

# Association of active and passive smoking with risk of breast cancer among postmenopausal women: a prospective cohort study

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## ABSTRACT

**Objective** To examine the association between smoking and risk of invasive breast cancer using quantitative measures of lifetime passive and active smoking exposure among postmenopausal women.

**Design** Prospective cohort study.

**Setting** 40 clinical centres in the United States.

**Participants** 79 990 women aged 50–79 enrolled in the Women's Health Initiative Observational Study during 1993–8.

**Main outcome measures** Self reported active and passive smoking, pathologically confirmed invasive breast cancer.

**Results** In total, 3520 incident cases of invasive breast cancer were identified during an average of 10.3 years of follow-up. Compared with women who had never smoked, breast cancer risk was elevated by 9% among former smokers (hazard ratio 1.09 (95% CI 1.02 to 1.17)) and by 16% among current smokers (hazard ratio 1.16 (1.00 to 1.34)). Significantly higher breast cancer risk was observed in active smokers with high intensity and duration of smoking, as well as with initiation of smoking in the teenage years. The highest breast cancer risk was found among women who had smoked for ≥50 years or more (hazard ratio 1.35 (1.03 to 1.77)) compared with all lifetime non-smokers, hazard ratio 1.45 (1.06 to 1.98) compared with lifetime non-smokers with no exposure to passive smoking). An increased risk of breast cancer persisted for up to 20 years after smoking cessation. Among women who had never smoked, after adjustment for potential confounders, those with the most extensive exposure to passive smoking (≥10 years' exposure in childhood, ≥20 years' exposure as an adult at home, and ≥10 years' exposure as an adult at work) had a 32% excess risk of breast cancer compared with those who had never been exposed to passive smoking (hazard ratio 1.32 (1.04 to 1.67)). However, there was no significant association in the other groups with lower exposure and no clear dose response to cumulative passive smoking exposure.

**Conclusions** Active smoking was associated with an increase in breast cancer risk among postmenopausal women. There was also a suggestion of an association between passive smoking and increased risk of breast cancer.

## INTRODUCTION

Experimental studies have shown that compounds found in tobacco smoke, such as polycyclic hydrocarbons, aromatic amines, and N-nitrosamines, may induce mammary tumours.<sup>1</sup> The detection of smoking-specific DNA adducts and mutations in the p53 gene in the breast tissue of smokers supports the biological plausibility of a positive association between cigarette smoking and breast cancer.<sup>2–6</sup> Despite this mechanistic evidence, systematic reviews of epidemiological studies published as of 2002 concluded that there was no overall association between active smoking and breast cancer risk, and attributed conflicting results of individual studies in part to the confounding effects of alcohol.<sup>1 7–10</sup> However, recent reappraisals of evidence<sup>11 12</sup> from recent cohort studies<sup>13–19</sup> have suggested an increased risk of breast cancer that is independent of the effects of alcohol among women who smoked cigarettes for a long period of time or who started smoking at a young age.

Similarly, studies of passive smoking suggest an increased risk of breast cancer, particularly case-control studies that have conducted a more thorough assessment of exposure.<sup>20–24</sup> These reports found stronger evidence of an association of passive smoking and breast cancer in younger, primarily premenopausal women, whereas the evidence in older or postmenopausal women was inconclusive. Although cohort studies are generally considered less subject to bias than case-control studies, assessment of passive smoking has generally been less comprehensive in large cohort studies than in case-control studies.<sup>16 18 25–29</sup> Despite limited evaluations of exposure, which may lead to exposure misclassification and bias the relative risk towards the null, several of these studies found

associations of passive smoking with breast cancer in subgroups of women who reported being more heavily exposed.<sup>27–29</sup> Only one published cohort study includes full information on three major aspects of lifetime exposure to passive smoking (childhood, adult residential, and adult occupational), and this study found increased breast cancer risk in a heavily exposed subgroup.<sup>30</sup> Thus, prospective studies with more detailed assessment of passive smoking are needed to elucidate the association of passive smoking and breast cancer risk, especially in postmenopausal women.

In the Women's Health Initiative Observational Study, detailed information regarding active and passive smoking exposure was collected, including questions on passive exposure to cigarette smoke in childhood and adult exposure both in the home and at work. In the present study, we used data from the Women's Health Initiative that included a large number of breast cancer cases and detailed exposure information to address the relation between active and passive smoking and the risk of breast cancer in postmenopausal women. Further, given that breast cancer is a heterogeneous disease with subtypes that may vary in their aetiologies, we explored whether the effects of active or passive smoking differed by disease subgroups defined by hormone receptor status or histology.

## METHODS

### Women's Health Initiative

The Women's Health Initiative Observational Study was designed to address major causes of morbidity and mortality in postmenopausal women.<sup>31</sup> A total of 93 676 women aged 50–79 were recruited at 40 clinical centres throughout the United States from 1993 to 1998.<sup>32,33</sup> Women were excluded if they did not plan to reside in the area for at least three years, if they had conditions predictive of survival less than three years, or had complicating conditions such as alcoholism, drug dependency, or dementia. The study was overseen by ethics committees at all 40 clinical centres and at the coordinating centre, as well as by a data and safety monitoring board. All participants in the study gave informed consent and were followed prospectively. Of the original cohort of 93 676, we excluded 12 075 with a history of cancer (except non-melanoma skin cancer) at baseline, 443 who had no follow-up time, and 1168 with missing values of smoking status. This yielded a sample of 79 990 women for further analysis.

### Measurement of exposures and confounders

All information on exposures and confounders used in this analysis was collected at baseline. Information on active smoking included smoking status (never, former, or current), and women who were current or former smokers were also asked the age at which they started smoking, the number of cigarettes smoked per day, and the duration of smoking in years. Among former smokers, age at quitting smoking was also collected. Pack years of smoking were calculated by

multiplying the total years of smoking by the number of cigarettes smoked a day divided by 20.

Questions on exposure to passive smoking related to each of the following: childhood (<18 years old), adult home, adult work, current home, and current work (yes or no). Women who reported any of these exposures were also asked the number of years of exposure in childhood, as an adult at home, and as an adult at work (predefined categories <1, 1–4, 5–9, 10–18 for childhood exposure; <1, 1–4, 5–9, 10–19, 20–29, 30–39, ≥40 for adult exposure at home or work).

The potential confounders used in multivariable analyses included age at enrolment (<55, 55–59, 60–64, 65–69, 70–74, ≥75 years), ethnicity (American Indian or Alaska native, Asian or Pacific Islander, black or African-American, Hispanic or Latino, non-Hispanic white, and other), education (high school or less, some college or technical training, college or some post-college, and master's degree or higher), body mass index (<18.5, 18.5–24.9, 25.0–29.9, 30.0–34.9, 35.0–39.9, ≥40), physical activity (metabolic equivalent tasks per week <5, 5–<10, 10–<20, 20–<30, ≥30), alcohol intake (non-drinker, former drinker, <1 drink/month, 1 drink/month–<1 drink/week, 1–<7 drinks/week, ≥7 drinks/week), parity (never pregnant, never had term pregnancy, 1, 2, 3, 4, ≥5), family history of breast cancer (yes/no), history of hormone therapy use (none, oestrogen alone, oestrogen and progestin, mixed), age at menarche (≤12, 12–13, 14–15, ≥16 years), and age of first live birth (never had term pregnancy, <20, 20–29, ≥30 years).

### Follow-up and ascertainment of cases

Initial reports of cancer were ascertained by annual self administered questionnaires, and all reports of breast cancer were confirmed by review of medical records, including pathology reports (if a biopsy or resection was done). The breast cancer cases were then coded by an experienced coder in accordance with the coding guidelines from Surveillance Epidemiology and End Results.<sup>34</sup> Primary tumour site and histology were coded in accordance with the ICD-O-2 (international classification of diseases for oncology, second edition). The completion rate of annual questionnaires was 93%–96%.

In this study, we included only invasive breast cancer cases; cases of carcinoma in situ were not included. By 14 August 2009, with an average 10.3 years of follow-up, 3520 incident cases of invasive breast cancer had been identified. Among 41 022 women who had never smoked, 1692 cases were identified, and these cases were used for the passive smoking analysis.

### Statistical analysis

Analysis of the association of passive smoking with breast cancer risk was limited to the 41 022 women who had never smoked. Before analysing the data, we developed a method to combine data on all passive smoking exposures, including exposure during childhood, adult home exposure, and work exposure. We constructed a new variable using the mutually

**Table 1** | Baseline characteristics of 79 990 postmenopausal women by diagnosis of invasive breast cancer (values are numbers (percentages) of women unless stated otherwise)

Variable	Cases	Non-cases	P value of difference*
Total No of women	3520	76 470	
Mean (SD) age at baseline (years)	63.7 (7.1)	63.4 (7.3)	0.006
Mean (SD) body mass index (kg/m <sup>2</sup> )	27.4 (5.7)	27.2 (5.8)	0.07
Mean (SD) physical activity (METs/week)	13.7 (13.6)	13.8 (14.4)	0.8
Non-Hispanic white ethnicity	3126 (88.8)	63 356 (82.9)	<0.0001
Higher educational status†	1674 (47.6)	31 827 (41.6)	<0.0001
Age at menarche <12 years	439 (12.5)	8792 (11.5)	0.2
Hormone therapy use:			
Oestrogen alone	986 (28.0)	23 714 (31.0)	
Oestrogen plus progestin	1082 (30.7)	17 842 (23.3)	<0.0001
Mixed use	262 (7.4)	4699 (6.1)	
Nulliparous	496 (14.1)	9484 (12.4)	<0.0001
Age at first live birth ≥30 years	337 (9.6)	5695 (7.5)	<0.0001
Alcohol intake ≥7 drinks/week	552 (15.7)	9418 (12.4)	<0.0001
Family history of breast cancer	812 (23.1)	13 379 (17.5)	<0.0001
Smoking status:			
Never smoked:	1692 (48.1)	39 330 (51.4)	
Exposure to passive smoking:			
None	145 (8.6)	3609 (9.2)	
Childhood only	110 (6.5)	2241 (5.7)	
Adult at home only	88 (5.2)	2436 (6.2)	
Adult at work only	199 (11.8)	4800 (12.2)	<0.0001
More than one type	1118 (66.1)	25 132 (63.9)	
Unknown	32 (1.9)	1112 (2.8)	
Former smokers	1619 (46.0)	32 371 (42.3)	
Current smokers	209 (5.9)	4769 (6.2)	

SD=standard deviation. METs=metabolic equivalent tasks.

\* $\chi^2$  test between cases and non-cases. Significant for age, race, education, hormone therapy use, parity, age of first live birth, alcohol intake, and family history of breast cancer (all P values <0.05).

†College graduate or above.

exclusive categories of no exposure, exposure in childhood only, adult exposure at home only, adult exposure at work only, exposure to any two combinations, and exposure to all three. Based on the distribution of responses, we dichotomised childhood exposure as <10 or ≥10 years, adult exposure at home as <20 or ≥20 years, and adult exposure at work as <10 or ≥10 years. Within the triple exposure category, we created an additional “extensive exposure” group, defined as exposure in childhood ≥10 years, adult home exposure ≥20 years, and adult work exposure ≥10 years. Of the 41 022 never smokers, 2956 (7.2%) had “extensive exposure” to passive smoking.

In addition to the above variables defined a priori, we derived some categories after initial analysis in order to estimate the cumulative dose of exposure to passive smoking. We assigned mean values for each exposure category in the original questionnaire response and simply summed up all three types of exposure years. We assessed exposure for ≥10 years in childhood plus summed mean years of adult exposure, weighting adult home exposure three times more heavily, based on the estimate of time spent at home compared with at work.<sup>35</sup> We also applied various multipliers to workplace exposure ranging from 0.8 to 1.5

based on the work of Hammond et al.<sup>35,36</sup> Finally, we examined additional categories of dual and triple exposure based on our original dichotomous variables, including the “extensive exposure” group defined a priori.

We estimated hazard ratios for active smoking compared with two reference groups. In the primary analysis, the reference group was lifetime non-smokers; in a secondary analysis we used the much smaller reference group of lifetime non-smokers with no history of passive smoking.

Cox proportional hazards regression models were used to estimate hazard ratios and 95% confidence intervals, adjusting for the potential confounders described above in measurement of exposure and confounders. Participants without complete data for all covariates in a given multivariable model were excluded from that analysis. Follow-up time for each woman was accrued from enrolment to the date of diagnosis of invasive breast cancer, death, loss to follow-up (including non-participation in an extension of the study starting in 2005), or the administrative censoring date (14 August 2009), whichever occurred first. Tests for trend were performed by using the ordered category, including the reference, as a continuous variable in the proportional hazard model. The proportionality assumption was satisfied for all exposure variables of interest and potential confounding variables based on graphs of scaled Schoenfeld residuals.<sup>37</sup> All statistical analyses were conducted using SAS (Version 9.0, SAS Institute, Cary, NC).

## RESULTS

Table 1 shows the baseline characteristics of the study subjects by disease status. Women who developed invasive breast cancer were slightly older than women who did not, but their body mass index and physical activity were similar. Compared with women who remained free of breast cancer, those who developed the disease were significantly more likely to be non-Hispanic white, more highly educated, nulliparous, older at first live birth, and to have a history of taking oestrogen plus progesterone hormone therapy, heavier alcohol intake, and a family history of breast cancer (all P values <0.05). Women who developed breast cancer were also less likely to have never smoked than those who did not develop cancer (48.1% v 51.4%). Among lifetime non-smokers, 88.1% were exposed to passive smoking, and most women had multiple types of passive smoking exposure. Childhood exposure only was slightly higher among the non-smokers who developed breast cancer than those who did not, as was multiple types of passive smoking exposure (table 1).

Among women who had never smoked, we did not observe a significant association between breast cancer risk and passive smoking except for the highest level of exposure (childhood exposure ≥10 years, adult home exposure ≥20 years, and adult work exposure ≥10 years) (table 2). We also did not observe a significant dose-response trend for any method of estimating

**Table 2** | Cases of incident invasive breast cancer and related hazard ratios associated with exposure to passive smoking among 41 022 postmenopausal women who had never smoked

Exposure to passive smoking	No of cases	Hazard ratio (95% CI)	
		Age adjusted	Multivariable adjusted*
None	145	1.00 (reference)	1.00 (reference)
Any:	1515	1.08 (0.91 to 1.28)	1.09 (0.92 to 1.29)
Childhood only	110	1.22 (0.95 to 1.56)	1.19 (0.93 to 1.53)
Adult at home only	88	0.89 (0.69 to 1.16)	0.91 (0.70 to 1.19)
Adult at work only	199	1.02 (0.83 to 1.27)	1.01 (0.82 to 1.26)
Childhood + adult at home	170	1.02 (0.81 to 1.27)	1.04 (0.83 to 1.30)
Childhood + adult at work	191	1.18 (0.95 to 1.47)	1.17 (0.94 to 1.45)
Adult at home + at work	236	1.12 (0.91 to 1.38)	1.15 (0.93 to 1.41)
Childhood + adult at home + at work	521	1.09 (0.90 to 1.31)	1.11 (0.92 to 1.34)
Cumulative exposure categories:			
No childhood + any adult	524	1.04 (0.86 to 1.24)	1.05 (0.87 to 1.26)
Childhood <10 years + any adult	154	1.10 (0.88 to 1.39)	1.13 (0.90 to 1.42)
Childhood ≥10 years + adult at home <20 years + adult at work <10 years	421	1.14 (0.95 to 1.38)	1.13 (0.94 to 1.37)
Childhood ≥10 years + adult at home <20 years + adult at work ≥10 years	151	0.99 (0.79 to 1.24)	0.98 (0.77 to 1.23)
Childhood ≥10 years + adult at home ≥20 years + adult at work <10 years	124	0.99 (0.78 to 1.25)	1.05 (0.82 to 1.33)
Childhood ≥10 years + adult at home ≥20 years + adult at work ≥10 years†	146	1.27 (1.01 to 1.60)	1.32 (1.04 to 1.67)
P value for trend§		0.16	0.10

\*Variables used in multivariable models included age at enrolment, ethnicity, education, body mass index, physical activity, alcohol intake, parity, family history of breast cancer, hormone therapy use, age at menarche, and age of first live birth.  
†The "extensive exposure" variable defined a priori.  
§Including the reference group (no exposure) and the 6 rows of cumulative exposure categories.

cumulative exposure, with all the estimates being qualitatively similar to the categorical results shown in table 2. In the women who had never smoked with the most extensive exposure to passive smoking, we observed a 32% (95% confidence interval 4% to 67%) excess risk of breast cancer (table 2).

We assessed the risk of breast cancer associated with different measures of active smoking by including or excluding passive smokers in the reference category who had never smoked (table 3). In the primary analysis, with all lifetime non-smokers as the reference group, breast cancer risk was elevated by 9% (2% to 17%) among former smokers and by 16% (0% to 34%,  $P=0.05$ ) among current smokers. Compared with never smoking, the risk of breast cancer was positively associated with smoking intensity, smoking duration, and pack years of cigarette smoking, and inversely associated with age at smoking initiation. Compared with never smokers who had at least one full term pregnancy, women who began smoking before their first full term pregnancy had a 21% (11% to 33%) elevated risk of breast cancer. All trend tests were statistically significant. The highest breast cancer risk was found among women who had smoked for ≥50 years (hazard ratio 1.35 (1.03 to 1.77)). Among former smokers, the years since quitting smoking was significantly inversely associated with breast cancer risk. An increased risk of breast cancer persisted for up to 20 years after smoking cessation.

Since extensive exposure to passive smoking was associated with increased breast cancer risk, which could result in underestimating the risks of active smoking, we conducted a secondary analysis using lifetime non-smokers with no history of any exposure to

passive smoke as the reference group. In general, the point estimates of active smoking risks in this secondary analysis were higher than those based on inclusion of passive smokers in the reference group but were less precise because of the much smaller size of the reference group. The hazard ratio for breast cancer among current smokers was significantly elevated ( $P<0.05$ ) regardless of the reference group used, as was the risk among women who started smoking before their first term pregnancy, smoked for ≥50 years, or had ≥50 pack years of smoking (table 3).

Finally, we assessed the association between smoking exposure and the risk of breast cancer for different types of breast cancer (table 4). There was no significant association of extensive passive smoking with any breast cancer type, but statistical power was limited by the small sample sizes. There was a non-significant trend towards a stronger association of current smoking with lobular cancer than with ductal cancer ( $P$  for interaction=0.13). Although there was a significant association of former and current smoking with tumours that were hormone receptor positive (for both oestrogen and progesterone, the dominant subtype), statistical power was probably too limited to assess whether this was true with other hormone receptor subtypes ( $P$  for interaction>0.2) (table 4). When we examined the associations of the different variables of active smoking (including age of smoking initiation, intensity, duration, and pack years of smoking) on the risk of breast cancer for these subgroups, we found similar patterns to that of breast cancer overall for dominant subtypes (ductal cancer and hormone receptor positive) and no clear pattern for other subtypes (data not shown).

**Table 3** | Cases of incident invasive breast cancer and related hazard ratios associated with smoking status among 79 990 postmenopausal women

Smoking status	No of cases	Hazard ratio (95% CI)	
		Multivariable adjusted*	Multivariable adjusted†
Never smoked:	1692	1.00 (reference)	—
With no exposure to passive smoking	145	—	1.00 (reference)
With exposure to passive smoking	1515	—	1.08 (0.91 to 1.28)
Former smokers	1619	1.09 (1.02 to 1.17)	1.16 (0.98 to 1.38)
Current smokers	209	1.16 (1.00 to 1.34)	1.24 (1.00 to 1.54)
Age at starting smoking (years):			
<15	112	1.12 (0.92 to 1.36)	1.19 (0.93 to 1.53)
15–19	940	1.13 (1.04 to 1.23)	1.21 (1.01 to 1.44)
20–24	581	1.09 (0.99 to 1.20)	1.16 (0.97 to 1.40)
25–29	110	1.02 (0.84 to 1.24)	1.09 (0.85 to 1.40)
≥30	80	0.94 (0.75 to 1.18)	1.00 (0.76 to 1.32)
P value for trend		0.002	0.04
Smoking started before first term pregnancy‡:			
Yes	833	1.21 (1.11 to 1.33)	1.28 (1.06 to 1.55)
No	116	1.12 (0.92 to 1.36)	1.17 (0.90 to 1.52)
Uncertain	618	1.07 (0.97 to 1.18)	1.13 (0.93 to 1.37)
Average No of cigarettes smoked/day:			
<5	383	1.05 (0.94 to 1.17)	1.12 (0.92 to 1.35)
5–14	560	1.11 (1.01 to 1.23)	1.19 (0.99 to 1.43)
15–24	503	1.14 (1.03 to 1.27)	1.22 (1.01 to 1.47)
≥25	301	1.08 (0.95 to 1.22)	1.15 (0.94 to 1.41)
P value for trend		0.01	0.16
Total No of smoking years:			
<5	220	0.97 (0.84 to 1.11)	1.03 (0.86 to 1.28)
5–9	182	1.05 (0.90 to 1.22)	1.12 (0.90 to 1.40)
10–19	401	1.09 (0.98 to 1.22)	1.17 (0.96 to 1.42)
20–29	387	1.09 (0.97 to 1.22)	1.16 (0.96 to 1.41)
30–39	341	1.21 (1.07 to 1.36)	1.29 (1.06 to 1.58)
40–49	178	1.14 (0.98 to 1.34)	1.22 (0.98 to 1.53)
≥50	56	1.35 (1.03 to 1.77)	1.45 (1.06 to 1.98)
P value for trend		0.0002	0.002
No of smoking pack years:			
<10	702	1.04 (0.95 to 1.14)	1.11 (0.93 to 1.33)
10–<20	359	1.21 (1.08 to 1.36)	1.29 (1.06 to 1.58)
20–<30	225	1.13 (0.98 to 1.30)	1.21 (0.98 to 1.50)
30–<40	168	1.01 (0.86 to 1.19)	1.08 (0.86 to 1.36)
40–<50	69	1.16 (0.91 to 1.47)	1.24 (0.92 to 1.65)
≥50	218	1.18 (1.02 to 1.37)	1.26 (1.02 to 1.57)
P value for trend		0.005	0.08
Years since quitting smoking (former smokers):			
<10	295	1.15 (1.01 to 1.31)	1.19 (0.96 to 1.47)
10–<20	416	1.14 (1.02 to 1.27)	1.17 (0.96 to 1.43)
20–<30	391	1.08 (0.96 to 1.21)	1.11 (0.91 to 1.36)
≥30	414	1.06 (0.95 to 1.18)	1.08 (0.89 to 1.32)
P value for trend		0.005	0.08

\*Reference category was lifetime non-smokers.

†Reference category was lifetime non-smokers with no history of exposure to passive smoking.

‡Among 69 533 women who had at least one full term pregnancy, 1429 cases were among never smokers and 125 cases in no exposure group.

## DISCUSSION

In this large prospective study in postmenopausal women, we observed an elevated risk of breast cancer in former smokers (9%) and current smokers (16%), particularly with smoking of high intensity, of long duration, and which started at an early age. Among former smokers, the time since quitting smoking was significantly inversely associated with breast cancer risk, and it took up to 20 years for a former smoker's risk to return to baseline. Finally, we observed a 32% excess risk of breast cancer associated with the most extensive exposure to passive smoking among women who had never been active smokers.

### Comparison with other studies

Until recently, most scientists agreed that there was not enough consistent evidence to determine whether active smoking plays a causal role in breast cancer.<sup>17–19</sup> However, a Canadian panel of experts reviewed the extensive new research in this area and concluded that the relations between active smoking and breast cancer, both premenopausal and postmenopausal, are consistent with causality, based on the weight of evidence from epidemiological and toxicological studies and on an understanding of biological mechanisms.<sup>12</sup> Our findings on active smoking are consistent with the most recent studies reporting a risk elevation of around 20%–50% in association with high intensity smoking of long duration and with early age at smoking initiation or smoking initiation before their first pregnancy among postmenopausal women.<sup>13–19,38–40</sup>

Our results also indicate that the elevated risk of breast cancer that smokers experience declined over time after smoking cessation, and it took up to 20 years for a former smoker's risk to return to baseline. Few studies have evaluated this relation in postmenopausal women, and their results were inconsistent. Some studies of predominantly postmenopausal women observed no relation between years since quitting smoking and breast cancer risk,<sup>15,16,18</sup> whereas other studies found results similar to ours.<sup>20,38</sup>

Our study is one of few prospective studies to observe a significantly increased risk of breast cancer associated with passive smoking among postmenopausal women. It seems unlikely that this could be explained by bias or confounding for several reasons. Firstly, our study is a prospective study, which circumvents problems of recall and selection bias common to case-control studies. In addition, the results remained similar when we excluded the first two years of follow-up (relative risk 1.38 (1.07 to 1.78) for the most extensive passive exposure). Secondly, we were able to adjust for all commonly known or suspected confounders for breast cancer. Thirdly, our study collected data on three major aspects of lifetime exposure (childhood, adult residential, and occupational), which enabled us to examine a reference group that had been unexposed to either active or passive smoking. Although our extensive variable of passive smoking exposure was

**Table 4** | Cases of incident invasive breast cancer and related hazard ratios among 79 990 postmenopausal women by cancer subtypes and exposure to tobacco smoke

	Passive smoking exposure*							Active smoking status				
	Never† No of cases	Any exposure		Non-extensive exposure‡		Extensive exposure‡		Never† No of cases	Former		Current	
		No of cases	Hazard ratio (95% CI)	No of cases	Hazard ratio (95% CI)	No of cases	Hazard ratio (95% CI)		No of cases	Hazard ratio (95% CI)	No of cases	Hazard ratio (95% CI)
<b>Cases defined by histology</b>												
Ductal cancer	98	969	1.02 (0.83 to 1.26)	878	0.99 (0.80 to 1.23)	91	1.20 (0.90 to 1.61)	1088	1013	1.07 (0.98 to 1.18)	91	1.16 (0.97 to 1.40)
Lobular cancer	14	157	1.22 (0.70 to 2.11)	147	1.25 (0.72 to 2.17)	10	0.97 (0.43 to 2.22)	175	164	1.08 (0.86 to 1.35)	27	1.64 (1.09 to 2.49)
<b>Cases defined by hormone receptor</b>												
OR+ PR+	93	941	1.04 (0.84 to 1.29)	851	1.01 (0.82 to 1.26)	90	1.23 (0.91 to 1.65)	1052	1049	1.12 (1.03 to 1.22)	139	1.28 (1.07 to 1.53)
OR+ PR-	18	184	1.14 (0.70 to 1.86)	165	1.09 (0.67 to 1.78)	19	1.53 (0.79 to 2.94)	207	213	1.18 (0.97 to 1.44)	13	0.98 (0.63 to 1.53)
OR- PR-	20	160	1.10 (0.69 to 1.75)	193	1.08 (0.68 to 1.72)	21	1.39 (0.74 to 2.58)	237	194	1.00 (0.82 to 1.22)	21	0.98 (0.65 to 1.49)

OR+ = oestrogen receptor positive. OR- = oestrogen receptor negative. PR+ = progesterone receptor positive. PR- = progesterone receptor negative.

\*Analyses performed only among lifetime non-smokers.

†Reference categories for hazard ratios. For active smoking, all non-smokers formed the reference category regardless of passive smoking exposure.

‡Extensive exposure defined as exposure in childhood for ≥10 years, in adult home for ≥20 years, and in adult work for ≥10 years.

defined a priori, there was no clear dose response to cumulative exposure, raising the possibility that the result was due to chance.

Previous epidemiological studies examining the association between passive smoking and breast cancer provided little evidence for a relation in postmenopausal women.<sup>16 18 25-29 41 42</sup> However, all but one of these cohort studies had important limitations in terms of exposure assessment. Given the high prevalence of passive smoking exposure in the population (more than 80% of women in Western populations have been exposed to regular residential or occupational passive smoking exposure),<sup>20</sup> inadequate assessment of exposure to passive smoking could have resulted in underestimating the passive smoking exposure status of most of those categorised as unexposed, thus contaminating the reference group and leading to an underestimate of risk. Johnson conducted a literature review and found that studies with more complete ascertainment of lifetime exposure, and which excluded women with passive smoking exposure from the reference group, consistently show higher breast cancer risks in association with passive smoking.<sup>20-24 43</sup> Recently, one cohort study that included a comprehensive measure of lifetime exposure to passive smoking<sup>30</sup> also observed an increased risk in the most highly exposed subgroup of postmenopausal women whose passive smoking began after 20 years of age (hazard ratio 1.25 (1.01 to 1.56)).

#### Possible biological mechanisms

The association of both active and passive smoking with increased risk of breast cancer is biologically plausible, especially when exposure occurs during certain critical periods. Animal models have shown that mammary tissue may be particularly susceptible to carcinogenic exposures during the years from start of puberty

to first full term pregnancy.<sup>1</sup> Breast epithelial cells do not become fully differentiated until after the first full term pregnancy. Thus, the period between the onset of puberty and first full term pregnancy may be a time of higher risk of cancer initiation.<sup>44</sup> Our findings of an elevated risk of breast cancer associated with early age of starting smoking (<15 years old) and starting smoking before first full term pregnancy support the hypothesis that smoking during the early teenage years increases the risk of breast cancer.<sup>44</sup> Alternatively, these results may simply be correlated with a long duration of smoking, which was also associated with increased risk in our cohort.

In addition, human biomarker studies have strongly suggested that breast tissue is a target for the carcinogenic effects of tobacco smoke.<sup>2</sup> Studies have found that DNA adducts (DNA covalently bonded to a carcinogenic chemical) with derivatives of tobacco smoke are more common in the breast tissue of smokers than that of non-smokers.<sup>6 45 46</sup> The wide variation between people in DNA adduct formation and repair is likely related to a combination of factors, including the properties of individual carcinogens and host genetic polymorphisms.<sup>47</sup> In 1996 Ambrosone et al reported that postmenopausal women who smoked and had the slow acetylator genotype for N-acetyltransferase 2 had a significantly elevated risk of breast cancer.<sup>48</sup> A recent meta-analysis of 13 studies of this association reached the same conclusion.<sup>49</sup>

Breast cancer is a heterogeneous disease with subtypes that may vary in their aetiologies. Our data suggest that current smoking was more strongly associated with lobular cancer than with ductal cancer and with tumours that were hormone receptor positive (for both oestrogen and progesterone) than the other three combinations of receptor status. One study examining smoking in relation to the risk of different histological

**WHAT IS ALREADY KNOWN ON THIS TOPIC**

Recent cohort studies have shown an increased risk of breast cancer among women who started smoking at a young age or who smoked for many years

An increased risk of breast cancer in women exposed to passive smoking has also been suggested but remains controversial

**WHAT THIS STUDY ADDS**

This cohort of postmenopausal women showed an increased risk of breast cancer among smokers, particularly those with high intensity and long duration of smoking

The study also found an increased risk of breast cancer among non-smokers with extensive exposure to passive smoking. However, the lack of significant association in other non-smokers with lower exposure and no clear dose response to cumulative exposure mean the association between passive smoking and breast cancer should be considered suggestive only

subtypes of breast cancer among postmenopausal women reported that recent smoking had a borderline association with risk of ductal cancer (odds ratio 1.2 (95% confidence interval 1.0 to 1.4)) but not with other histological types.<sup>50</sup> Our findings suggesting a particular association with lobular cancer should be considered preliminary. A few studies examining smoking in relation to subtypes of breast cancer by hormone receptor status have yielded mixed results.<sup>51-54</sup> Some have reported an association with hormone receptor positive cancer,<sup>53,54</sup> but others have reported the opposite<sup>51,52</sup> or no association.<sup>55</sup> Finally, Kabat et al studied active smoking and ductal carcinoma in situ (487 cases among 63 393 women enrolled in the Women's Health Initiative) and did not find an association.<sup>56</sup> More large studies are needed to clarify these relations.

**Strengths and limitations of the study**

Our study's strengths include the prospective design, the large size and broad geographical distribution of the cohort, the large number of cancer cases, pathological confirmation of cases by trained adjudicators and cancer coders, detailed information on potential confounders, and, particularly, detailed information on passive smoking, including quantitative measures of exposure in childhood and adult exposure in residential and workplace settings.

However, our study also has limitations. One is that we used only baseline values for women's smoking status and all covariates and did not account for changes in exposure during follow-up, which may have caused some exposure misclassification and biased our results towards the null. However, given that our study population consisted of postmenopausal women and that few were current smokers, the effect of changes in smoking status during follow-up is likely to have been minimal. Based on yearly reassessment of smoking status in the Women's Health Initiative, about 60% of smokers continued to smoke for six years of follow-up and 99% of non-smokers remained abstinent.

Secondly, there was possible recall bias with regard to passive smoking exposure in childhood. In addition,

data were not collected for the intensity or frequency of passive smoke exposure. The crude measures and potential misclassification of exposure may make our estimate more conservative. The lack of intensity or frequency data also may have hampered our ability to detect a dose-response relation.

**Conclusions and policy implications**

Our study supports the hypothesis that active smoking is associated with an increased risk of breast cancer among postmenopausal women. In addition, our data suggest that extensive exposure to passive smoking may increase breast cancer risk. However, since risk of breast cancer was restricted to the most extensive passive smoking category with no clear dose response, the association with passive smoking should be considered suggestive only and needs confirmation from other studies. Our findings highlight the need for interventions to prevent initiation of smoking, especially at an early age, and to encourage smoking cessation at all ages. Future studies examining how genetic polymorphisms and other risk factors modify the effect of tobacco exposure on breast cancer risk are likely to further our understanding of this important issue.

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- 1 International Agency for Research on Cancer. *IARC monographs on the evaluation of carcinogenic risk of chemicals to humans*. IARC, 2004.
- 2 Conway K, Edmiston SN, Cui L, Drouin SS, Pang J, He M, et al. Prevalence and spectrum of p53 mutations associated with smoking in breast cancer. *Cancer Res* 2002;62:1987-95.
- 3 Li D, Zhang W, Sahin AA, Hittelman WN. DNA adducts in normal tissue adjacent to breast cancer: a review. *Cancer Detect Prev* 1999;23:454-62.
- 4 Rundle A, Tang D, Hibshoosh H, Estabrook A, Schnabel F, Cao W, et al. The relationship between genetic damage from polycyclic aromatic hydrocarbons in breast tissue and breast cancer. *Carcinogenesis* 2000;21:1281-9.
- 5 Li D, Wang M, Dhingra K, Hittelman WN. Aromatic DNA adducts in adjacent tissues of breast cancer patients: clues to breast cancer etiology. *Cancer Res* 1996;56:287-93.
- 6 Firozi PF, Bondy ML, Sahin AA, Chang P, Lukmanji F, Singletary ES, et al. Aromatic DNA adducts and polymorphisms of CYP1A1, NAT2, and GSTM1 in breast cancer. *Carcinogenesis* 2002;23:301-6.
- 7 Hamajima N, Hirose K, Tajima K, Rohan T, Calle EE, Heath CW Jr, et al. Alcohol, tobacco and breast cancer—collaborative reanalysis of individual data from 53 epidemiological studies, including 58,515

- women with breast cancer and 95,067 women without the disease. *Br J Cancer* 2002;87:1234-45.
- 8 Palmer JR, Rosenberg L. Cigarette smoking and the risk of breast cancer. *Epidemiol Rev* 1993;15:145-56.
  - 9 Terry PD, Rohan TE. Cigarette smoking and the risk of breast cancer in women: a review of the literature. *Cancer Epidemiol Biomarkers Prev* 2002;11:953-71.
  - 10 The 2004 United States Surgeon General's Report: the health consequences of smoking. *N S W Public Health Bull* 2004;15:107.
  - 11 Kopans DB. Standardized mammography reporting. *Radiol Clin North Am* 1992;30:257-64.
  - 12 Collishaw NE, Boyd NF, Cantor KP, Hammond SK, Johnson KC, Millar J, et al. *Canadian expert panel on tobacco smoke and breast cancer risk*. Ontario Tobacco Research Unit, 2009. [www.otru.org/pdf/special/expert\\_panel\\_tobacco\\_breast\\_cancer.pdf](http://www.otru.org/pdf/special/expert_panel_tobacco_breast_cancer.pdf).
  - 13 Olson JE, Vachon CM, Vierkant RA, Sweeney C, Limburg PJ, Cerhan JR, et al. Prepregnancy exposure to cigarette smoking and subsequent risk of postmenopausal breast cancer. *Mayo Clin Proc* 2005;80:1423-8.
  - 14 Gram IT, Braaten T, Terry PD, Sasco AJ, Adami HO, Lund E, et al. Breast cancer risk among women who start smoking as teenagers. *Cancer Epidemiol Biomarkers Prev* 2005;14:61-6.
  - 15 Cui Y, Miller AB, Rohan TE. Cigarette smoking and breast cancer risk: update of a prospective cohort study. *Breast Cancer Res Treat* 2006;100:293-9.
  - 16 Reynolds P, Hurley S, Goldberg DE, Anton-Culver H, Bernstein L, Deapen D, et al. Active smoking, household passive smoking, and breast cancer: evidence from the California Teachers Study. *J Natl Cancer Inst* 2004;96:29-37.
  - 17 Al-Delaimy WK, Cho E, Chen WY, Colditz G, Willet WC. A prospective study of smoking and risk of breast cancer in young adult women. *Cancer Epidemiol Biomarkers Prev* 2004;13:398-404.
  - 18 Egan KM, Stampfer MJ, Hunter D, Hankinson S, Rosner BA, Holmes M, et al. Active and passive smoking in breast cancer: prospective results from the Nurses' Health Study. *Epidemiology* 2002;13:138-45.
  - 19 Ha M, Mabuchi K, Sigurdson AJ, Freedman DM, Linet MS, Doody MM, et al. Smoking cigarettes before first childbirth and risk of breast cancer. *Am J Epidemiol* 2007;166:55-61.
  - 20 Johnson KC, Hu J, Mao Y. Passive and active smoking and breast cancer risk in Canada, 1994-97. *Cancer Causes Control* 2000;11:211-21.
  - 21 Kropp S, Chang-Claude J. Active and passive smoking and risk of breast cancer by age 50 years among German women. *Am J Epidemiol* 2002;156:616-26.
  - 22 Morabia A, Bernstein M, Heritier S, Khatchatrian N. Relation of breast cancer with passive and active exposure to tobacco smoke. *Am J Epidemiol* 1996;143:918-28.
  - 23 Smith SJ, Deacon JM, Chilvers CE. Alcohol, smoking, passive smoking and caffeine in relation to breast cancer risk in young women. UK National Case-Control Study Group. *Br J Cancer* 1994;70:112-9.
  - 24 Zhao Y, Shi Z, Liu L. [Matched case-control study for detecting risk factors of breast cancer in women living in Chengdu.] *Zhonghua Liu Xing Bing Xue Za Zhi* 1999;20:91-4.
  - 25 Wartenberg D, Calle EE, Thun MJ, Heath CW Jr, Lally C, Woodruff T. Passive smoking exposure and female breast cancer mortality. *J Natl Cancer Inst* 2000;92:1666-73.
  - 26 Pirie K, Beral V, Peto R, Roddam A, Reeves G, Green J. Passive smoking and breast cancer in never smokers: prospective study and meta-analysis. *Int J Epidemiol* 2008;37:1069-79.
  - 27 Hanaoka T, Yamamoto S, Sobue T, Sasaki S, Tsugane S. Active and passive smoking and breast cancer risk in middle-aged Japanese women. *Int J Cancer* 2005;114:317-22.
  - 28 Wells AJ. Breast cancer, cigarette smoking, and passive smoking. *Am J Epidemiol* 1991;133:208-10.
  - 29 Jee SH, Ohrr H, Kim IS. Effects of husbands' smoking on the incidence of lung cancer in Korean women. *Int J Epidemiol* 1999;28:824-8.
  - 30 Reynolds P, Goldberg D, Hurley S, Nelson DO, Largent J, Henderson KD, et al. Passive smoking and risk of breast cancer in the California teachers study. *Cancer Epidemiol Biomarkers Prev* 2009;18:3389-98.
  - 31 The Women's Health Initiative Study Group. Design of the Women's Health Initiative clinical trial and observational study. *Control Clin Trials* 1998;19:61-109.
  - 32 Hays J, Hunt JR, Hubbell FA, Anderson GL, Limacher M, Allen C, et al. The Women's Health Initiative recruitment methods and results. *Ann Epidemiol* 2003;13(suppl 9):18-77S.
  - 33 Langer RD, White E, Lewis CE, Kotchen JM, Hendrix SL, Trevisan M. The Women's Health Initiative Observational Study: baseline characteristics of participants and reliability of baseline measures. *Ann Epidemiol* 2003;13(suppl 9):107-21S.
  - 34 Surveillance Epidemiology and End Results. *SEER extent of disease: 1988 codes and coding instructions*. 3rd ed. National Cancer Institute, US National Institutes of Health, 1998. <http://seer.cancer.gov/manuals/EOD10Dig.3rd.pdf>.
  - 35 Hammond SK. Exposure of US workers to environmental tobacco smoke. *Environ Health Perspect* 1999;107(suppl 2):329-40.
  - 36 Hammond SK, Sorensen G, Youngstrom R, Ockene JK. Occupational exposure to environmental tobacco smoke. *JAMA* 1995;274:956-60.
  - 37 Hess KR. Graphical methods for assessing violations of the proportional hazards assumption in Cox regression. *Stat Med* 1995;14:1707-23.
  - 38 Li CI, Malone KE, Daling JR. The relationship between various measures of cigarette smoking and risk of breast cancer among older women 65-79 years of age (United States). *Cancer Causes Control* 2005;16:975-85.
  - 39 Band PR, Le ND, Fang R, Deschamps M. Carcinogenic and endocrine disrupting effects of cigarette smoke and risk of breast cancer. *Lancet* 2002;360:1044-9.
  - 40 Lissowska J, Brinton LA, Zatonski W, Blair A, Bardin-Mikolajczak A, Peplonska B, et al. Tobacco smoking, NAT2 acetylation genotype and breast cancer risk. *Int J Cancer* 2006;119:1961-9.
  - 41 Miller MD, Marty MA, Broadwin R, Johnson KC, Salmon AG, Winder B, et al. The association between exposure to environmental tobacco smoke and breast cancer: a review by the California Environmental Protection Agency. *Prev Med* 2007;44:93-106.
  - 42 Department of Health and Human Services. The health consequences of involuntary exposure to tobacco smoke: a report of the Surgeon General. US Department of Health and Human Services, 2006.
  - 43 Johnson KC. Accumulating evidence on passive and active smoking and breast cancer risk. *Int J Cancer* 2005;117:619-28.
  - 44 Palmer JR, Rosenberg L, Clarke EA, Stolley PD, Warshauer ME, Zauber AG, et al. Breast cancer and cigarette smoking: a hypothesis. *Am J Epidemiol* 1991;134:1-13.
  - 45 Perera FP, Estabrook A, Hewer A, Channing K, Rundle A, Mooney LA, et al. Carcinogen-DNA adducts in human breast tissue. *Cancer Epidemiol Biomarkers Prev* 1995;4:233-8.
  - 46 Faraglia B, Chen SY, Gammon MD, Zhang Y, Teitelbaum SL, Neugut AI, et al. Evaluation of 4-aminobiphenyl-DNA adducts in human breast cancer: the influence of tobacco smoke. *Carcinogenesis* 2003;24:719-25.
  - 47 Harris CC. Interindividual variation among humans in carcinogen metabolism, DNA adduct formation and DNA repair. *Carcinogenesis* 1989;10:1563-6.
  - 48 Ambrosone CB, Freudenheim JL, Graham S, Marshall JR, Vena JE, Brasure JR, et al. Cigarette smoking, N-acetyltransferase 2 genetic polymorphisms, and breast cancer risk. *JAMA* 1996;276:1494-501.
  - 49 Ambrosone CB, Kropp S, Yang J, Yao S, Shields PG, Chang-Claude J. Cigarette smoking, N-acetyltransferase 2 genotypes, and breast cancer risk: pooled analysis and meta-analysis. *Cancer Epidemiol Biomarkers Prev* 2008;17:15-26.
  - 50 Rosenberg LU, Magnusson C, Lindstrom E, Wedren S, Hall P, Dickman PW. Menopausal hormone therapy and other breast cancer risk factors in relation to the risk of different histological subtypes of breast cancer: a case-control study. *Breast Cancer Res* 2006;8:R11.
  - 51 Manjer J, Malina J, Berglund G, Bondeson L, Game JP, Janzon L. Smoking associated with hormone receptor negative breast cancer. *Int J Cancer* 2001;91:580-4.
  - 52 Cooper JA, Rohan TE, Cant EL, Horsfall DJ, Tilley WD. Risk factors for breast cancer by oestrogen receptor status: a population-based case-control study. *Br J Cancer* 1989;59:119-25.
  - 53 London SJ, Colditz GA, Stampfer MJ, Willett WC, Rosner BA, Speizer FE. Prospective study of smoking and the risk of breast cancer. *J Natl Cancer Inst* 1989;81:1625-31.
  - 54 Yoo KY, Tajima K, Miura S, Takeuchi T, Hirose K, Risch H, et al. Breast cancer risk factors according to combined estrogen and progesterone receptor status: a case-control analysis. *Am J Epidemiol* 1997;146:307-14.
  - 55 Althuis MD, Fergenbaum JH, Garcia-Closas M, Brinton LA, Madigan MP, Sherman ME. Etiology of hormone receptor-defined breast cancer: a systematic review of the literature. *Cancer Epidemiol Biomarkers Prev* 2004;13:1558-68.
  - 56 Kabat GC, Kim M, Kakani C, Tindle H, Wactawski-Wende J, Ockene JK, et al. Cigarette smoking in relation to risk of ductal carcinoma in situ of the breast in a cohort of postmenopausal women. *Am J Epidemiol* 2010;172:591-9.

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