



# Periodontal Disease and Breast Cancer: A Meta-Analysis of 1,73,162 Participants

Jun Shao<sup>1†</sup>, Lan Wu<sup>2,3,4†</sup>, Wei-Dong Leng<sup>5</sup>, Cheng Fang<sup>2,3</sup>, You-Jia Zhu<sup>4</sup>, Ying-Hui Jin<sup>2,3\*</sup> and Xian-Tao Zeng<sup>1,2,3,4,5\*</sup>

<sup>1</sup> Department of Stomatology, Guangzhou Hospital of Integrated Traditional and West Medicine, Guangzhou, China, <sup>2</sup> Center for Evidence-Based and Translational Medicine, Zhongnan Hospital of Wuhan University, Wuhan, China, <sup>3</sup> Department of Evidence-Based Medicine and Clinical Epidemiology, The Second Clinical College of Wuhan University, Wuhan, China, <sup>4</sup> Department of Stomatology, Zhongnan Hospital of Wuhan University, Wuhan, China, <sup>5</sup> Department of Stomatology, Taihe Hospital, Hubei University of Medicine, Shiyan, China

## OPEN ACCESS

### Edited by:

Sarah M. Temkin,  
Virginia Commonwealth University,  
United States

### Reviewed by:

Tarah Ballinger,  
Indiana University, Purdue University  
Indianapolis, United States  
Alessandro Igor Cavalcanti Leal,  
Johns Hopkins Medicine,  
United States

### \*Correspondence:

Ying-Hui Jin  
jinyinghuiebm@163.com  
Xian-Tao Zeng  
zengxiantao1128@163.com

<sup>†</sup>These authors have contributed  
equally to this work and are joint first  
authors

### Specialty section:

This article was submitted to  
Women's Cancer,  
a section of the journal  
Frontiers in Oncology

**Received:** 03 September 2018

**Accepted:** 26 November 2018

**Published:** 12 December 2018

### Citation:

Shao J, Wu L, Leng WD, Fang C, Zhu  
YJ, Jin YH and Zeng XT (2018)  
Periodontal Disease and Breast  
Cancer: A Meta-Analysis of 1,73,162  
Participants. *Front. Oncol.* 8:601.  
doi: 10.3389/fonc.2018.00601

**Objective:** To investigate the correlation between periodontal disease and breast cancer.

**Materials and Methods:** PubMed and China National Knowledge Infrastructure (CNKI) databases were searched up to February 8, 2018 for observational studies examining the association between periodontal disease and breast cancer. Study selection was conducted according to predesigned eligibility criteria, and two authors independently extracted data from included studies. Meta-analysis was performed using the Comprehensive Meta-Analysis v2 software and risk estimates were calculated as relative risks (RRs) with corresponding 95% confidence intervals (CIs).

**Results:** A total of 11 study were included. Meta-analysis indicated that periodontal disease significantly increased the risk of breast cancer by 1.22-fold (RR = 1.22, 95% CI = 1.06–1.40). Amongst participants with periodontal patients and a history of periodontal therapy, the risk of developing breast cancer was not significant (RR = 1.23; 95% CI = 0.95–1.60). The association results between periodontal diseases and breast cancer were found to be robust, as evident in the leave-one-out sensitivity analysis.

**Conclusions:** Periodontal disease may be a potential risk factor for the development of breast cancer among women, and thus effective periodontal therapy may present as a valuable preventive measure against breast cancer.

**Keywords:** periodontal disease, breast cancer, risk factor, periodontal therapy, meta-analysis

## INTRODUCTION

Periodontal disease, a complex polymicrobial inflammatory disease, is a public health burden and is partially responsible for tooth loss. Existing epidemiological studies have linked periodontal disease to numerous systemic conditions, such as cardiovascular disease (1), preterm birth (2), osteoporosis (3), and diabetes mellitus (4), all of which may be attributed to systemic infection and inflammation (5). In recent years, there is increased interests in exploring the relationship between periodontal disease and cancer risk, particularly for cancers in the head and neck, upper gastrointestinal system, lung, and pancreas (6–9).

Cancer is a leading cause of mortality worldwide. Among the female-specific malignancies, breast cancer remains the most common cancer type in developed countries, accounting for one-third of newly diagnosed cancers (10). Available data indicated a number of risk factor associated with this cancer, such as endogenous hormone levels, immune factors and genetic susceptibility and lifestyle habits (11–14). Periodontal health status has also been identified as a risk factor for breast cancer, a study of Chung et al. (15), reported an increased risk of breast cancer among patients with chronic periodontitis [Hazard ratio (HR) = 1.23, 95% confidence interval (CI) = 1.11–1.36] although no such observation was found amongst subjects who underwent gingivectomy or periodontal flap operation (HR = 1.17, 95% CI = 0.86–1.58). However, Published studies reported inconsistent findings on the association between periodontal disease and breast cancer, with some supporting their significant association while others claiming none. For example, Freudenheim et al. (16), reported that periodontal disease significantly increased breast cancer risk (HR = 1.14, 95% CI = 1.03–1.26), particularly among former smokers who quit within 20 years (HR = 1.36, 95% CI = 1.05–1.77); while Mai et al. (17) and Han et al. (18) reported different observations. Given that single epidemiologic studies may be inadequate for identifying etiological influence of periodontal disease on breast cancer. We consequently conducted this meta-analysis of case-control and cohort studies to provide updated evidence on the relationship between periodontal disease and female-specific breast cancer.

## MATERIALS AND METHODS

Our study was conducted and reported according to the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) statement (19).

## Eligible Criteria

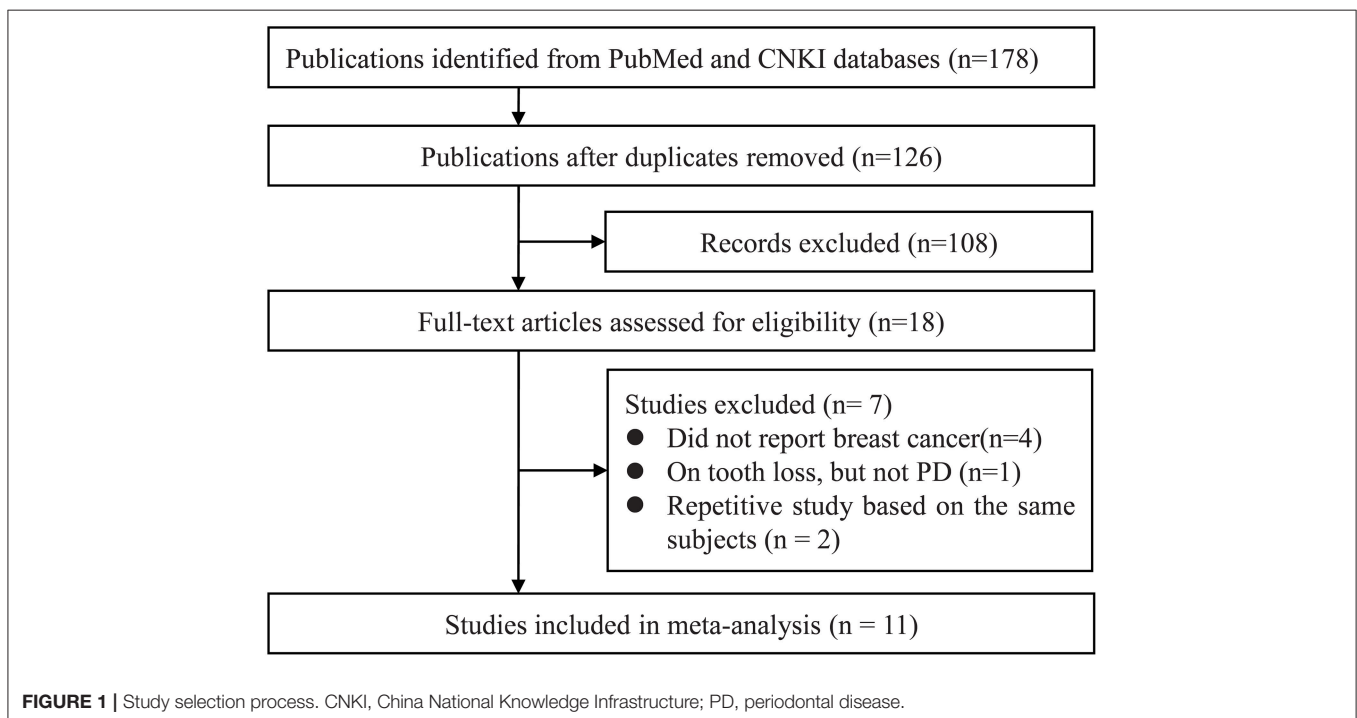
Studies were eligible for inclusion if they met the following criteria: (1) adopting a cohort or case-control design; (2) study participants with periodontal disease, as the exposure of interest; (3) female-specific breast cancer, as the outcome of interest; (4) providing relative risks (RRs), HRs or odds ratios (ORs) and corresponding 95% CIs, or data for their calculation; (5) published in full in English-language or Chinese-language peer reviewed journals. For multiple study reports with the same study population, data from the longest follow-up duration were extracted and analyzed.

## Search Strategy

PubMed and China National Knowledge Infrastructure (CNKI) databases were searched up to February 8, 2018 for case-control and cohort studies examining the association between periodontal disease and breast cancer. The following key words and subject terms were used: “periodontitis,” “periodontal disease,” “oral status,” “dental health,” “breast cancer,” “cancer,” and “carcinoma.”

## Data Extraction

Two authors independently extract the following data extraction from all eligible studies: the first author’s name, year of publication, country, study design, follow-up period, age, sample size, periodontal disease, and breast cancer ascertainment, type of periodontal therapy, reference of control, dental and smoking status, risk estimates with corresponding 95% CIs, and corresponding adjustments. Any disagreements over the extracted data were resolved by consensus or by consulting a third author.



**TABLE 1** | Characteristics of included studies.

Study ID	Sample	Study design (study period)	Follow-up period (yrs)	Age (yrs)	Cases/sample	PD ascertainment	Breast cancer ascertainment	Periodontal therapy	Control	Dental /smoking status	Risk estimates (95% CI)
Hujoel et al. (28)	The NHANES I Epidemiologic Follow-up Study (NHEFS); USA	Prospective cohort (1971/1975–1992)	10	25–74	79/6862	Clinical diagnosis	ICD-9	Unreported	No PD	PD	OR = 1.32 (0.74–2.38)
Arora et al. (29)	The Swedish Twin Registry; Sweden	Prospective cohort (1963–2004)	27	38–77	531/8433	Self-report	ICD	Unreported	No PD	PD Minor mobility	OR = 0.62 (0.32–1.21) OR = 0.59 (0.32–1.10) HR = 1.12(0.75–1.68) HR = 1.22(0.92–1.61)
Söder et al. (30)	A registry database of all inhabitants of Stockholm County; Sweden	Prospective cohort (1985–2001)	16	30–40	39/1586	Clinical diagnosis	ICD-7, ICD-9, ICD-10	Unreported	No PD	PD	OR = 13.08 (3.09–55.32)
Chung et al. (15)	The Taiwan National Health Insurance (NHI) program; China	Retrospect cohort (1995–2004)	5	54.1±11.5	367/42548	Clinical diagnosis	ICD-9-CM	Included	No PD	CP	HR = 1.23(1.11–1.36)
Mai et al. (17)	The Buffalo OsteoPerio Study; USA	Prospective cohort (1997–2014)	12.2 ± 4.2	53–85*	89/1337	Measured ACH	Medical records	Unreported	No PD	Periodontal therapy Mild/moderate P Severe P Over all#	HR = 1.15 (0.67–1.99) HR = 0.94 (0.49–1.82) HR = 1.06 (0.70–1.61)#
Dizdar et al. (31)	The Hacettepe University Dentistry and Oncology hospitals in Ankara; Turkey	Retrospect cohort (2001–2010)	12	49.6	5/151	Clinical and radiographic parameters	Patient files and hospital registries	Included	No control, use national cancer registry figures	CP (moderate or severe)	RR = 2.40 (0.88–6.55)
Han (18)	The sixth Korean National Health and Nutrition Examination Survey (KNHANES); Korea	Prospective cohort (2013–2015)	2	>19	84/1979	Community periodontal index	Health interview	Unreported	No PD	PD	OR = 1.56(0.79–3.06)
Nwizu et al. (32)	Women's Health Initiative Observational Study (WHI-OS); USA	Prospective cohort (1994/1998–2013)	8.32	50–79*	2416/65869	Self-report and medical records	ICD-O-2	Unreported	No PD	PD	HR = 1.13 (1.03–1.23)

(Continued)

TABLE 1 | Continued

Study ID	Sample	Study design (study period)	Follow-up period (yrs)	Age (yrs)	Cases/sample	PD ascertainment	Breast cancer ascertainment	Periodontal therapy	Control	Dental /smoking status	Risk estimates (95% CI)
Sfreddo et al. (33)	The University Hospital of Santa Maria (UHSM); Brazil	Case-control (2013–2015)	/	55.0 ± 8.9/ 54.6 ± 8.7	67/201	Clinical diagnosis	ICD-10 C50	Unreported	No PD	Never smokers Former smokers Current smokers PD	1.05 (0.92–1.21) 1.22 (1.08–1.37) 1.32 (0.89–1.97) OR = 2.72(1.18–6.27) Corrected RR = 2.07(1.13–3.80) OR = 5.08(1.66–15.61) OR = 2.39(1.03–5.52) RR = 1.19 (0.66–2.12)
Heikkilä et al. (34)	The patient register of the Public Dental Service of the City of Helsinki; Finland	Prospective cohort (2001/2002–2013)	10.1	>29	74/40108	periodontal pocket depth	ICD-10	Included	No PD	Never smokers Non-drinkers PD	HR = 1.32(0.81–2.14)
Michaud et al. (24)	The Atherosclerosis Risk in Communities (ARIC) study; USA	Prospective cohort (1987–2012)	14.7	44–66	202/4088	Clinical diagnosis	Patient files and hospital registries	Unreported	No PD	Mild Moderate Severe Over all# Edentulism	HR = 0.89(0.61–1.29) HR = 0.84(0.49–1.43) HR = 0.98(0.76–1.27) HR = 0.88(0.56–1.36)

\*Postmenopausal women; #Over all data was calculated by pooling mild/moderate P and severe P using a fixed-effects model for the final analysis. ACH, Oral alveolar crestal height; ICD, International classification of diseases; CP, chronic periodontitis; PD, periodontal disease; P, Periodontitis; CI, Confidence interval; HR, Hazard ratio; OR, Odds ratio; RR, Relative risk.

## Statistical Analysis

Association between periodontal disease and breast cancer risk was assessed using risk ratios (RRs) and corresponding 95% CIs across the included studies. HRs and incidence density ratios (IDRs) were directly considered to be equivalent to RRs. Where ORs were reported they were transformed into RRs using this formula:  $RR = OR / [(1 - P_0) + (P_0 \times OR)]$ , in which  $P_0$  is the incidence of the outcome of interest in the non-exposed group (20), standard errors (SEs) of the corrected RRs were calculated from this formula:  $SE_{log}(\text{relative risk}) = SE_{log}(\text{odds ratio}) \times \log(\text{relative risk}) / \log(\text{odds ratio})$  (21). When these data could not be obtained through calculation, the ORs were directly pooled with tolerance. Since these transformations may underestimate the variance of the RRs derived from the ORs (22, 23), we performed a sensitivity analysis by excluding studies for which this transformation had been applied. For two studies (17, 24) that reported HRs for mild/moderate periodontitis and severe periodontitis, respectively, we constructed subgroups and calculated a combined HR using a fixed-effect model for the main analysis. Heterogeneity among included studies was investigated using the  $I^2$  statistics and Q test, and low statistical heterogeneity was defined as  $I^2 \leq 50\%$  and  $p > 0.1$ , respectively. A fixed-effects model was utilized in analyses where statistical heterogeneity was shown to be low; otherwise a random-effect model was adopted (25). Subgroup analyses were conducted according to different characteristics of the included studies. The leave-one-out sensitivity analysis was performed to detect the robustness of the association results by removing one study at a time. Publication bias among the included studies was examined visually using funnel plots (26). Statistical analyses were undertaken using the Comprehensive Meta-Analysis v2.2 software (CMA v2.2) (27).

## RESULTS

### Study Selection and Characteristics

Our systematic literature search yielded a total of 178 publications. After de-duplication 126 titles and abstracts were assessed for eligibility, of these, full texts articles of 18 records were retrieved for further assessment. Eventually, 11 studies were included in the qualitative synthesis (15, 17, 24, 28–34) (Figure 1).

The 11 included studies were published in the period of 2003 and 2018. Of these, 7 were prospective cohort studies, 3 were retrospective cohort studies and one was case-control study, involving a total of 1,73,162 of whom 3,953 were diagnosed with breast cancer. Among these studies, one focused on co-twin (29), two on postmenopausal woman (17, 32) and on individuals who had a history of periodontal therapy and on females who are currently treated by periodontal therapy (15, 31, 34). Three included studies (15, 31, 32) also provided information on the participants' baseline dental or smoking status. Majority of the included studies (15, 17, 24, 28–34) chose healthy subjects without periodontal disease as controls, however one study (31) enrolled national cancer registry figures as the control group. All of studies performed covariate adjustment. Characteristics of included studies are summarized in Tables 1, 2.

**TABLE 2 |** Covariate adjustments performed in the included studies.

Study ID	Adjusted covariates
Hujoel et al. (28)	Age
Arora et al. (29)	Age, education, employment, number of siblings, smoking status, smoking status of partner, alcohol status, and body mass index
Söder et al. (30)	Age, education level, socio economic status, working history, yearly income, smoking in pack-years, dental appointments, dental plaque index, gingival bleeding index, and loss of any molar tooth in the mandible
Chung et al. (15)	Patient's monthly income, geographic region, and diabetes
Mai et al. (17)	Age at visit (continuous) and pack-years of smoking (continuous)
Dizdar et al. (31)	Age
Han (18)	Household income, education, smoking status, drinking frequency, and chronic disease
Nwizu et al. (32)	Age, BMI
Sfreddo et al. (33)	Age, parity, total breastfeeding time, hormonal replacement therapy, BMI
Heikkilä et al. (34)	Calendar time, age, socio-economic status, number of teeth, dental treatments (gingivitis, caries, endodontic caries, surgery and prosthesis), oral health indices (I, D, DMF), need of periodontal treatment and diabetes
Michaud et al. (24)	Age, field center, education level, smoking status, smoking duration, drinking status, body mass index, and diabetes status (diagnosed diabetes, undiagnosed diabetes, at risk for diabetes [reference is normal]), hormone replacement therapy use, joint terms for field center and race

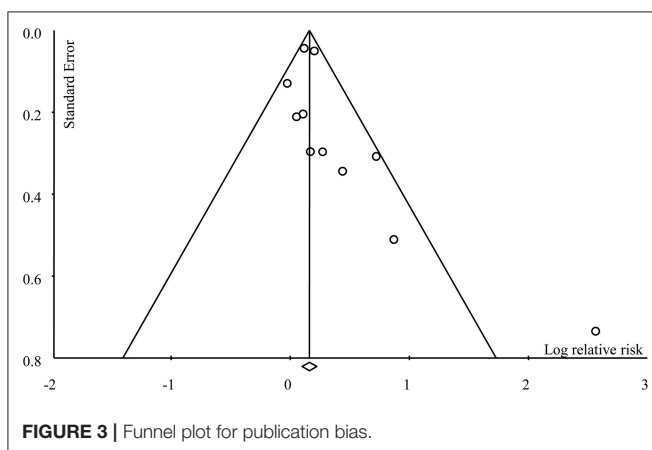
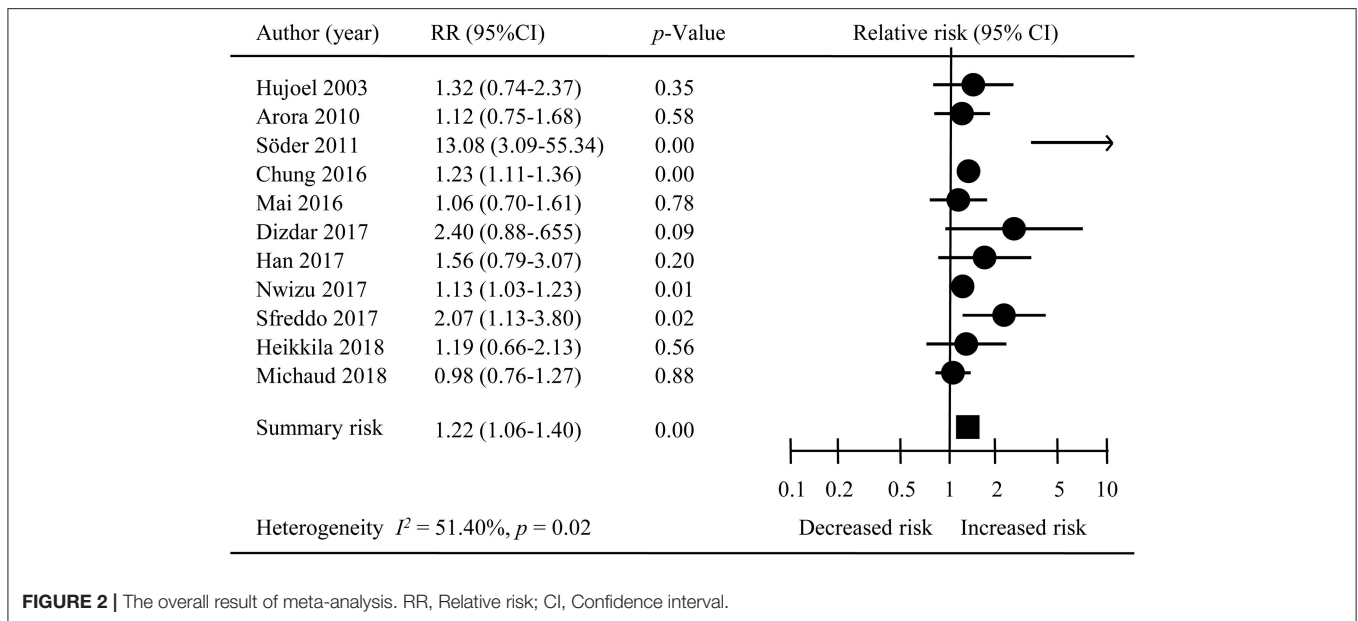
BMI, Body mass index; I, Initial caries; D, Decayed teeth; DMF, Decayed-missing-filled teeth; NSAID, Non-steroidal anti-inflammatory drug.

## Overall Analysis

The overall meta-analysis of 11 studies suggested that the risk of developing female-specific breast cancer was 1.22 times greater amongst patients with periodontal disease than those without periodontal disease (RR = 1.22, 95% CI = 1.06–1.40; Figure 2). Moderate statistical heterogeneity was noted across the 11 included studies ( $I^2 = 51.40\%$ ;  $P = 0.04$ ). Visual inspection of a funnel plot revealed a degree of publication bias (Figure 3).

## Subgroup Analyses

A number of pre-specified subgroup analyses, including study design, follow-up period, pausimonia status, medical history of periodontal therapy, and dental status were performed (Table 3). The impact of study design on final results was investigated by stratifying analyses according to prospective cohort study design ( $n = 8$ ) (17, 18, 24, 28–30, 32, 34), retrospect cohort study design ( $n = 2$ ) (15, 31) and case-control study design (33), and the results showed statistical difference (Prospective cohort: RR = 1.17, 95% CI = 0.97–1.41; retrospective cohort: RR = 1.24, 95% CI = 1.12–1.37). In terms of follow-up duration, three studies (15, 18, 32) that followed study participants for <10-year indicated a significant impact of periodontal disease on the risk of developing breast cancer (RR = 1.18, 95% CI = 1.10–1.26); in the 7 studies (17, 24, 29–34) with follow-up period of  $\geq 10$  years, the association of periodontal disease and breast cancer risk was



not statistically significant (RR = 1.45, 95% CI = 0.95–1.76). And these results suggested that study design and follow-up period might have certain effect on study results. Amongst three studies describing the use of periodontal therapy (15, 31, 34), results of subgroup analysis indicated a non-significant association of periodontal disease and breast cancer (RR = 1.23; 95% CI = 0.95–1.60); analysis of eight studies (17, 18, 24, 28–30, 32, 33) without clear reporting of revealed a distinct relationship of periodontal disease and breast cancer risk (RR = 1.25, 95% CI = 1.01–1.55). Results of the subgroup analyses according to pausimemia status and dental status were consistent with the overall results. Due to limited data, we were unable to pursue subgroup analysis by study participants' smoking status.

### Sensitivity Analyses

We conducted sensitivity analysis by removing one study at a time from the overall met-analysis (Figure 4), Omission of a study (31), which did not precisely state whether its

control individuals had periodontal disease, did not lead to any substantial changes in the overall results (RR = 1.20; 95% CI = 1.05–1.37); similar observation was noted upon exclusion of a study on co-twin (29) (RR = 1.23; 95% CI = 1.06–1.43). We also sequentially removed four studies (18, 28, 30, 33) reporting their association results, and a noticeable change in statistical heterogeneity was observed, (overall analysis:  $I^2 = 51.40\%$ ,  $P = 0.04$ ; sensitivity analysis:  $I^2 = <0.01$ ,  $P = 0.48$ ) although the pooled estimates (RR = 1.16; 95% CI = 1.09–1.23) remained consistent between the overall results.

The leave-one-out sensitivity analysis led to a decline of the level of statistical heterogeneity declined from moderate ( $I^2 = 51.40\%$ ,  $P = 0.04$ ) to low one ( $I^2 = 8.56\%$ ,  $P = 0.36$ ) upon removing the study by Söder et al. (30), indicating that this study contributed to a considerable to the statistical part of heterogeneity across studies.

## DISCUSSION

This meta-analysis of 11 observational studies suggested that periodontal disease might be a risk factor for female-specific breast cancer. Finding from sensitivity analyses confirmed the robustness of our association results. According to the findings from subgroup analyses, a number of confounding factors might be influencing the results of the overall analysis, including study design, follow-up period, and history of periodontal therapy.

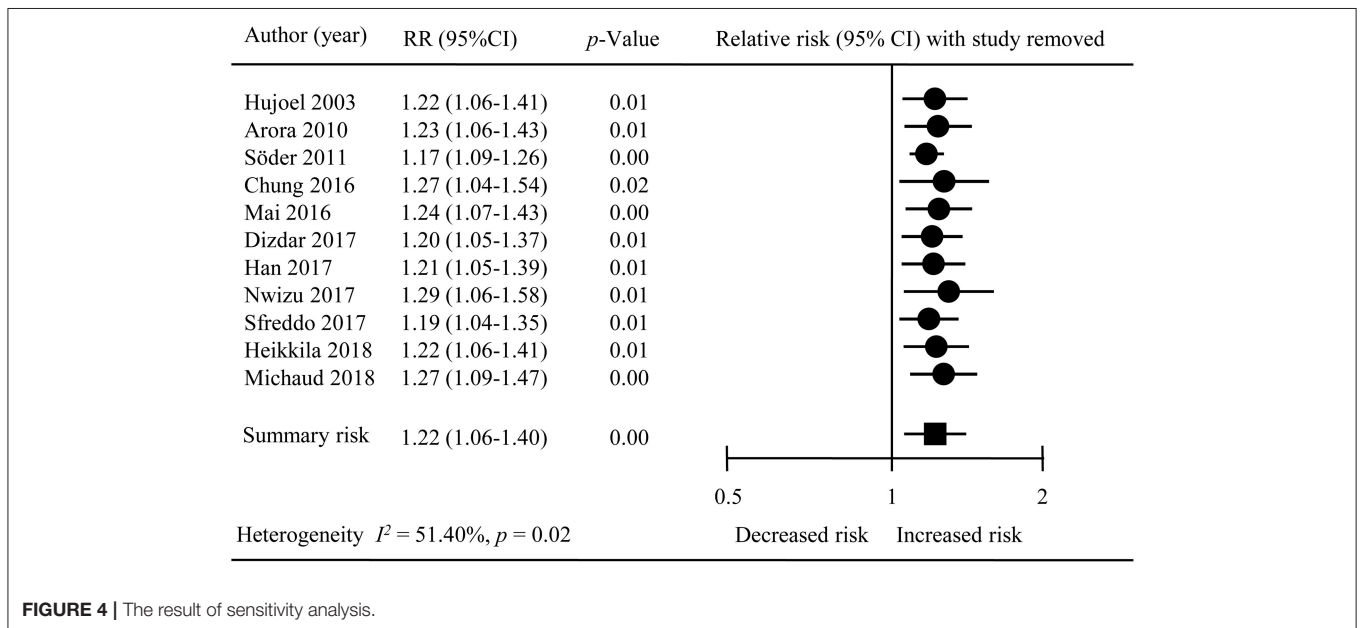
Periodontal disease is a complex infectious diseases (35), and its association with cancer risk has attracted increasing attentions over the years (36, 37). Existing research provided useful insights on the role of periodontal disease treatment in reducing the risk of different types of cancers (38). Possible mechanisms behind the association between periodontal disease and cancers have been proposed, and the most widely advocated theory is the alteration of the oral flora and subsequent influx of the



**TABLE 3** | Results of the overall meta-analysis and subgroup analyses.

Type of analysis	No. of studies	Heterogeneity		Model	Results		
		<i>p</i>	<i>I</i> <sup>2</sup> (%)		RR	95% CI	<i>P</i> -value
<b>Overall analysis</b>	11	0.04	51.40	Random	1.22	1.06–1.40	<0.01
<b>SUBGROUP: STUDY DESIGN</b>							
Prospective cohort	8	0.06	48.22	Random	1.17	0.97–1.41	0.09
Retrospective cohort	2	0.19	40.71	Fixed	1.24	1.12–1.37	<0.01
Case-control	1	-	-	-	2.07	1.13–2.91	0.02
<b>SUBGROUP: FOLLOW-UP PERIOD</b>							
<10 years	3	0.33	8.94	Fixed	1.18	1.10–1.26	<0.01
≥10 years	7	0.02	59.49	Random	1.29	0.95–1.76	0.11
<b>SUBGROUP: MENOPAUSAL STATUS</b>							
Pausimenia	2	0.77	<0.01	Fixed	1.13	1.03–1.23	<0.01
Unspecified	9	0.02	57.01	Random	1.35	1.08–1.68	0.01
<b>SUBGROUP: PERIODONTAL THERAPY</b>							
Included	3	0.40	<0.01	Fixed	1.23	0.95–1.60	0.26
Unreported	8	0.02	59.47	Random	1.25	1.01–1.55	<0.01
<b>SUBGROUP: DENTAL STATUS</b>							
CP	2	0.19	40.71	Fixed	1.24	1.12–1.37	<0.01
PD	9	0.03	52.74	Random	1.24	1.02–1.50	0.03
<b>SUBGROUP: SENSITIVITY ANALYSES</b>							
Excluded studies results reported as ORs	7	0.48	<0.01	Fixed	1.16	1.09–1.23	<0.01
Excluded studies based on co-twin	10	0.01	56.14	Random	1.23	1.06–1.43	0.09
Excluded studies with controls with PD	10	0.03	51.68	Random	1.20	1.05–1.37	0.01

CP, Chronic periodontitis; PD, Periodontal disease; OR, Odds ratio; RR, Relative risk; CI, Confidence interval.

**FIGURE 4** | The result of sensitivity analysis.

inflammatory markers into the systemic circulation (35, 39). Numerous studies suggested that periodontal disease might be a potential risk factor for cancers in the oral, digestive, respiratory and reproductive systems (6–9, 40, 41). Moreover, existing data revealed a prevalence of periodontal disease among females during different time periods, including puberty, menstruation,

pregnancy, and menopause (42–44); and periodontal disease in women has been found to be positively associated with autoimmune diseases, disorders infertility, adverse pregnancy, low birth weight, and preterm birth, and breast cancer (45, 46). Consequently, it has been suggested that oral health amongst females should be a cause for concern.

It is worth highlighting that a similar meta-analysis by Shi et al. (47) was published recently. Though this meta-analysis was conducted with sound methodology and reported clearly according to validated guidelines, our current meta-analysis has a number of strengths. First, the mentioned meta-analysis (47) studies whereas our findings were based on a large evidence base of 11 observational studies. This is due to a more comprehensive search period and study selection approach. Second, Shi et al. (47) only investigated the association between periodontal disease and breast cancer; however, we also specifically investigated the influence of confounding factors such as paucimonia status and history of periodontal therapy on the association between periodontal disease and breast cancer risk. Third, in regards to the calculation of risk estimates, Shi et al. (47) considered HRs/ORs as RRs, whereas we took an extra step to transform ORs into RRs, thereby providing more reliable implications for clinical practice and further researches.

Despite our efforts, however, there are some limitations of our meta-analysis. First, although we were able to identify one (30) contributing to statistical heterogeneity, we were unable to establish the precise causes of such influence. Second, definition of periodontal disease varied between the included studies. Where certain studies used tooth mobility as disease marker, some were self-reported disease status, some used patient medical records for confirmation, and some studies employed clinical or radiographic examinations as final diagnostic status. Third, consideration of confounding factors varied across studies and certain valuable factors such as history of periodontal therapy, baseline smoking status and dental health status, were not consistently reported, and thus findings of the corresponding subgroup analyses remained inconclusive. Fourth, there could be potential control selection bias since we included studies that recruited participants without periodontal disease as controls, as well as one study which adopted national cancer registry figures as the reference (31), although findings from sensitivity analysis revealed robustness of our overall results. Finally, owing to limited data, we were unable to further explore association between various severity of periodontal disease and different subtypes of breast cancer.

## REFERENCES

- Leng WD, Zeng XT, Kwong JS, Hua XP. Periodontal disease and risk of coronary heart disease: an updated meta-analysis of prospective cohort studies. *Int J Cardiol.* (2015) 201:469–72. doi: 10.1016/j.ijcard.2015.07.087
- Chambrone L, Guglielmetti MR, Pannuti CM, Chambrone LA. Evidence grade associating periodontitis to preterm birth and/or low birth weight: I. A systematic review of prospective cohort studies. *J Clin Periodontol.* (2011) 38:795–808. doi: 10.1111/j.1600-051X.2011.01755.x
- Penoni DC, Fidalgo TK, Torres SR, Varela VM, Masterson D, Leao AT, et al. Bone density and clinical periodontal attachment in postmenopausal women: a systematic review and meta-analysis. *J Dent Res.* (2017) 96:261–9. doi: 10.1177/0022034516682017
- Artese HP, Foz AM, Rabelo Mde S, Gomes GH, Orlandi M, Suvan J, et al. Periodontal therapy and systemic inflammation in type 2 diabetes mellitus: a meta-analysis. *PLoS ONE* (2015) 10:e0128344. doi: 10.1371/journal.pone.0128344

Based on currently available evidence, we urge healthcare professionals and the general public to be further aware of the potential association between periodontal disease and breast cancer. The current meta-analysis provided useful insights on the potential effect of periodontal disease on the risk of developing breast cancer, however, further research establishing the precise mechanisms underlying such effect, such as the relationship between altered oral flora, inflammatory marker influx and hormonal levels is warranted. Furthermore, the influences of potential confounding factors, such as smoking status, dietary history, use of contraceptives, marriage, and childbirth status, stress, types of breast cancer, severity of periodontal disease should also be explored.

## CONCLUSION

This meta-analysis indicated that periodontal disease might be a potential risk factor for the development of breast cancer in women, and effective periodontal therapy could pose as a viable preventive measure against breast cancer. However, the association between different severity of periodontal disease and sub-types of breast cancer remain to be ascertained. Nonetheless, we are confident that the findings of the current meta-analysis could be helpful in increasing awareness and importance of oral health maintenance, which may lead to a reduced risk of developing breast cancer.

## AUTHOR CONTRIBUTIONS

JS, LW, and X-TZ designed this study. CF and Y-HJ performed search and collected data. Y-JZ re-checked data. LW, CF, and Y-HJ performed analysis. JS and LW wrote the manuscript. X-TZ reviewed the manuscript.

## ACKNOWLEDGMENTS

We thank Joey S. W. Kwong (JC School of Public Health and Primary Care, Faculty of Medicine, The Chinese University of Hong Kong) for her help with editing and revising our manuscript.

- Mawardi HH, Elbadawi LS, Sonis ST. Current understanding of the relationship between periodontal and systemic diseases. *Saudi Med J.* (2015) 36:150–8. doi: 10.15537/smj.2015.2.9424
- Zeng XT, Deng AP, Li C, Xia LY, Niu YM, Leng WD. Periodontal disease and risk of head and neck cancer: a meta-analysis of observational studies. *PLoS ONE* (2013) 8:e79017. doi: 10.1371/journal.pone.0079017
- Yin XH, Wang YD, Luo H, Zhao K, Huang GL, Luo SY, et al. Association between tooth loss and gastric cancer: a meta-analysis of observational studies. *PLoS ONE* (2016) 11:e0149653. doi: 10.1371/journal.pone.0149653
- Zeng XT, Xia LY, Zhang YG, Li S, Leng WD, Kwong JS. Periodontal disease and incident lung cancer risk: a meta-analysis of cohort studies. *J Periodontol.* (2016) 87:1158–64. doi: 10.1902/jop.2016.150597
- Maisonneuve P, Amar S, Lowenfels AB. Periodontal disease, edentulism, and pancreatic cancer: a meta-analysis. *Ann Oncol.* (2017) 28:985–95. doi: 10.1093/annonc/mdx019
- Ganz PA, Goodwin PJ. Breast cancer survivorship: where are we today? *Adv Exp Med Biol.* (2015) 862:1–8. doi: 10.1007/978-3-319-16366-6\_1



11. Bhatelia K, Singh K, Singh R. TLRs: linking inflammation and breast cancer. *Cell Signal.* (2014) 26:2350–7. doi: 10.1016/j.cellsig.2014.07.035
12. Dartois L, Fagherazzi G, Boutron-Ruault MC, Mesrine S, Clavel-Chapelon F. Association between five lifestyle habits and cancer risk: results from the E3N cohort. *Cancer Prev Res.* (2014) 7:516–25. doi: 10.1158/1940-6207.CAPR-13-0325
13. Mcnamara KM, Moore NL, Hickey TE, Sasano H, Tilley WD. Complexities of androgen receptor signalling in breast cancer. *Endocr Relat Cancer* (2014) 21:T161–181. doi: 10.1530/ERC-14-0243
14. Pinto R, De Summa S, Pilato B, Tommasi S. DNA methylation and miRNAs regulation in hereditary breast cancer: epigenetic changes, players in transcriptional and post-transcriptional regulation in hereditary breast cancer. *Curr Mol Med.* (2014) 14:45–57. doi: 10.2174/1566524013666131203101405
15. Chung SD, Tsai MC, Huang CC, Kao LT, Chen CH. A population-based study on the associations between chronic periodontitis and the risk of cancer. *Int J Clin Oncol.* (2016) 21:219–23. doi: 10.1007/s10147-015-0884-6
16. Freudenheim JL, Genco RJ, Lamonte MJ, Millen AE, Hovey KM, Mai X, et al. Periodontal disease and breast cancer: prospective cohort study of postmenopausal women. *Cancer Epidemiol Biomarkers Prev.* (2016) 25:43–50. doi: 10.1158/1055-9965.EPI-16-0076
17. Mai X, Lamonte MJ, Hovey KM, Freudenheim JL, Andrews CA, Genco RJ, et al. Periodontal disease severity and cancer risk in postmenopausal women: the Buffalo OsteoPerio Study. *Cancer Causes Control* (2016) 27:217–28. doi: 10.1007/s10552-015-0699-9
18. Han MA. Oral health status and behavior among cancer survivors in Korea using nationwide survey. *Int J Environ Res Public Health* (2018) 15:14. doi: 10.3390/ijerph15010014
19. Moher D, Liberati A, Tetzlaff J, Altman DG. Preferred reporting items for systematic reviews and meta-analyses: the PRISMA statement. *PLoS Med.* (2009) 6:e1000097. doi: 10.1371/journal.pmed.1000097
20. Zhang J, Yu KF. What's the relative risk? a method of correcting the odds ratio in cohort studies of common outcomes. *J Am Med Assoc.* (1998) 280:1690–1.
21. Ronskley PE, Brien SE, Turner BJ, Mukamal KJ, Ghali WA. Association of alcohol consumption with selected cardiovascular disease outcomes: a systematic review and meta-analysis. *BMJ* (2011) 342:d671. doi: 10.1136/bmj.d671
22. McNutt LA, Wu C, Xue X, Hafner JP. Estimating the relative risk in cohort studies and clinical trials of common outcomes. *Am J Epidemiol.* (2003) 157:940–3. doi: 10.1093/aje/kwg074
23. Greenland S. Model-based estimation of relative risks and other epidemiologic measures in studies of common outcomes and in case-control studies. *Am J Epidemiol.* (2004) 160:301–5. doi: 10.1093/aje/kwh221
24. Michaud DS, Lu J, Peacock-Villada AY, Barber JR, Joshi CE, Prizment AE, et al. Periodontal disease assessed using clinical dental measurements and cancer risk in the Aric study. *J Natl Cancer Inst.* (2018) 110:843–54. doi: 10.1093/jnci/djx278
25. Higgins JP, Thompson SG, Deeks JJ, Altman DG. Measuring inconsistency in meta-analyses. *BMJ* (2003) 327:557–60. doi: 10.1136/bmj.327.7414.557
26. Egger M, Davey Smith G, Schneider M, Minder C. Bias in meta-analysis detected by a simple, graphical test. *BMJ* (1997) 315:629–34. doi: 10.1136/bmj.315.7109.629
27. Borenstein M, Hedges L, Higgins JP, Rothstein HR. *Comprehensive Meta-Analysis (Version 2.2.027)*. [Computer software] Englewood, NJ: Biostat (2005).
28. Hujuel PP, Drangsholt M, Spiekerman C, Weiss NS. An exploration of the periodontitis-cancer association. *Ann Epidemiol.* (2003) 13:312–6. doi: 10.1016/S1047-2797(02)00425-8
29. Arora M, Weuve J, Fall K, Pedersen NL, Mucci LA. An exploration of shared genetic risk factors between periodontal disease and cancers: a prospective co-twin study. *Am J Epidemiol.* (2010) 171:253–9. doi: 10.1093/aje/kwp340
30. Soder B, Yakob M, Meurman JH, Andersson LC, Klinge B, Soder PO. Periodontal disease may associate with breast cancer. *Breast Cancer Res Treat* (2011) 127:497–502. doi: 10.1007/s10549-010-1221-4
31. Dizdar O, Hayran M, Guven DC, Yilmaz TB, Taheri S, Akman AC, et al. Increased cancer risk in patients with periodontitis. *Curr Med Res Opin.* (2017) 33:2195–200. doi: 10.1080/03007995.2017.1354829
32. Nwizu NN, Marshall JR, Moysich K, Genco RJ, Hovey KM, Mai X, et al. Periodontal disease and incident cancer risk among postmenopausal women: results from the women's health initiative observational cohort. *Cancer Epidemiol Biomarkers Prev.* (2017) 26:1255–65. doi: 10.1158/1055-9965.EPI-17-0212
33. Sfreddo CS, Maier J, De David SC, Susin C, Moreira CHC. Periodontitis and breast cancer: a case-control study. *Community Dent Oral Epidemiol.* (2017) 45:545–51. doi: 10.1111/cdoe.12318
34. Heikkilä P, But A, Sorsa T, Haukka J. Periodontitis and cancer mortality: register-based cohort study of 68,273 adults in 10-year follow-up. *Int J Cancer* (2018) 142:2244–53. doi: 10.1002/ijc.31254
35. Sledziewski TK, Glinka K. Proinflammatory cytokines in periodontal diseases and certain systemic disorders. *Przegl Lek.* (2015) 72:354–7.
36. Michaud DS, Liu Y, Meyer M, Giovannucci E, Joshipura K. Periodontal disease, tooth loss, and cancer risk in male health professionals: a prospective cohort study. *Lancet Oncol.* (2008) 9:550–8. doi: 10.1016/S1470-2045(08)70106-2
37. Fitzpatrick SG, Katz J. The association between periodontal disease and cancer: a review of the literature. *J Dent.* (2010) 38:83–95. doi: 10.1016/j.jdent.2009.10.007
38. Hwang IM, Sun LM, Lin CL, Lee CF, Kao CH. Periodontal disease with treatment reduces subsequent cancer risks. *QJM* (2014) 107:805–12. doi: 10.1093/qjmed/hcu078
39. Mohangi GU, Singh-Rambirich S, Volchansky A. Periodontal disease: mechanisms of infection and inflammation and possible impact on miscellaneous systemic diseases and conditions. *SADJ* (2013) 68:464–467.
40. Michaud DS, Kelsey KT, Papanasiou E, Genco CA, Giovannucci E. Periodontal disease and risk of all cancers among male never smokers: an updated analysis of the health professionals follow-up study. *Ann Oncol.* (2016) 27:941–7. doi: 10.1093/annonc/mdw028
41. Ye L, Jiang Y, Liu W, Tao H. Correlation between periodontal disease and oral cancer risk: a meta-analysis. *J Cancer Res Ther.* (2016) 12:C237–C240. doi: 10.4103/0973-1482.200746
42. Meurman JH, Tarkkila L, Tiitinen A. The menopause and oral health. *Maturitas* (2009) 63:56–62. doi: 10.1016/j.maturitas.2009.02.009
43. Masamatti SS, Kumar A, Virdi MS. Periodontal diseases in children and adolescents: a clinician's perspective part. *Dent Update* (2012)39:551–42. doi: 10.12968/denu.2012.39.9.639
44. Armitage GC. Bi-directional relationship between pregnancy and periodontal disease. *Periodontol* (2013) 2000:160–76. doi: 10.1111/j.1600-0757.2011.00396.x.
45. Lamonte MJ, Hovey KM, Millen AE, Genco RJ, Wactawski-Wende J. Accuracy of self-reported periodontal disease in the women's health initiative observational study. *J Periodontol.* (2014) 85:1006–18. doi: 10.1902/jop.2013.130488
46. Martelli ML, Brandi ML, Martelli M, Nobili P, Medico E, Martelli F. Periodontal disease and women's health. *Curr Med Res Opin.* (2017) 33:1005–15. doi: 10.1080/03007995.2017.1297928
47. Shi T, Min M, Sun C, Zhang Y, Liang M, Sun Y. Periodontal disease and susceptibility to breast cancer: a meta-analysis of observational studies. *J Clin Periodontol.* (2018) 45:1025–33. doi: 10.1111/jcpe.12982

**Conflict of Interest Statement:** The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Copyright © 2018 Shao, Wu, Leng, Fang, Zhu, Jin and Zeng. This is an open-access article distributed under the terms of the Creative Commons Attribution License (CC BY). The use, distribution or reproduction in other forums is permitted, provided the original author(s) and the copyright owner(s) are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.