i>	12	37.8; 0.055	1.63 (1.34, 1.97)	
Sample size				
<i>>368[¶]</i>	17	72.7; <0.001	1.60 (1.36, 1.87)	0.110
<i>≤368</i>	15	24.4; 0.161	1.99 (1.60, 2.46)	
Study location				
<i>China</i>	17	43.5; 0.016	1.77 (1.56, 2.00)	
<i>Taiwan</i>	3	34.8; 0.163	2.34 (1.39, 3.94)	
<i>India</i>	2	73.5; 0.010	1.30 (0.70, 2.42)	
<i>Other countries</i>	6	76.4; <0.001	1.49 (1.05, 2.13)	
Year study conducted				
<i>2000 onwards</i>	2	80.3; 0.006	1.85 (0.93, 3.67)	0.813
<i>Before 2000</i>	26	61.7; <0.001	1.70 (1.49, 1.95)	
Year study published				
<i>2000 onwards</i>	13	72.7; <0.001	1.70 (1.39, 2.08)	1.000
<i>Before 2000</i>	15	43.7; 0.020	1.70 (1.45, 2.01)	
Language of publication				
<i>Chinese</i>	8	0.0; 0.468	2.16 (1.81, 2.59)	0.006
<i>English</i>	33	62.7; <0.001	1.56 (1.35, 1.81)	
Newcastle-Ottawa score				
<i>>6⁺</i>	28	64.3; <0.001	1.58 (1.36, 1.85)	0.116
<i>≤6</i>	13	48.6; 0.025	1.97 (1.57, 2.47)	
Quality of exposure assessment[§]				
<i>1 star</i>	10	55.5; 0.017	1.91 (1.45, 2.53)	
<i>2 stars</i>	27	68.0; <0.001	1.64 (1.41, 1.91)	
<i>3 stars</i>	4	0.0; 0.754	1.78 (0.94, 3.37)	

*The total number of studies is 28 but as some studies have reported more than one sub-group type (mentioned above): hence the number of studies does not add up to 28 in all sub-group types.

[#]Studies reporting risk estimates from both genders combined.

[¶]The median sample size of all 28 studies

⁺The median Newcastle-Ottawa score

[§]The Newcastle-Ottawa score assigns a maximum of stars on the exposure assessment

Table 3: Sub-group analyses of the lung cancer risk according to fuel type

Sub-group types	Studies (n)	Heterogeneity (I ² (%); p-value)	OR (95% CI)	p-value
Exposure to biomass smoke*				
Gender				
Female only (F)	5	0.0; 0.434	1.98 (1.44, 2.73)	0.881 (F vs. M)
Male only (M)	1	N/A	1.78 (0.46, 6.93)	
Male and female	2	0.0; 0.404	1.20 (1.03, 1.39)	
Cancer histological sub-type				
Unspecified	6	15.8; 0.309	1.31 (1.09, 1.58)	0.942 (Adeno. vs. Squamous)
Adenocarcinoma	1	N/A	3.30 (1.36, 8.00)	
Squamous carcinoma	1	N/A	3.50 (0.95, 12.90)	
Adjustment for smoking				
Yes	3	44.1; 0.111	1.36 (0.99, 1.86)	0.183
Non-smokers only	4	0.0; 0.814	1.89 (1.31, 2.73)	
Study design				
Population-based	1	N/A	1.11 (0.44, 2.80)	0.327
Hospital-based	5	54.3; 0.053	1.84 (1.23, 2.76)	
Sample size				
>368	5	55.6; 0.061	1.45 (1.10, 1.91)	0.485
≤368	3	10.8; 0.339	1.88 (0.96, 3.70)	
Language of publication				
Chinese	0	N/A	N/A	
English	9	41.2; 0.092	1.50 (1.17, 1.94)	
Newcastle-Ottawa score				
>6	7	43.0; 0.104	1.42 (1.04, 1.94)	0.326
≤6	2	0.0; 0.860	1.82 (1.24, 2.68)	
Quality of exposure assessment				
1 star	1	N/A	1.90 (1.03, 3.50)	
2 stars	7	43.9; 0.098	1.42 (1.07, 1.87)	
3 stars	1	N/A	2.70 (0.82, 8.90)	
Exposure to coal smoke*				
Gender				
Female only (F)	10	45.5; 0.043	1.70 (1.40, 2.06)	0.490 (F vs. M)
Male only (M)	3	26.0; 0.259	1.54 (1.25, 1.88)	
Male and female	12	39.9; 0.068	2.19 (1.74, 2.76)	
Cancer histological sub-type				
Unspecified	16	38.1; 0.047	1.70 (1.51, 1.92)	0.324 (Adeno. vs. squamous)
Adenocarcinoma	4	0.0; 0.501	2.22 (1.60, 3.08)	
Squamous carcinoma	5	61.2; 0.035	3.81 (1.37, 10.58)	
Adjustment for smoking				
Yes	19	33.4; 0.054	1.82 (1.62, 2.06)	0.909
Non-smokers only	3	76.7; 0.014	1.73 (0.73, 4.10)	
Study design				
Population-based	13	60.4; 0.001	1.89 (1.59, 2.25)	0.730
Hospital-based	9	43.4; 0.008	1.82 (1.60, 2.06)	
Sample size				
>368	13	36.0; 0.087	2.04 (1.59, 2.61)	0.246
≤368	11	49.3; 0.019	1.72 (1.49, 2.00)	
Language of publication				
Chinese	8	0.0; 0.468	2.16 (1.81, 2.59)	0.022
English	20	42.3; 0.024	1.65 (1.43, 1.91)	
Newcastle-Ottawa score				
>6	20	42.3; 0.024	1.65 (1.43, 1.91)	0.022
≤6	8	0.0; 0.468	2.16(1.81, 2.59)	
Quality of exposure assessment				
1 star	7	0.0; 0.584	2.11 (1.75, 2.56)	
2 stars	18	53.8; 0.004	1.72 (1.47, 2.02)	
3 stars	3	0.0; 0.765	1.50 (0.71, 3.20)	

*The total number of biomass studies is 7 but as some studies have reported more than one sub-group types (mentioned above) the number of studies does not add up to 7 in all sub-group types; similarly, the total number of coal studies is 22.

Publication bias

Funnel plots suggested potential publication bias for both biomass (Supplementary Figure S7) and coal smoke (Supplementary Figure S8) studies. However, Egger's test showed substantial publication bias only in coal smoke studies (bias=1.04, $p=0.016$) (Supplementary Figure S10), which disappeared when two outlying studies [26, 29] were removed (bias=0.76, $p=0.093$). The pooled effect estimate (OR=1.64; 95% CI 1.45, 1.86) was slightly attenuated after excluding the two outliers.

Heterogeneity by meta-regression

Heterogeneity was initially explored by graphical display (Galbraith plot) (Supplementary Figure S11 for biomass and Figure S12 for coal), then by meta-regression to assess contributions by gender, histological sub-type, smoking, adjustment for ETS exposure, sample size, study location, year in which the study was carried out, year of publication, and language of publication. In studies of biomass smoke exposure significant but small heterogeneity was observed in gender (coefficient=-0.253, $p=0.025$), although there was a non-significant heterogeneity in lung cancer histology (coefficient=0.636, $p=0.057$). On the other hand, in studies of coal smoke exposure language of publication (coefficient=0.308, $p=0.032$) and histology (coefficient=0.273, $p=0.058$) had similar magnitude of heterogeneity, although the latter was not statistically significant. We did not find evidence of heterogeneity ($p=0.116$) between the studies of better quality (Newcastle-Ottawa score>6) and the poorer quality ones (≤ 6).

DISCUSSION

This meta-analysis included studies conducted in China, Taiwan, Japan, India, Mexico, Morocco, the United States and Canada, as well as a study carried out jointly in seven European countries (Czech Republic, Hungary, Poland, Romania,

Russia, Slovakia and United Kingdom). The pooled effect estimates that the risk of lung cancer among users of solid fuels is 70% (95% CI 50%, 94%) higher than non-users.

The magnitude of association between coal use and lung cancer (OR=1.82, 95% CI 1.60, 2.06) was greatest followed by biomass (predominantly wood, OR=1.50, 95% CI 1.17, 1.94) and mixed fuel (OR=1.13, 95% CI 0.52, 2.46), although the differences were not statistically significant. The higher risk of lung cancer in coal users was not surprising as combustion products obtained from in-home coal burning contain a range of Group 1 carcinogenic PAHs [4]. While there is sufficient evidence to suggest exposure to biomass smoke is a risk factor for chronic obstructive pulmonary disease (COPD) in adults [40] and acute respiratory infection in children [41], the International Agency for Research on Cancer (IARC) has classified combustion products from biomass (primarily wood) as probable human lung carcinogen (Group 2A), citing there was "limited evidence" in humans and experimental animals [4]. The pooled effect size obtained from studies using population- (OR=1.83, 95% CI 1.51, 2.21) based controls (carrying 56% weight) was similar to that using hospital- (OR=1.63, 95% CI 1.34, 1.97) based controls (39% weight). Among the 28 studies included, two population-based [16, 38] and two hospital-based studies [33, 36] did not find an increased risk of lung cancer. Of these, three being related to biomass use [16, 33, 38] and the other to coal use [36], supporting the IARC notion that the evidence of the carcinogenicity of biomass smoke is still not conclusive.

The association between lung cancer and solid fuel use persisted even after stratifying for gender, fuel types, smoking, and study location. The duration of exposure in most of the studies was not clearly defined and there was marked variability in reported exposure intensity across studies but the number of studies were too small to determine any dose-response relationship. Of the 28 studies

included in this meta-analysis, two studies scored the maximum of three stars on the Newcastle-Ottawa criteria for exposure whereas 18 studies scored two and eight studies scored only one. The studies with the highest quality in exposure assessment have lower effect sizes suggesting that misclassification and residual confounding might be operating, thereby inflating the risk estimate. Users of biomass often switch from one type of biomass to another. A detailed history on the type, duration and intensity of fuel use (such as average number of hours exposed) must be gathered in future studies to better estimate the risks from particular biomass fuels as combustion products from different types of biomass burning have variable toxicity [42].

Cigarette smoking has been widely accepted as the main contributory factor to lung cancer worldwide [43, 44]. We excluded two papers on the basis that smoking had not been allowed for in the risk estimates [45, 46], and all studies included in this review have either adjusted for smoking or studied a population of non-smokers. A recent meta-analysis [8] included effect estimates from Chinese studies that did not adjust for smoking. The extent of confounding is, however, difficult to predict. While it is accepted that self-reported smoking history is the best that can be achieved when considering life-long smoking details, objective measurement of smoking, such as salivary cotinine, is becoming more easily usable in field studies and provides information on current smoking, which may to a certain extent help reduce exposure misclassification. This is particularly the case for women from countries who hesitate to admit to smoking for the fear of marginalisation.

Although half of the studies included in the meta-analysis measured ETS but only a quarter of them presented data with adjusted ETS exposure. In studies that did, the pooled effect size (OR=1.47, 95% CI 1.13, 1.91) was smaller than (but not statistically significant, $p = 0.230$) those that did not (OR=1.74, 95% CI 1.60, 1.89). In

women the pooled effect estimate with adjusted ETS was significantly lower compared to non-adjusted suggesting the overall pooled effect estimate particularly in women might be lower than what is presented here. Only one study out of eight related to biomass smoke exposure adjusted for ETS and had effect size higher than the other that were not adjusted for ETS. Thus, ambiguity regarding the combined effect of smoking, combustion products of solid fuels and ETS exposure still prevails and future studies need to address this issue particularly in women from Asian sub-continent as they are highly likely to be exposed to ETS. There is evidence from occupational studies that smoking and some occupational exposures (e.g. asbestos, PAHs) have a multiplicative rather than an additive effect on lung cancer risk [47, 48] and it is therefore possible that such a potentiating effect may be seen with respect to smoke from solid fuel burning, especially that from coal.

Women in developing countries do most of the cooking and thus are more likely to be exposed to indoor air pollution than men. The pooled effect size shows that the risk of lung cancer is greater in women (OR=1.81, 95% CI 1.54, 2.12) compared to men (OR=1.16, 95% CI 0.79, 1.69), similar to that reported in a limited earlier meta-analysis [7] for women only (OR=1.83, 95% CI 0.62, 5.41). Many published meta-analyses reported data for men and women combined. In this study, the pooled effect size for both genders was 1.93 (95% CI 1.53, 2.44), smaller than that reported by Zhao *et al* [7] (OR=2.66, 96% CI 1.39, 5.07), probably because the latter was obtained from the coal using population in China. The pooled effect size in our study would have been reduced to 1.80 (95% CI 1.46, 2.22) if the two studies with effect sizes of 24.34 (95% CI 2.97, 199.48) [29] and 14.10 (95% CI 1.37, 145.61) [17] were excluded.

The pooled effect estimate in studies published in Chinese language (OR=2.16; 95% CI 1.81, 2.59) was significantly greater ($p=0.006$) than studies published in English.

When scrutinising the Chinese papers, we found a consistently large effect size. While the effect could be real, as Chinese papers reported focused on the coal-using Chinese population and that coal has been recognised by the IARC as a carcinogen, this raises a concern on the overall quality of the research published in Chinese journals.

Table 4 presents the main findings from previously published meta-analyses (including the current study). Over 60% of these (5/8) included studies either from China or the Chinese population only and examined only the effects of coal use. In contrast, the current meta-analysis presents the pooled results from various geographical regions, and has investigated the effects of biomass and coal exposures separately. In addition, we have specified in our inclusion criteria that only those studies that have adjusted for smoking or used non-smoking sample would be included, therefore minimising potential confounding from smoking.

Table 4: Pooled effect estimates from previous meta-analyses on solid fuel use and lung cancer

References	Publication year	No. of papers	Fuel type	Study location	Gender	Pooled effect estimate (95% CI)	Remarks
Current study	-	28	Biomass & coal	No limitation	F	1.81 (1.54, 2.12)	All studies included were adjusted for smoking or were from non-smokers
					M	1.16 (0.79, 1.69)	
					M & F	1.93 (1.53, 2.44)	
Hosgood <i>et al</i> [8]	2011	25	Coal	No limitation	F	2.50 (1.56, 4.00)	Covers all geographic areas; risk estimates of some studies were not adjusted for smoking
					M	2.76 (1.44, 5.27)	
					Non-smoking F	2.93 (1.40, 6.12)	
Hosgood <i>et al</i> [9]	2010	7	Wood & coal	Asia, Europe and N. America	M & F	2.15 (1.61, 2.89)	Large sample size; not systematic review
					F	1.60 (1.41, 1.82)	
					M	1.42 (1.27, 1.59)	
Hosgood <i>et al</i> [49]	Results not reported here as the meta-analysis focused on genotypes						
Zhao <i>et al</i> [7]	2006	27	Coal	China	F	1.83 (0.62, 5.41)	Studies based in China only
					M & F	2.66 (1.39, 5.07)	
Smith <i>et al</i> [50]	2004	12	Coal	China	F	1.94 (1.09, 3.47)	Not systematic review
					M	1.51 (0.97, 2.46)	
					M & F	2.55 (1.58, 4.10)	
Yao and Shi [5]	2003	5	Coal	China	M & F	3.20 (1.79, 5.71)	Chinese population only
Yao <i>et al</i> [51]	2002	Not specified	Coal	China	F	1.84 (0.94, 3.59)	Chinese population only
Zhang <i>et al</i> [6]	2001	4	Coal	China	Non-smoking F	1.42 (1.30, 1.55)	Studies based in China only

To our knowledge, this is the first assessment of whether solid fuel smoke is associated with specific histological sub-type. Cell type was reported in eight papers but the criteria for histological classification were not provided. The pooled effect size for squamous cell carcinoma (OR=3.58, 95% CI 1.58, 8.12) was greatest followed by adenocarcinoma (OR=2.33, 95% CI 1.72, 3.17) and unspecified type of lung cancer (OR=1.57, 95% CI 1.38, 1.80). Squamous cell lung cancer is more commonly associated with cigarette smoking [52] although reported series of lung cancers have recently shown an increase in the proportion of adenocarcinoma which cannot simply be attributed to changes in classification/grading [53]. If cell type reflects different carcinogenic properties of different exposures then future studies studying the risk of lung cancer from solid fuel would benefit by classifying the types of lung cancer by fuel type.

Most of the studies included in this meta-analysis are from China where coal is the main fuel. The pooled effect size in Taiwan (3 studies, OR=2.34, 95% CI 1.39, 3.94) is greater than that in China (17 studies, OR=1.77, 95% CI 1.57, 2.00). None of the studies included from China and Taiwan have looked at the association between coal type and lung cancer risk. Nevertheless, evidence from a community with high lung cancer mortality in China suggested that bituminous or “smoky” coal, with a high volatiles content (23.1%) was more carcinogenic compared to smokeless coal which contains relatively high sulphur (1.9%) but low volatiles (13.8%) [54]. Further investigation [54, 55] concluded that compared to wood and smokeless coal, smoky coal contains more methylated PAH compounds, nitrogen heterocyclic compounds and dibenzo[a,l]pyrene, a potent carcinogen with the highest mutagenic activity in mice.

Most studies did not measure exposure quantitatively. Understanding the shape of the dose-response curve has been a challenge for a range of outcomes arising from

biomass smoke exposure (e.g. COPD [40], and acute respiratory infections in children [41]), but is crucial in determining to what extent exposures would need to be reduced in order to confer a significant health benefit. However, measuring current exposures may only partially reflect historical exposures, even though in many areas where solid fuel is burnt, practice and therefore exposures have likely remained similar for decades. Nevertheless, if formal quantification of exposures can be undertaken in future studies this will provide relevant information to address this issue.

Our results suggested an element of publication bias which could be due to fewer positive studies being rejected and more positive studies some with flawed methodology being accepted. The meta-regression showed that there was significant heterogeneity among studies reporting different types of lung cancer.

CONCLUSION

Our meta-analysis suggested that coal is highly associated with lung cancer compared to other types of biomass. The risk was greater in women and in China which could be because Chinese women used coal. Future studies need to look at objective measurements of smoking and also the carcinogenic potential of different coal subtypes to explain some of the variability seen in the risk estimates.

Competing interests None

REFERENCES

1. Parkin DM, Bray F, Ferlay J, Pisani P. Global cancer statistics, 2002. *CA Cancer J Clin* 2005; 55: 74-108.
2. Sun S, Schiller JH, Gazdar AF. Lung cancer in never smokers--a different disease. *Nat Rev Cancer* 2007; 7: 778-790.
3. International Energy Agency. World Energy Outlook. Chapter 13: Energy and Poverty. 2nd ed, 2002; pp. 365-395.
4. IARC. IARC Monographs on the evaluation of carcinogenic risks to humans, Vol. 95. Household use of solid fuels and high-temperature frying. IARC, Lyon, 2010; p. 392.
5. Yao HY, Shi LY. Meta-analysis of the risk factors on lung cancer in Chinese people. *Zhonghua Liu Xing Bing Xue Za Zhi* 2003; 24: 45-49.
6. Zhang Y, Chen K, Zhang HL, Zhu YM. Meta-analysis of risk factors on lung cancer in non-smoking Chinese female. *Zhonghua Liu Xing Bing Xue Za Zhi* 2001; 22: 119-121.
7. Zhao Y, Wang S, Aunan K, Seip HM, Hao J. Air pollution and lung cancer risks in China--a meta-analysis. *Sci Total Environ* 2006; 366: 500-513.
8. Hosgood HD, 3rd, Wei H, Sapkota A, Choudhury I, Bruce N, Smith KR, Rothman N, Lan Q. Household coal use and lung cancer: systematic review and meta-analysis of case-control studies, with an emphasis on geographic variation. *Int J Epidemiol* 2011.
9. Hosgood HD, 3rd, Boffetta P, Greenland S, Lee YC, McLaughlin J, Seow A, Duell EJ, Andrew AS, Zaridze D, Szeszenia-Dabrowska N, Rudnai P, Lissowska J, Fabianova E, Mates D, Bencko V, Foretova L, Janout V, Morgenstern H, Rothman N, Hung RJ, Brennan P, Lan Q. In-home coal and wood use and lung cancer risk: a pooled analysis of the International Lung Cancer Consortium. *Environ Health Perspect* 2010; 118: 1743-1747.
10. Wells GA, Shea B, O'Connell D, Peterson P, Welch V, Losos M, Tugwell P. The Newcastle-Ottawa Scale (NOS) for assessing the quality of nonrandomised studies in meta-analyses. 2011 [cited; Available from: http://www.ohri.ca/programs/clinical_epidemiology/oxford.asp
11. Borenstein M, Hedges LV, Higgins JPT, Rothstein HR. Introduction to meta-analysis. Wiley, Oxford, 2009.
12. Huang C, Zhang X, Qiao Z, Guan L, Peng S, Liu J, Xie R, Zheng L. A case-control study of dietary factors in patients with lung cancer. *Biomed Environ Sci* 1992; 5: 257-265.
13. Lan Q, He X. Molecular epidemiological studies on the relationship between indoor coal burning and lung cancer in Xuan Wei, China. *Toxicology* 2004; 198: 301-305.
14. Lan Q, Chapman RS, Schreinemachers DM, Tian L, He X. Household stove improvement and risk of lung cancer in Xuanwei, China. *J Natl Cancer Inst* 2002; 94: 826-835.
15. Liu Q, Sasco AJ, Riboli E, Hu MX. Indoor air pollution and lung cancer in Guangzhou, People's Republic of China. *Am J Epidemiol* 1993; 137: 145-154.
16. Liu ZY, He XZ, Chapman RS. Smoking and other risk factors for lung cancer in Xuanwei, China. *Int J Epidemiol* 1991; 20: 26-31.
17. Luo RX, Wu B, Yi YN, Huang ZW, Lin RT. Indoor burning coal air pollution and lung cancer--a case-control study in Fuzhou, China. *Lung Cancer* 1996; 14 Suppl 1: S113-119.

18. Sun XW, Dai XD, Xu GJ, Yu SF. Heating fuels and respiratory diseases in the risks of female lung cancer. *Zhonghua Zhong Liu Za Zhi* 1991; 13: 413-415.
19. Sun XW, Dai XD, Shi YB, Lin YG. A case-control study on the relationship among indoor air pollution, depression and oncogenesis of lung cancer. *Chin J Lung Cancer* 2002; 5: 101-103.
20. Zhong L, Goldberg MS, Gao YT, Jin F. Lung cancer and indoor air pollution arising from Chinese-style cooking among nonsmoking women living in Shanghai, China. *Epidemiology* 1999; 10: 488-494.
21. Wu-Williams AH, Dai XD, Blot W, Xu ZY, Sun XW, Xiao HP, Stone BJ, Yu SF, Feng YP, Ershow AG, et al. Lung cancer among women in north-east China. *Br J Cancer* 1990; 62: 982-987.
22. Xu ZY, Blot WJ, Xiao HP, Wu A, Feng YP, Stone BJ, Sun J, Ershow AG, Henderson BE, Fraumeni JF, Jr. Smoking, air pollution, and the high rates of lung cancer in Shenyang, China. *J Natl Cancer Inst* 1989; 81: 1800-1806.
23. Galeone C, Pelucchi C, La Vecchia C, Negri E, Bosetti C, Hu J. Indoor air pollution from solid fuel use, chronic lung diseases and lung cancer in Harbin, Northeast China. *Eur J Cancer Prev* 2008; 17: 473-478.
24. Lin CY, Sun XW, Shi YB, Dai XD, Zhang YH, Wang Y. Indoor coal smoke pollution and female adenocarcinoma. *Bulletin of Chinese cancer* 1996; 5: 21-22.
25. Hao LY, Liu JZ. A study on the risk factors for lung cancer mortality among farmers in Shunyi County, Beijing. *Chin J Public Health* 1998; 14: 457-458.
26. Lu J, Zhu J, Wang Y, Zeng WM, Zeng BH, Wu ZL, Shi LY, Liao YD. The risk factors of human lung cancer with 445 paired cases and controls. *Practical Preventive Medicine* 2003; 10: 275-279.
27. Liang GH, Pu YP, Yin LH. Case-control study on environmental risk factors of lung cancer in Nanjing. *Chin J Public Health* 2004; 20: 260-261.
28. Huang ZB. A study on the risk factors and population attributable risk for primary lung cancer. *J Guangxi Med Univ* 1999; 16: 447-450.
29. Ger LP, Hsu WL, Chen KT, Chen CJ. Risk factors of lung cancer by histological category in Taiwan. *Anticancer Res* 1993; 13: 1491-1500.
30. Ko YC, Lee CH, Chen MJ, Huang CC, Chang WY, Lin HJ, Wang HZ, Chang PY. Risk factors for primary lung cancer among non-smoking women in Taiwan. *Int J Epidemiol* 1997; 26: 24-31.
31. Lee CH, Ko YC, Cheng LS, Lin YC, Lin HJ, Huang MS, Huang JJ, Kao EL, Wang HZ. The heterogeneity in risk factors of lung cancer and the difference of histologic distribution between genders in Taiwan. *Cancer Causes Control* 2001; 12: 289-300.
32. Sapkota A, Gajalakshmi V, Jetly DH, Roychowdhury S, Dikshit RP, Brennan P, Hashibe M, Boffetta P. Indoor air pollution from solid fuels and risk of hypopharyngeal/laryngeal and lung cancers: a multicentric case-control study from India. *Int J Epidemiol* 2008; 37: 321-328.
33. Gupta D, Boffetta P, Gaborieau V, Jindal SK. Risk factors of lung cancer in Chandigarh, India. *Indian J Med Res* 2001; 113: 142-150.
34. Sobue T. Association of indoor air pollution and lifestyle with lung cancer in Osaka, Japan. *Int J Epidemiol* 1990; 19 Suppl 1: S62-66.

35. Hernandez-Garduno E, Brauer M, Perez-Neria J, Vedal S. Wood smoke exposure and lung adenocarcinoma in non-smoking Mexican women. *Int J Tuberc Lung Dis* 2004; 8: 377-383.
36. Sasco AJ, Merrill RM, Dari I, Benhaim-Luzon V, Carriot F, Cann CI, Bartal M. A case-control study of lung cancer in Casablanca, Morocco. *Cancer Causes Control* 2002; 13: 609-616.
37. Wu AH, Henderson BE, Pike MC, Yu MC. Smoking and other risk factors for lung cancer in women. *J Natl Cancer Inst* 1985; 74: 747-751.
38. Ramanakumar AV, Parent ME, Siemiatycki J. Risk of lung cancer from residential heating and cooking fuels in Montreal, Canada. *Am J Epidemiol* 2007; 165: 634-642.
39. Lissowska J, Bardin-Mikolajczak A, Fletcher T, Zaridze D, Szeszenia-Dabrowska N, Rudnai P, Fabianova E, Cassidy A, Mates D, Holcatova I, Vitova V, Janout V, Mannetje A, Brennan P, Boffetta P. Lung cancer and indoor pollution from heating and cooking with solid fuels: the IARC international multicentre case-control study in Eastern/Central Europe and the United Kingdom. *Am J Epidemiol* 2005; 162: 326-333.
40. Kurmi OP, Semple S, Simkhada P, Smith WC, Ayres JG. COPD and chronic bronchitis risk of indoor air pollution from solid fuel: a systematic review and meta-analysis. *Thorax* 2010; 65: 221-228.
41. Dherani M, Pope D, Mascarenhas M, Smith KR, Weber M, Bruce N. Indoor air pollution from unprocessed solid fuel use and pneumonia risk in children aged under five years: a systematic review and meta-analysis. *Bull World Health Organ* 2008; 86: 390-398C.
42. Semple S, Devakumar D, Fullerton DG, Thorne PS, Metwali N, Costello A, Gordon SB, Manandhar DS, Ayres JG. Airborne endotoxin concentrations in homes burning biomass fuel. *Environ Health Perspect* 2010; 118: 988-991.
43. Peto R, Darby S, Deo H, Silcocks P, Whitley E, Doll R. Smoking, smoking cessation, and lung cancer in the UK since 1950: combination of national statistics with two case-control studies. *BMJ* 2000; 321: 323-329.
44. Hirayama T. Non-smoking wives of heavy smokers have a higher risk of lung cancer: a study from Japan. *Br Med J (Clin Res Ed)* 1981; 282: 183-185.
45. Lei YX, Cai WC, Chen YZ, Du YX. Some lifestyle factors in human lung cancer: a case-control study of 792 lung cancer cases. *Lung Cancer* 1996; 14 Suppl 1: S121-136.
46. Li J, Wang GX, Shen XB, Zhang XJ, Shen QJ, Zhang XD, Chang W, Liu HT, Song LP, Yang B, . The analysis of risk factors of primary lung adenocarcinoma patients in Nanjing City. *Modern Railway Journal* 1993; 21: 329-331.
47. Whitwell F, Newhouse ML, Bennett DR. A study of the histological cell types of lung cancer in workers suffering from asbestosis in the United Kingdom. *Br J Ind Med* 1974; 31: 298-303.
48. Tang DL, Rundle A, Warburton D, Santella RM, Tsai WY, Chiamprasert S, Hsu YZ, Perera FP. Associations between both genetic and environmental biomarkers and lung cancer: evidence of a greater risk of lung cancer in women smokers. *Carcinogenesis* 1998; 19: 1949-1953.

49. Hosgood HD, 3rd, Berndt SI, Lan Q. GST genotypes and lung cancer susceptibility in Asian populations with indoor air pollution exposures: a meta-analysis. *Mutat Res* 2007; 636: 134-143.
50. Smith KR, Mehta S, Feuz M. Indoor smoke from household solid fuels. *In: Ezzati M, Rodgers AD, Lopez AD, Murray CJL, eds. Comparative quantification of health risks: global and regional burden of disease due to selected major risk factors. World Health Organisation, Geneva, 2004.*
51. Yao HY, Lü K, Shi LY. An analysis of risk factors in female lung cancer. *Bulletin of Chinese cancer* 2002; 11: 508-510.
52. Nakachi K, Imai K, Hayashi S, Watanabe J, Kawajiri K. Genetic susceptibility to squamous cell carcinoma of the lung in relation to cigarette smoking dose. *Cancer Res* 1991; 51: 5177-5180.
53. Kusano C, Gotoda T, Khor CJ, Katai H, Kato H, Taniguchi H, Shimoda T. Changing trends in the proportion of adenocarcinoma of the esophagogastric junction in a large tertiary referral center in Japan. *J Gastroenterol Hepatol* 2008; 23: 1662-1665.
54. Mumford JL, He XZ, Chapman RS, Cao SR, Harris DB, Li XM, Xian YL, Jiang WZ, Xu CW, Chuang JC, et al. Lung cancer and indoor air pollution in Xuan Wei, China. *Science* 1987; 235: 217-220.
55. Mumford JL, Li X, Hu F, Lu XB, Chuang JC. Human exposure and dosimetry of polycyclic aromatic hydrocarbons in urine from Xuan Wei, China with high lung cancer mortality associated with exposure to unvented coal smoke. *Carcinogenesis* 1995; 16: 3031-3036.

Figure 1: Flow chart showing studies related to lung cancer and exposure to solid fuel

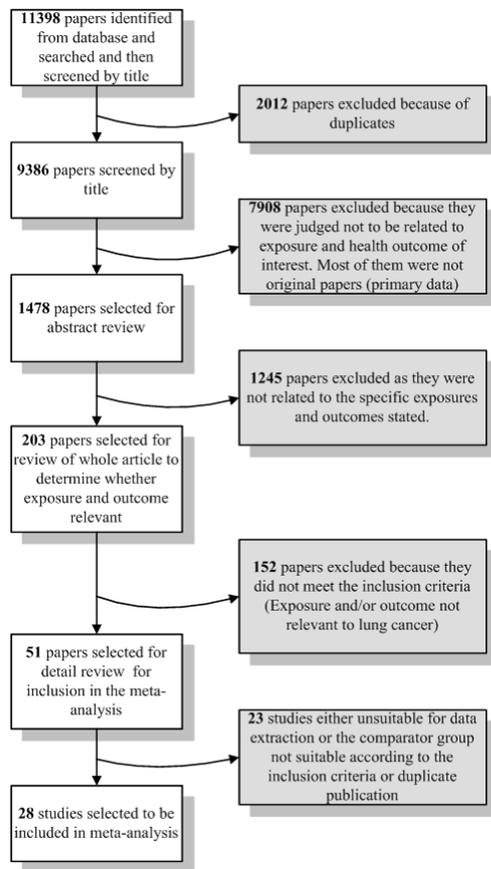


Figure 2: Forest plot of studies reporting lung cancer associated with exposure to solid fuels (stratified by types of fuel used)

