

Research Article

Periodontitis and Breast Cancer: A Bidirectional Two-Sample Mendelian Randomization Study

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Received: August 29, 2023**Accepted:** September 23, 2023**Published:** September 30, 2023**Abstract**

Background: Observational studies linking periodontitis and breast cancer may be unreliable due to potential reverse causation and confounders.

Methods: A two-sample bidirectional Mendelian Randomization (MR) analysis was conducted. Techniques such as Inverse-Variance Weighted (IVW) analysis, weighted median, weighted mode, simple mode, and MR-Egger regression were utilized. Sensitivity analyses were also executed.

Results: The IVW results indicated no significant genetic link between periodontitis and increased breast cancer risk (OR=1.01, 95% CI: 0.90-1.13, P=0.92). Four other MR methods concurred. Reverse MR analysis also revealed no association. The IVW findings (OR=1.01, 95% CI: 0.90-1.13, P=0.92) were further validated by four additional MR techniques. As a validation step, periodontitis Genome-Wide Association Study (GWAS) data from the FinnGen consortium and breast cancer GWAS data from UKB were used.

Conclusions: The MR study confirms no bidirectional genetic causality between periodontitis and breast cancer. This aids in understanding the relationship between the two conditions and emphasizes further exploration into how periodontitis might relate to other systemic diseases.

Keywords: Breast cancer; Periodontitis; Mendelian randomization

Abbreviations: GWAS: Genome-Wide Association Study; IVW: Inverse Variance Weighted; MR: Mendelian Randomization; UKB: UK Biobank; SNPs: Single-Nucleotide Polymorphisms; IVs: Instrumental Variables; CI: Confidence Interval; OR: Odds Ratio; R²: Coefficient of Determination; RCT: Randomized Controlled Trial; WM: Weighted Median; EUR-UKB: European Ancestry-UK Biobank.

Introduction

As a public health burden and one of the primary causes of tooth loss, periodontitis is a chronic inflammatory condition with a complex multifactorial etiology [1]. In epidemiological investigations that have previously been performed, periodontal disease has been demonstrated to be causally related to an assortment of systemic illnesses in addition to its local effects, such as cardiovascular disease (Zhou, Dong, Zha, & Liao, 2021), rheumatoid arthritis (Bae & Lee, 2020), Parkinson's Disease (Botelho, Machado, Mendes, & Mascarenhas, 2021), depression (Nolde et al., 2022) and diabetes mellitus (Shah, Schooling, & Borrell, 2021). The investigation of the relationship between periodontal disease and the risk of suffering from cancer has become extremely popular. According to published studies,

colorectal, lung, and pancreatic cancers have been associated with the pathogenesis of periodontitis [2-4].

Globally, the risk of contracting breast cancer is the highest among cancers and the second highest in terms of mortality [5]. Numerous risk variables, including endogenous hormone levels, immunological factors, genetic predisposition, and lifestyle choices, have been associated with this cancer according to the data that was accessible. A risk factor for breast cancer has also been associated with periodontal health. Studies that have been published on the possible association between periodontitis and breast cancer have resulted in varying results, with some arguing for a strong association and others arguing against it.

Table 1: Summary of the GWAS included in this Mendelian randomization study.

	Exposures/Outcomes	Consortium	Ethnicity	Sample Sizes	Year
Training set	Periodontitis	GLIDE consortium and UKB	European	17,353 cases;	2019
				28,210 controls	
	Breast cancer	BCAC	European	133,384 cases;	2020
				113,789 controls	
Validation set	Periodontitis	FinnGen R8	European	3837 cases;	2022
				242518 controls	
	Breast cancer	UKB	European	13,879 cases;	2021
				198523 controls	

GWAS: Genome-Wide Association Studies; SNPs: Single Nucleotide Polymorphisms; IVs: Instrumental Variables; GLIDE: Gene-Lifestyle Interactions in Dental Endpoints; UKB: UK Biobank; BCAC: Breast Cancer Association Consortium

A meta-analysis involving eight studies, 168111 individuals confirmed periodontitis did increase susceptibility to breast cancer (RR=1.18, 95%CI: 1.11-1.26, I²=17.6%) [6]. Freudenheim et al. [7] and Sfreddo et al. [8] reported the same conclusion, while Mai et al. [9], Jia et al. [10], and Han et al. [11] reported different observations. Likewise, breast cancer treatment such as radiotherapy, chemotherapy, and endocrine therapy has an impact on periodontal health [12]. These observational studies are generally limited by residual confounders and reverse causality bias. Therefore, studying the possible causal relationship between periodontitis and breast cancer is a valuable tool to improve the treatment of both diseases.

Mendelian Randomization (MR) is a method based on whole genome sequencing data, similar to randomized controlled trial (RCT) studies, and has developed into a multifaceted approach to assessing causal relationships in epidemiology [13,14]. Using genetic variation closely related to exposure as Instrumental Variables (IVs) to evaluate causality and limit bias due to confounders, MR provides reliable insights into the effects of modifiable exposures on traits of interest, transforming phenotype-to-phenotype causality studies into genotypic studies, compared to traditional observational studies [15,16].

The present study is based on the hypothesis that there may be a bidirectional causal relationship between periodontitis and breast cancer. Therefore, two-sample summary data MR analysis was implemented in the study to investigate the possible bidirectional causal relationship between periodontitis and breast cancer.

Materials and Methods

Data Sources and Selection of Genetic Variants

For the validity of each IV, the Single-Nucleotide Polymorphisms (SNPs) used in MR analyses must fulfil three key assumptions [4]:

(1) Relevance assumption: there is robust association between the instrument and the exposure; (2) Independence assumption: the SNP cannot be related to any confounders of the exposure-outcome association; (3) Exclusion restriction assumption: the SNP has to affect the outcome only through the exposure of interest and not through any other pathway. Figure 1 shows a schematic of the Mendelian randomization study of periodontitis and breast cancer.

We performed the two sample MR analysis on the basis of aggregated statistics from the largest available Genome-Wide Association Study (GWAS) on periodontitis, which included 17353 periodontitis cases and 28210 controls [17]. For breast cancer, the data was from the GWAS including 133384 breast cancer cases and 113789 controls [18]. Both GWAS were performed among people of White European descent. As a vali-

date set, the second data of periodontitis were obtained from the FinnGen consortium R8 release data (3837 cases and 242518 controls) [19,20]. The phenotype “periodontitis” was adopted in the current study. The data of breast cancer was from the UKB (EUR-UKB), including 13879 cases and 198523 controls [21]. The summary of the GWAS included in this Mendelian randomization study was in Table 1.

Selection of Genetic Instrumental Variables

Generally, the SNPs were selected at a genome-wide significance of $P < 5 \times 10^{-8}$ to satisfy the first assumption. Considering that few SNPs were associated with periodontitis at the level of $P < 5 \times 10^{-8}$, the genetic instruments with $P < 5 \times 10^{-6}$ were selected. Eight SNPs were proposed as being strongly related with periodontitis based on a significant P -value ($< 5 \times 10^{-6}$ threshold) and were used as IVs [17]. To investigate the causal effect of breast cancer on periodontitis, 12 index SNPs reported significantly associated with breast cancer in a meta-analysis of GWAS on breast cancer ($P < 5 \times 10^{-10}$) were included as candidate genetic instruments [18].

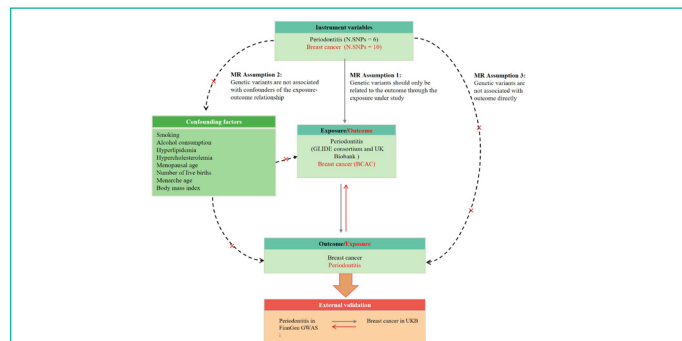


Figure 1: The flowchart of Mendelian randomization analysis. IVW: Inverse Variance Weighted; MR: Mendelian Randomization; SNPs: Single Nucleotide Polymorphisms; IVs: Instrumental Variables; GLIDE: Gene-Lifestyle Interactions in Dental Endpoints; BCAC: Breast Cancer Association Consortium; UKB: UK Biobank

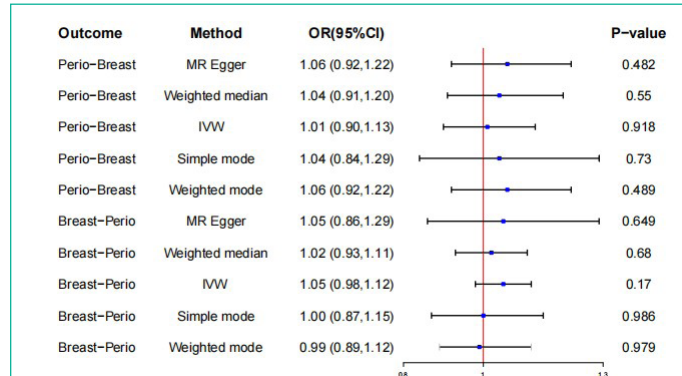


Figure 2: Mendelian randomization estimates for the relationship between genetically instrumented periodontitis and breast cancer, and vice versa. CI: Confidence Interval; IVW: Inverse Variance-Weighted; OR: Odds Ratio.

To exclude SNPs that were in strong Linkage Disequilibrium (LD), we performed the clumping procedure with $R^2 < 0.001$ and a window size = 10,000 kb with the European ancestral individuals from the 1000 Genomes Project to ensure the independence of the instruments [22,23].

Finally, we harmonized the exposure and result datasets to eliminate ambiguous SNPs with non-concordant alleles and SNPs with intermediate allele frequencies in order to effectively ensure that the effect alleles correspond to the same allele. These meticulously selected SNPs served as the final genetic IVs for the ensuing MR analysis.

Furthermore, the F statistics for each SNP have been solely and cumulatively calculated with the following equation: $F = R^2 * (N-2) / (1-R^2)$. R^2 denotes the variance of exposure explained by each IV. IVs with F statistics of less than ten were considered weak instruments and would be excluded for MR analysis [13,24]. An illustration of the Mendelian randomization study of periodontitis and breast cancer is shown in Figure 1.

Statistical Analyses

Multiple statistical methods of MR for causal effect investigation of breast cancer and periodontitis were employed, including the Inverse Variance Weighted (IVW) [25], the Weighted Median (WM) [26], Simple Mode, Weighted Mode, and the MR-Egger [27] methods. Due to its robust causal estimates, IVW is the main method commonly used for two-sample MR studies [25]. WM is the median of the distribution function obtained by sorting all individual SNP effect values according to the weight. The WM can be estimated robustly when at least 50% of the information comes from valid instrumental variables [26]. MR-Egger is a multivariate MR method modified from IVW, assessing potential asymmetry for bias from the pleiotropic effect of the multiple genetic variants and estimating the causal effect [28].

Pleiotropy and Sensitivity Analysis

To assess the effect of risk of bias on the pooled results, we performed the pleiotropy and sensitivity analysis in which we excluded studies with a high risk of bias and assessed whether this changed the results appreciably, using the pleiotropy test, heterogeneity test, and leave-one-out sensitivity test. The intercept term of the MR-Egger regression and the asymmetry of the funnel plot can be utilized for calculating the IVs' average horizontal pleiotropy [29]. MR-Pleiotropy Residual Sum and Outlier (MR-PRESSO) analysis was conducted as well to determine whether pleiotropy was present, correct for horizontal pleiotropy by eliminating the outlier, and deter-

mine whether there are any substantial disparities between the causal effects prior to and following outlier removal [28]. We used the IVW and MR-Egger methods to detect heterogeneity, and a significant P -value (< 0.05) indicates the presence of heterogeneity. In addition, we utilized the leave-one-out analysis to detect the robustness and consistency of the results.

All results are presented as an estimate or Odds Ratio (OR) with a 95% Confidence Interval (CI) of the outcomes OR or per predicted increase/decrease. All statistical tests were two-sided, and the evidence of association was cutoff at a prespecified P -value below 0.05. The open-source statistical software R (version: 4.2.0) was used for all analyses. The TwoSampleMR (version: 0.5.6) [28], MR-PRESSO (version: 1.0), and were mainly used to perform all analyses.

Results

Causal Effects of periodontitis on breast cancer

After removing two SNPs, rs2921075, rs4811024, for being palindromic with intermediate allele frequencies, we incorporated 6 independent SNPs from periodontitis GWAS as IVs for breast cancer. Figure 2 depicts the MR estimations obtained using various different approaches. Overall, no causal associations have been identified between genetically predicted periodontitis and breast cancer risk. The predominant findings of IVW revealed that an increase in the risk of developing periodontitis did not correlate significantly with an increased risk of contracting breast cancer (OR=1.01, 95% CI: 0.90-1.13, $P=0.92$, Table 2). In addition, the MR-Egger, Weighted Median, and Weighted Mode approaches also yield consistent results. Forest plot showing MR results testing for causal relationships between periodontitis and breast cancer (Figure 2). The scatter plot and funnel plots for effect sizes of SNPs for periodontitis and those for breast cancer were shown in Figures 3A and 3B. There was no pleiotropy between periodontitis (MR-Egger regression test, intercept = -0.018, $P=0.305$; Cochran's Q report, $P>0.05$) and breast cancer (Table 2). No high-impact points were found in the leave-one-out analysis (Figures 3C). The F-statistic values were all greater than 10, ranging from 22.257 to 24.303, which indicated that the selection of instrumental variables was effective. The Supplementary Table S1, encompasses detailed information on instrumental variables for periodontitis.

Causal Effects of breast cancer on periodontitis

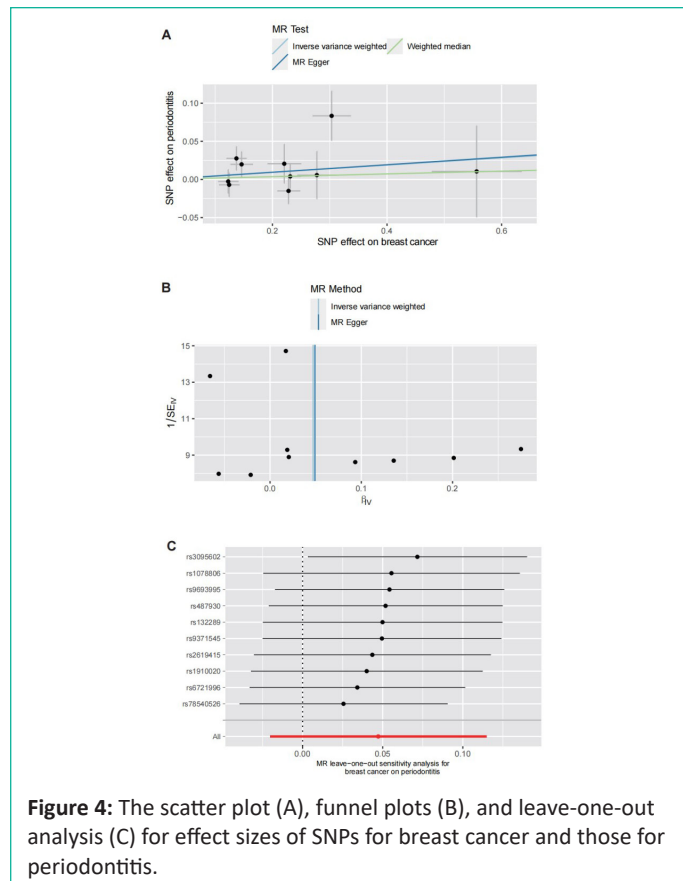
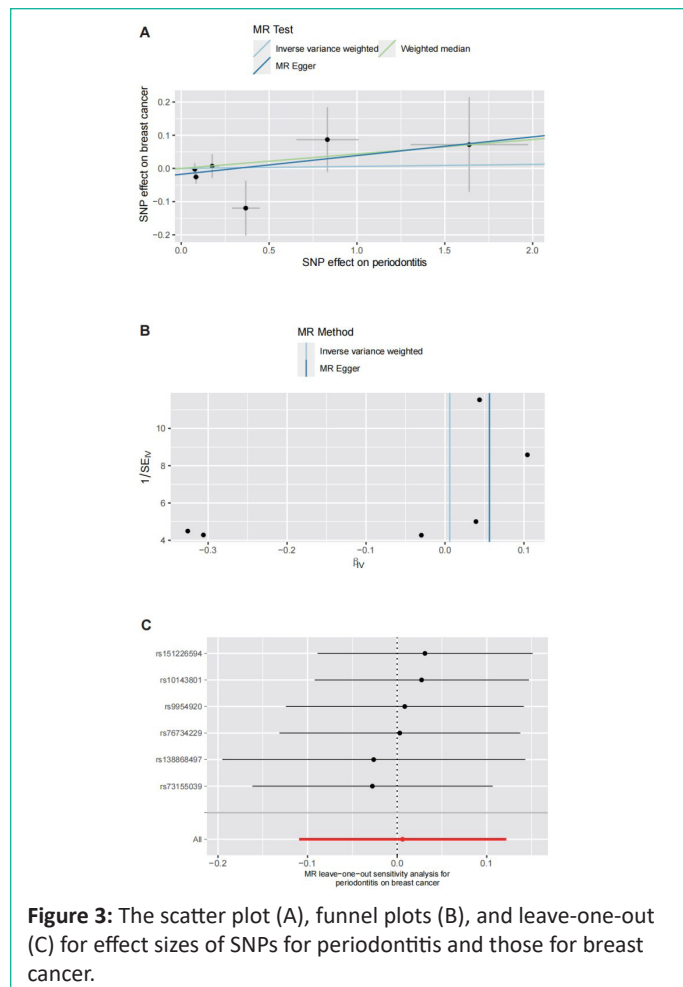
After removing two SNPs, rs10483813, rs10995181, for being palindromic with intermediate allele frequencies, we incorporated 10 independent SNPs from breast cancer GWAS as

Table 2: Mendelian randomization estimates for the relationship between genetically instrumented periodontitis and breast cancer.

Outcome	Exposure	Method	Beta	SE	P value	OR (95% CI)	Cochran's Q			Horizontal pleiotropy		
							Q	Q_df	Q_pval	Egger Intercept	SE	P value
Periodontitis	Breast cancer	MR Egger	0.056	0.073	0.482	1.058(0.917-1.220)	3.579	4	0.466	-0.018	0.015	0.305
		IVW	0.006	0.059	0.918	1.006(0.896-1.129)	4.963	5	0.42			
		Weighted median	0.044	0.071	0.542	1.044(0.908-1.201)						
		Simple mode	0.042	0.108	0.711	1.043(0.845-1.288)						
		Weighted mode	0.055	0.072	0.481	1.057(0.917-1.218)						
Breast cancer	Periodontitis	MR Egger	0.049	0.104	0.649	1.050(0.856-1.289)	10.688	8	0.22	0	0.02	0.984
		IVW	0.047	0.034	0.17	1.048(0.980-1.122)	10.688	9	0.298			
		Weighted median	0.018	0.046	0.69	1.018(0.931-1.114)						
		Simple mode	0.001	0.071	0.987	1.001(0.871-1.151)						
		Weighted mode	-0.002	0.058	0.979	0.998(0.892-1.118)						

MR: Mendelian Randomization; IVW: Inverse Variance Weighted; df: Degree of Freedom; MR: Mendelian Randomization; Q: Heterogeneity Statistic Q.

IVs for periodontitis. The F-statistics ranged between 49.160-168.415, indicating that the selection of instrumental variables was effective (Supplementary file, Table S1). There was no association between breast cancer and periodontitis (OR=1.01, 95% CI: 0.90-1.13, $P=0.92$, Table 2), based on the IVW estimate,



which was supported by the MR-Egger, the Weighted Median, Simple Mode, and the Weighted Mode methods (Table 2, Figure 2). The scatter plot and funnel plots for effect sizes of SNPs for breast cancer and those for periodontitis were shown in Figure 4A and 4B. The leave-one-out analysis did not reveal any leverage points with high influence (Figures 4C). There was no pleiotropy between breast cancer (MR-Egger regression test, intercept = -0.013, $P=0.270$; Cochran's Q report, $P>0.05$) and periodontitis (Table 2).

The validation of bidirectional Mendelian Randomization

The validation using 30 non-overlapping SNPs from the FinnGen consortium R8 release data (19, 20) as periodontitis IVs and 28 SNPs from UKB (21) as breast cancer IVs confirmed the aforementioned results (Supplementary Table S2).

Discussion

In the realm of understanding the complex interplay between chronic inflammatory conditions and cancer susceptibilities, our rigorous two-way MR analysis offers a novel perspective on the potential causal relationship between periodontitis and breast cancer. While previous epidemiological studies painted an inconsistent picture of this relationship, our data-driven approach, grounded in large-scale GWAS datasets, provides a compelling narrative.

It's well-established that periodontitis, a chronic inflammatory disease, has the potential to influence systemic immune and inflammatory responses, primarily through the release of inflammatory mediators [30]. Such chronic inflammation can compromise the immune system's equilibrium, which may subsequently impair tumor cell recognition and elimination, thereby fostering tumor progression [31].

Regarding the association between periodontitis and malignancies, existing observational studies have yielded inconsistent findings. While some studies have reported significant associations between periodontitis and malignancies such as colorectal [32,33], lung [4,34], prostate cancers [35], and pancreatic cancers [36], others have failed to confirm such associations. There is evidence that periodontitis may promote the occurrence and development of breast cancer, and even inflammation of the periodontal tissues promotes breast cancer metastases [37]. In turn, the treatment of breast cancer, radiation therapy, chemotherapy, and endocrine therapy, can affect periodontal health [12]. These conflicting results may be attributed to variations in sample size, study design, population characteristics, and definitions of periodontitis and neoplasia.

The underlying mechanisms influencing the relationship between periodontitis and malignancy are multifaceted and encompass chronic inflammation, the role of bacteria and periodontal pathogens, and alterations in the immune system [38,39]. A particularly intriguing facet of this relationship is the role of the oral microbiome. Disruptions in the oral microbiota have been spotlighted as risk factors for a plethora of diseases, including gastrointestinal cancers, lung cancer, and notably, breast cancer [40,41]. *Fusobacterium nucleatum*, an oral bacterium, has emerged as a noteworthy player, being linked with various tumor types [42]. Its proposed mechanisms span from modulating the tumor microenvironment to intricate interactions with host cells and inducing host inflammation. Although our results, consistent across multiple MR methods, suggest that the genetic underpinnings of periodontitis may not have a direct causal influence on breast cancer risk, the alterations

in the oral microbiome remain a tantalizing area warranting deeper exploration.

Our revelations seem to diverge from some observational studies that have asserted a causal interplay between periodontitis and breast cancer [6,8,10,43-45]. Confounding and reverse causation bias are two fundamental drawbacks of observational research that ought to be acknowledged. In particular, age, lifestyle choices, and underlying genetic predisposition constitute common risk factors between periodontitis and breast cancer, which may obscure relationships discovered in observational research. In observational studies, where the existence of breast cancer may influence changes in oral health and the emergence of periodontitis, reverse causality can also be an issue that might arise. Contrarily, confounding and reverse causation biases, which can be prevalent in observational research, have been effectively eliminated by the MR analysis employed in this research. MR analysis is able to produce far more precise causal estimates for investigating the potential causal relationship between periodontitis and breast cancer through the utilization of genetic variation as a proxy for exposure. Nevertheless, it is crucial to reiterate that although our results utilize a robust causal inference technique, it is not sufficient to completely rule out the possibility of an association between periodontitis and breast cancer.

Despite the beneficial effects of MR analysis in resolving confounding variables and reverse causation, it is contingent on certain presumptions that could result in potential constraints to our investigation. As such, it is contingent upon the availability of genetic tools, for instance, and may not be able to adequately account for all genetic heterogeneity. Although we conducted sensitivity analyses and employed MR-Egger and MR-PRESSO tests to detect genetic heterogeneity effects, the possibility of unmeasured genetic heterogeneity cannot be entirely ruled out. Additionally, the GWAS data utilized in our analysis predominantly represented European Caucasians, and the genetic structure and underlying biological mechanisms may vary across different populations. Therefore, the generalizability of our findings to other populations with distinct genetic backgrounds necessitates further investigation.

Although our two-way MR analysis did not provide evidence supporting a causal relationship between periodontitis and breast cancer, the insights gained from this investigation shed light on common risk factors and potential biological pathways shared between these two conditions. Understanding the interactions between periodontitis and breast cancer could inform public health interventions and personalized treatment strategies. Future studies should continue to explore the underlying biological mechanisms involving genetic factors, inflammation, and cancer risk, while also encompassing diverse populations to attain a more comprehensive understanding of the association between periodontitis and breast cancer.

Conclusions

Our bidirectional two-sample MR analysis doesn't buttress a direct causal link between periodontitis and breast cancer. These findings punctuate the ongoing dialogue surrounding periodontitis and breast cancer, emphasizing the intricate choreography between oral health, the microbiome, and systemic maladies like cancer. This underscores the imperative for a more integrative healthcare paradigm, recognizing the symbiosis of seemingly discrete physiological systems.

Author Statements

Conflict of Interest

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.

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Data Availability Statement

The datasets analyzed during the current study are available from <https://data.bris.ac.uk/data/dataset/2j2rqgzdxlq02oqbb4vmycnc2> for periodontitis GWAS, <https://bcac.ccge.medschl.cam.ac.uk/bcacdata/> for breast cancer GWAS, gs://finngen-public-data-r8/summary_stats/finngen_R8_K11_PERIODON_CHRON.gz for validation of periodontitis GWAS and <https://gwas.mrcieu.ac.uk/datasets/ieu-b-4810/> for validation of breast cancer GWAS.

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