

## Relationship between oral health and glaucoma traits in the United Kingdom

### Short title: Oral health and glaucoma

Rachel H Lee, MD, MPH,<sup>1</sup> Jae H Kang, ScD,<sup>2</sup> Janey L Wiggs, MD, PhD,<sup>3</sup> Siegfried K Wagner, BM, BCh, MA, FRCOphth,<sup>4</sup> Anthony P Khawaja, MB BS, MRCOphth,<sup>5</sup> Louis R Pasquale, MD,<sup>1,6</sup> for the Modifiable Risk Factors for Glaucoma Collaboration, the UK Biobank Eye and Vision Consortium, and the International Glaucoma Genetics Consortium

<sup>1</sup>New York Eye and Ear Infirmary of Mount Sinai, New York, NY, USA

<sup>2</sup>Channing Division of Network Medicine, Brigham and Women's Hospital, Harvard Medical School, Boston, MA, USA

<sup>3</sup>Department of Ophthalmology, Massachusetts Eye and Ear, Harvard Medical School, Boston, MA, USA

<sup>4</sup>NIHR Birmingham Biomedical Research Centre, University Hospital Birmingham and University of Birmingham, Birmingham, UK

<sup>5</sup>NIHR Biomedical Research Centre, Moorfields Eye Hospital NHS Foundation Trust & UCL Institute of Ophthalmology, London, UK

<sup>6</sup>Department of Ophthalmology, Icahn School of Medicine at Mount Sinai, New York, NY, USA

Address for reprints:

Icahn School of Medicine, Mount Sinai  
1 Gustave L. Levy Place, NY, NY 10029  
Fax: 212-241-5764

This work was funded by the American Glaucoma Society Mentoring for Advancement of Physician Scientists grant, the Eye and Vision Research Institute of Mount Sinai, The Glaucoma Foundation (NYC), Research to Prevent Blindness (NYC), and NIH R01 EY015473 and EY032559.

COIs: JHK received research support from Pfizer; JLW is a consultant for Editas, CRISPR Therapeutics, Broadwing Bio, Regenxbio; LRP is a consultant to Twenty Twenty, Character Bio; RHI, SKW, APK- none

## **Precis:**

In this cross-sectional analysis of UK Biobank participants, we find no adverse association between self-reported oral health conditions and either glaucoma or elevated intraocular pressures.

## **Abstract:**

### *Purpose*

Poor oral health may cause inflammation that accelerates the progression of neurodegenerative diseases. We investigated the relationship between oral health and glaucoma.

### *Patients*

United Kingdom (UK) Biobank participants

### *Methods*

This is a cross-sectional analysis of participants categorized by self-reported oral health status. Multivariable linear and logistic regression models were employed. Primary analysis examined the association with glaucoma prevalence. Secondary analyses examined associations with IOP, macular retinal nerve fiber layer (mRNFL), and ganglion cell inner plexiform layer (mGCIPL) thicknesses, and interaction terms with multi-trait glaucoma polygenic risk scores (MTAG PRS) or intraocular pressure (IOP) PRS.

### *Results*

170,815 participants (34.3%) reported current oral health problems, including painful or bleeding gums, toothache, loose teeth, and/or denture wear. 33,059, 33,004, 14,652, and 14,613 participants were available for analysis of glaucoma, IOP, mRNFL, and mGCIPL, respectively. No association between oral health and glaucoma was identified (odds ratio (OR): 1.04, 95% confidence interval (CI): 0.95, 1.14). IOPs were slightly lower among those with oral disease (-0.08 mmHg, 95% CI: -0.15, -0.009); specifically, among those with loose teeth ( $p=0.03$ ) and denture-wearers ( $p<0.0001$ ). mRNFL measurements were lower among those with oral health conditions (-0.14 microns, 95% CI: -0.27, -0.0009), but mGCIPL measurements ( $p=0.96$ ) were not significantly different. A PRS for IOP or glaucoma did not modify relations between oral health and IOP nor glaucoma ( $p\text{-for-interactions}\geq 0.17$ ).

### *Conclusions*

Self-reported oral health was not associated with elevated IOP nor increased risk of glaucoma. Future studies should confirm the null association between clinically diagnosed oral health conditions and glaucoma.

**Keywords:** oral health; dental health; glaucoma; intraocular pressure

## I. Introduction

A growing body of research suggests that oral infections may lead to chronic inflammation of distant tissues. Poor oral health has been linked to a variety of systemic diseases, including heart disease, dementia, diabetes, chronic kidney disease, rheumatoid arthritis, and several malignancies.<sup>1</sup> Two possible mechanisms for these associations have been described previously. First, it is postulated that chronic periodontitis could allow oral bacteria to enter the systemic circulation, thereby enabling damage to distant organs,<sup>1</sup> including cardiovascular tissues. Additionally, chronic periodontitis may serve as a source of chronic inflammation, thus accelerating other disease processes,<sup>2,3</sup> including neurodegenerative conditions such as Alzheimer's disease.<sup>4</sup>

Glaucoma is a chronic neurodegenerative disease that may lead to permanent vision loss and blindness. To date, intraocular pressure (IOP) is the only known modifiable risk factor for glaucoma. Thus, all medical and surgical interventions for glaucoma aim to lower IOP. The discovery of additional modifiable risk factors for glaucoma would therefore be significant. Several studies suggest there is a relationship between oral health and glaucoma. A prospective cohort study conducted using data from the Health Professionals Follow-up Study found that a history of tooth loss during the preceding two years was associated with a 1.5-fold increase in the risk of primary open-angle glaucoma.<sup>5</sup> A study based on Taiwan's National Health Insurance Research Database reported that patients with a history of periodontitis were found to have a 30% increased risk of developing glaucoma as compared to those without dental disease.<sup>6</sup> In line with these findings, a cross-sectional study conducted in Korea noted over a three-fold increase in odds of glaucoma among patients with a history of periodontitis.<sup>7</sup> Although these initial studies are somewhat consistent, conclusions on the specific types of oral diseases associated with glaucoma vary from study to study. Furthermore, it is unknown whether individuals with poor oral health have higher IOPs.

Importantly, the mechanism underlying the potential link between oral health and glaucoma is also not fully understood. Previously, two studies investigated the possibility that dental health might affect the oral microbiome, and thereby increase the risk of glaucoma. While one study found that an increased total oral bacterial load was associated with glaucoma,<sup>8</sup> the other suggested that the prevalence of specific bacterial strains contributed to glaucomatous disease.<sup>9</sup> Therefore, while changes in the oral microbiome may be associated with an increased risk of glaucoma, an alternative and/or complementary mechanism to explain a possible link between poor oral health and glaucoma is also plausible.

The goal of this study was to investigate a possible relationship between poor oral health and glaucoma by leveraging data from the UK Biobank. We hypothesized that poor oral health may be linked to an altered metabolome,<sup>10,11</sup> that could lead to an upregulated inflammatory response, impaired microvascular flow, and ultimately elevated IOP and/or glaucoma. Therefore, we examined the association between poor oral health and the prevalence of glaucoma in the UK Biobank. We also investigated the relationship between oral health and intraocular pressures and ocular coherence tomography measurements.

## II. Methods

This study was exempt from review by the Mount Sinai Hospital Institutional Review Board due to the utilization of de-identified data. The North West Multi-Center Research Ethics Committee approved the study, according to the principles of the Declaration of Helsinki and the Icahn School of Medicine has entered into a data use agreement with the UK Biobank under UK Biobank application number 36741 for the use of the de-identified data files.

### A. Design

The UK Biobank is a cohort study including 502,389 adults recruited between 2006 and 2010. Participants completed questionnaires, in-person interviews, physical measurements, and the collection of biospecimens, including blood, urine, and saliva samples. Consent was obtained to link baseline data to health records. The UK Biobank, therefore, includes data on participant demographics, self-reported and clinician-confirmed health status and diseases, biometric measurements including IOPs, as well as genomes, proteomes, and metabolomes.

### B. Participants

Subjects included for analysis were between the ages of 37 and 73 years at the time of recruitment. Those participants missing data on oral health or ophthalmic measurements/glaucoma disease status were excluded from the analysis. Those participants with missing covariable data were excluded from the analysis, given small rates of missing data across the majority of covariables (0-3.5%). Data on metabolic equivalents (METS) and calorie intake were only available for ~80% and 14% of participants, respectively, but were included in analyses.

### C. Oral health and other exposure variables

Oral health history was ascertained based on data from baseline health questionnaires. Subjects were presented with a touchscreen question, “Do you have any of the following? (You can select more than one answer).” The choices included the presence of mouth ulcers, painful gums, bleeding gums, loose teeth, or toothache, none of the above, or prefer not to answer. Prior studies have suggested that bleeding and/or painful gums, and tooth mobility have acceptable validity in detecting moderate to severe periodontitis; self-reported toothache has validity in identifying patients with periodontitis and pulpitis.<sup>12–15</sup> Because periodontal disease is the leading cause of tooth loss leading to denture wear, and self-reported denture wear has been associated with excellent validity and reliability,<sup>16,17</sup> this variable was also included for analysis. Therefore, for this study, the presence of painful or bleeding gums, loose teeth, toothache, or denture wear was analyzed both as a composite exposure variable for ‘poor oral health’ and as individual exposures. Participants without oral health issues were defined by the lack of self-reported toothache, bleeding or painful gums, tooth loss, or denture wear.

Baseline demographic characteristics including age, sex, and self-reported ethnicity, were recorded. The Townsend deprivation index, a single numerical value quantifying material deprivation including unemployment, lack of car or home ownership, and household overcrowding,<sup>18</sup> was also documented for each subject. Additionally, data on multiple covariables including diagnosis of diabetes, alcohol use, smoking history, systolic blood pressure, body mass index, estimated caloric intake, and physical activity levels (as estimated via

metabolic equivalents in hours per week, or METS), spherical equivalents, and use of systemic beta-blockers were extracted and derived from the database for analysis.

#### *D. Outcome variables*

Ophthalmic data was obtained from 122,143 participants in 2009 and 2010 at UK Biobank assessment centers. A single IOP measurement was recorded for each eye using the Reichert Ocular Response Analyzer noncontact tonometer. Corneal-compensated IOP measurements of the right and left eyes were averaged to calculate a subject-level outcome. Subjects were excluded from analysis if they reported a history of eye surgery within four weeks of the measurement, or if they reported an eye infection at the time of measurement. The lowest and highest 0.5% of measurements were discarded to minimize bias due to artefactual extreme measures. For those patients on IOP-lowering therapy, pretreatment IOPs were imputed by dividing the measured IOP by 0.7.<sup>19–21</sup> Patients with a history of glaucoma surgery and laser were excluded from the analysis.

Baseline health questionnaires included the query, “Has a doctor told you that you have any of the following problems with your eyes?” Study participants were categorized as having self-reported glaucoma if the response “glaucoma” was selected for this query. Patients whose health records also revealed an associated ICD-9 or ICD-10 diagnosis code of open-angle glaucoma were also categorized as having diagnosis-confirmed glaucoma (ICD-9 codes 365.0\*, 365.1\*, 365.7\*; ICD-10 codes H40.1\*).

Retinal ocular coherence tomography (OCT) measurements in the macula region were obtained in 67,321 subjects. High-resolution spectral domain OCT images of undilated nerves and retinas were performed using the Topcon 3D OCT 1000 Mk2. Quality control steps for included OCT scans have been previously described.<sup>22,23</sup> In brief, scans with poor signal strength, and/or those with scan quality or segmentation indices in the bottom 20% of all images were also excluded.

#### *E. Genetic data, Glaucoma, and IOP polygenic risk scores*

Genotyping data were obtained on 488,377 subjects using Affymetrix UK BiLEVE Axiom Array (49,950 participants) and the Affymetrix UK Biobank Axiom Array (438,427 participants). As described previously, quality control and imputation were performed jointly, as the two arrays shared over 95% of genetic markers.<sup>24</sup> In total, 92,693,895 genetic markers of 487,442 participants were made available for analysis in the UK Biobank database.

Data from genome-wide association studies (GWAS) of individuals of European descent were used to create a multi-trait glaucoma polygenic risk score (MTAG PRS) for each patient, consisting of 2,673 independent genetic loci.<sup>25</sup> We also created an IOP PRS consisting of 111 independent genome-wide significant loci based on results from the largest IOP GWAS to date.<sup>20</sup> Each PRS served as a single numeric score that summarizes the genetic risk for POAG for each subject. The methods for creating a glaucoma PRS has been described previously.<sup>20,25</sup>

Additional analyses including glaucoma PRS as a covariable and evaluating whether a glaucoma PRS modifies the relationship between oral health and glaucoma were evaluated. In addition to treating the PRS as a continuous variable, we conducted a sensitivity analysis where we classified participants into two genetic risk groups: those with the highest 25% genetic risk scores and those in the lowest 25% genetic risk scores. The remaining 50% of participants were

excluded from the sensitivity analysis. Interaction terms between categorical PRS variables and oral health were calculated, and used to determine whether extremes of genetic risk scores modified the relationship between oral health and glaucoma.

### *F. Statistical analysis*

Baseline characteristics were compared among subjects in the poor oral health group versus the comparison group using the student's T-test and 1-proportion Z-test. To evaluate associations with poor oral health as a risk factor for glaucoma, multiple logistic regression models were used, adjusting for multiple covariables extracted from baseline health survey questionnaire data and measurements: age, sex, self-reported ethnicity, smoking history, alcohol use, physical activity, Townsend deprivation index, BMI (kilograms per square meter), systolic blood pressure, diabetes, and total calorie intake.<sup>26</sup> Similarly, to determine whether poor oral health is associated with IOPs or differences in macula region retinal nerve fiber layer thickness (mRNFL) or ganglion cell inner plexiform layer (mGCIPL) thickness, multiple linear regression models were conducted, adjusting for the same covariables. Finally, for each glaucoma trait, we examined whether a glaucoma or IOP PRS modified the relation between oral health and the outcomes of interest by evaluating interaction variables (oral health variable \* genetic variable). The significance of the interaction term was assessed with a p-for interaction test statistic. Since total caloric intake and physical activity had a high missingness rate, a sensitivity analysis excluding these covariates was performed.

All statistical analyses were conducted using SAS 9.4 and R software.

## **III. Results**

### *A. Demographics*

A total of 498,713 subjects completed the oral health questionnaire. Of these, a total of 170,815 (34%) subjects reported a history of oral health conditions including bleeding or painful gums, toothache or loss, or need for dentures. Please see **Figure 1** for details on subject inclusion details.

Subjects with a history of self-reported oral health conditions were older, more likely to be female, of non-Caucasian descent, and were more likely to experience material deprivation than those without a history of oral health conditions. Additionally, those with self-reported oral health conditions were more likely to have a history of diabetes, reported consuming less alcohol, were more likely to report smoking, and had higher BMIs than those without oral health problems. Those with oral health conditions were also more likely to have increased calorie intake and were less sedentary than those without oral health conditions (**Table 1**). We adjusted for all these covariables in multivariable analysis.

### *B. Association with glaucoma prevalence*

A total of 33,059 subjects had data available for all covariables for analysis on the relationship between oral health and glaucoma. A total of 4,801 subjects (2.81%) with self-reported oral health conditions and 6,842 subjects (2.30%) without oral health conditions also had a history of

glaucoma at the time of survey collection. A statistically significant difference in the proportion of patients with glaucoma was noted in univariate analysis ( $p < 0.0001$ ).

After controlling for covariables, oral health conditions were not associated with an increased likelihood of glaucoma (**Table 2**). The overall odds of glaucoma among subjects with oral health conditions as compared to those without in this cohort was 1.04 (95% confidence interval (CI): 0.95, 1.14;  $p = 0.39$ ). The relationship between each oral health variable and glaucoma was also analyzed. None of the five oral health variables—the presence of painful gums, bleeding gums, toothache, loose teeth, or denture wear—were associated with increased odds of glaucoma (Supplemental Table 1, Supplemental Digital Content 1, <http://links.lww.com/IJG/A872>).

Sensitivity analysis excluding variables for physical activity and caloric intake showed that oral health problems resulted in no material differences (data not shown).

Each standard deviation increase in PRS was associated with a nearly 2.5-fold increased odds of glaucoma (odds ratio: 2.48, 95% confidence interval: 2.34, 2.63,  $p < 0.0001$ ). Overall, the MTAG PRS did not modify the relationship between oral health problems and glaucoma ( $p$  for interaction = 0.84; **Table 2**). Furthermore, no significant modification was noted for any of the five oral health variables in relation to glaucoma ( $p$  for interaction  $\geq 0.34$ , Supplemental Table 1, Supplemental Digital Content 1, <http://links.lww.com/IJG/A872>). Finally, no significant interaction between dental health problems and MTAG PRS were noted when only considering those in the lowest vs highest quartile of glaucoma risk ( $p = 0.98$ , Supplemental Table 2, Supplemental Digital Content 2, <http://links.lww.com/IJG/A873>).

### C. Association of oral health conditions with IOP

A total of 33,004 subjects had IOP data available for analysis. The average IOP among patients without oral health problems was  $16.0 \pm 3.3$  mmHg, while the average IOP among patients with oral health conditions was  $15.9 \pm 3.3$  mmHg ( $p = 0.03$  on univariate analysis). Self-reported oral health conditions were associated with a small, but statistically significant lower IOP, after adjusting for multiple covariables ( $p = 0.03$ , **Table 3**).

The relationship between each oral health variable with IOP was assessed. We found that participants reporting loose teeth or wearing dentures had lower IOPs than those without loose teeth ( $p = 0.03$ ) or without dentures ( $p < 0.0001$ , Supplemental Table 3, Supplemental Digital Content 3, <http://links.lww.com/IJG/A874>). By contrast, painful or bleeding gums, toothache, and loose teeth were not associated with IOP differences ( $p \geq 0.09$ ; Supplemental Table 3, Supplemental Digital Content 3, <http://links.lww.com/IJG/A874>). Sensitivity analysis excluding variables for physical activity and caloric intake showed that oral health problems resulted in no differences to the results described above (data not shown).

Every point increase in standardized IOP PRS was associated with a 0.72 mmHg increase in IOP ( $p < 0.0001$ ). The interaction between IOP PRS and oral health conditions was neither significant among patients across the spectrum of IOP PRS (**Table 3**;  $p = 0.37$ ), nor among those of the top 25% genetic risk score as compared to the lowest 25% risk scores ( $p = 0.32$ , Supplemental table 4, Supplemental Digital Content 4, <http://links.lww.com/IJG/A875>). Interaction terms with each of the individual oral health variables were also not statistically significant ( $p$ -for-interaction  $\geq 0.10$ ; Supplemental table 3, Supplemental Digital Content 3, <http://links.lww.com/IJG/A874>).

### D. Association of oral health conditions and macula region retinal nerve fiber layer (mRNFL) and macula region ganglion cell inner plexiform layer (mGCIPL)



14,652 and 14,613 subjects had mRNFL and mGCIPL data available for analysis, respectively. The average mRNFL thickness was  $28.0 \pm 3.8$  microns among subjects with oral health problems and  $28.4 \pm 3.8$  microns among those without oral health problems ( $p=0.38$  on univariate analysis). The average mGCIPL thickness was  $74.5 \pm 5.2$  microns among subjects with oral health problems and  $74.7 \pm 5.1$  microns among those without oral health problems ( $p=0.14$  on univariate analysis). There was an inverse association between self-reported oral health conditions and mRNFL thickness, after adjusting for multiple covariables ( $-0.14$  microns,  $p=0.04$ , **Table 4**), but no association between oral health conditions and mGCIPL thickness ( $p=0.96$ , **Table 5**).

The MTAG PRS did not modify the relationship between oral health conditions and macula inner retinal parameters (mRNFL and mGCIPL) ( $p$  for interaction  $\geq 0.17$ ; **Tables 4 and 5**). The interaction between the MTAG PRS and oral health conditions was weakly significant among those of the top 25% of MTAG PRS as compared to the lowest 25% risk scores for mGCIPL thickness (0.48 microns;  $p$  for interaction= $0.04$ ; Supplemental Table 4, Supplemental Digital Content 4, <http://links.lww.com/IJG/A875>).

Bleeding gums were associated with a small, but statistically significant decrease in mRNFL thickness ( $-0.21$  microns,  $p=0.02$ , Supplemental Table 5, Supplemental Digital Content 5, <http://links.lww.com/IJG/A876>). All other individual oral health problems were not associated with mRNFL nor mGCIPL ( $p \geq 0.13$ ; Supplemental Tables 5, Supplemental Digital Content 5, <http://links.lww.com/IJG/A876> and 6, Supplemental Digital Content 6, <http://links.lww.com/IJG/A877>). The MTAG PRS modified the relationship between toothache and mRNFL ( $p=0.003$  for interaction) despite a null primary relationship between toothache and mRNFL ( $p=0.14$ , Supplemental Table 5, Supplemental Digital Content 5, <http://links.lww.com/IJG/A876>). The MTAG PRS did not modify the associations between all other oral health variables and mRNFL nor mGCIPL (Supplemental Tables 5, Supplemental Digital Content 5, <http://links.lww.com/IJG/A876> and 6, Supplemental Digital Content 6, <http://links.lww.com/IJG/A877>). Sensitivity analysis excluding variables for physical activity and calorie intake revealed that oral health problems were associated with a small  $-0.11 \pm 0.04$  microns ( $p=0.01$ ) decrease in mRNFL thickness and  $-0.13 \pm 0.05$  microns ( $p=0.01$ ) decrease in mGCIPL thickness. Among the individual oral health variables, dentures were associated with slightly thinner mRNFL ( $p=0.01$ ) and mGCIPL ( $p=0.005$ ) thicknesses, and bleeding gums were associated with thinner mRNFL thickness ( $p=0.03$ , data not shown).

#### IV. Discussion

In this cross-sectional study of over 500,000 participants of the UK Biobank, we investigated a possible relationship between oral health and glaucoma. Consistent with prior reports outside the UK Biobank Study, over one-third of the study population reported some history of oral health conditions including loose teeth, use of dentures, and tooth or gum pain.<sup>27</sup> After controlling for multiple covariables, we found that self-reported oral health conditions were not associated with increased odds of glaucoma (odds ratio: 1.04;  $p=0.39$ ).

Oral health problems were weakly associated with lower IOP (**Table 3**;  $-0.08$  mmHg,  $p=0.03$ ). We found that this difference in IOP was driven by an association between both loose teeth and denture wear with IOP (Supplemental Table 3, Supplemental Digital Content 3, <http://links.lww.com/IJG/A874>;  $-0.13$  mmHg,  $p=0.03$ , and  $-0.22$  mmHg,

p<0.0001, respectively). Self-reported oral health conditions were also associated with a small decrease in mRNFL thickness (**Table 4**; -0.14 microns, p=0.04), but this difference in mRNFL was not consistently reproducible among the other component oral health variables. Furthermore, oral health conditions did not appear to affect the risk of glaucoma or elevated IOP among patients with low versus high genetic risk for glaucoma or elevated IOP. Thus, while oral health conditions may result in a small decrease in intraocular pressures, this is not protective against glaucoma. Additionally, oral health problems may be associated with thinning of inner retinal layers in a pressure-independent mechanism.

While prior studies suggest that recent loose teeth or periodontal disease is associated with an increased risk of primary open-angle glaucoma,<sup>5-7</sup> we suspect that these results likely reflect variations in the study population, covariable inclusion, and variable definitions. One study examined subjects who were older, male, and had more stringent glaucoma diagnosis criteria, including a review of visual fields.<sup>5</sup> That study reported that tooth loss within the last two years was associated with an increased risk of glaucoma. We found no association between loose teeth and glaucoma in this cross-sectional study—these differences may be attributed to differences in data collection. For instance, while patients in the UK Biobank reported a history of loose teeth or denture wear, data on the recency of tooth loss or denture requirement were not collected. Two other studies implicated an association between periodontal disease and glaucoma: one study examined a Taiwanese national health database without access to covariables including smoking, alcohol use, physical activity, and diet,<sup>6</sup> while a third study leveraged data from a Korean national health database.<sup>7</sup> Among our younger European population, we find that after accounting for multiple covariables, there is no consistent association between self-reported oral health problems and glaucoma in cross-sectional analysis.

The results here expand upon a prior study conducted by Lehrer *et al* on the association between dental disease and glaucoma utilizing the UK Biobank database.<sup>28</sup> In their work, Lehrer *et al* report that the presence of bleeding gums was associated with a decreased risk of primary open-angle glaucoma, and with lower IOP. Discrepancies in our results are likely related to differences in variable definitions and covariable inclusion. In their study, Lehrer *et al* defined glaucoma by the ICD-10 codes for primary open-angle glaucoma alone. In the UK Biobank, ICD-10 codes for POAG are recorded among individuals who undergo a procedure; therefore, the inclusion of individuals with an ICD-10 diagnosis of POAG likely excludes a significant number of patients with glaucoma. In our work, we defined glaucoma not only by the associated ICD-10 codes for glaucoma but also by self-reported glaucoma or the usage of glaucoma medications. Additionally, while the multivariate analysis conducted by Lehrer *et al* included age, gender, diabetes, and smoking history as covariables, our analysis includes a more comprehensive list of known risk factors for glaucoma, including ethnicity and component genetic PRS, among others. Finally, we examined the relationship between oral health conditions and glaucoma in detail, by also using imaging proxies including mRNFL and mGCIPL data. We found inconsistent relations between oral health problems and inner retinal biomarkers – participants with oral health problems had lower mRNFL but the relationship between oral health problems and mGCIPL was null.

The strengths of our study include the large sample available through the UK Biobank, a comprehensive collection of covariable and genetic data, availability of imaging data, and consistency with prior studies. Our results are supported by data collected previously on dental health in the United Kingdom outside of the UK Biobank. For instance, in line with data collected from the National Dental Public Health Team, we found that roughly 16% of

participants wear dentures. Similarly, we found that over 4.4% of participants reported loose teeth in the UK Biobank, comparable to the percentage of adults reporting missing anterior teeth according to data from the National Dental Public Health Team (7.6%), providing further indirect validation for the touchscreen questionnaire instrument used to assess oral health.<sup>29</sup> Furthermore, our univariate analysis suggests that oral health conditions may be more prevalent among patients with smoking history or diabetes, consistent with prior studies that link smoking to periodontitis,<sup>30–32</sup> and those that suggest a relationship between diabetes and oral health disease.<sup>33–36</sup> Additionally, despite the inclusion of self-reported glaucoma, we still found strong correlations between glaucoma PRS scores and patients with self-reported glaucoma. Finally, we did explore whether associations between oral health and glaucoma-related outcomes were modified by a genetic predisposition to higher IOP or glaucoma. Overall, our analyses revealed the minimal impact of our PRSs on the relationship between oral health problems and glaucoma traits.

This work is limited by the reliance upon self-reported oral disease. History of prior dental or oral problems may have been affected by recall bias by participants or misunderstandings of true oral health status. Our study likely underestimates the prevalence of oral health problems, particularly among those individuals with prior or minimal symptoms. Additionally, while our data are consistent with estimated rates of oral health problems in the United Kingdom, oral health problems were not confirmed by a clinician. Any misclassification of oral health may have decreased the power of our study to determine an association between oral disease and glaucoma, and may have biased our results towards the null. Similarly, misclassification of glaucoma due to improper use of diagnostic codes or errors in self-reported disease may have led to over or under-reporting of true glaucoma cases, and biased our results. Furthermore, because pretreatment IOPs were not available for all patients, these were estimated by imputation, as described previously.<sup>19–21</sup> Additionally, secondary subgroup analyses contained smaller sample sizes and may have been underpowered to determine associations. Although the majority of covariables had low missing rates, calorie intake and METS data were available for only a fraction of the participants, and may have affected results. Finally, results from this study are limited by a relatively homogenous population and may not be representative of more diverse populations.

In summary, in this large-scale cross-sectional study, we report no clear association between oral health and glaucoma. Although self-reported denture wear was associated with a small, clinically insignificant change in IOP, this was not protective against a diagnosis of glaucoma or in objective measures—including mRNFL and mGCIPL thickness—related to glaucoma.

## References

1. Ohki T, Itabashi Y, Kohno T, et al. Detection of periodontal bacteria in thrombi of patients with acute myocardial infarction by polymerase chain reaction. *Am Heart J*. 2012;163(2). doi:10.1016/j.ahj.2011.10.012
2. Linden GJ, McClean K, Young I, Evans A, Kee F. Persistently raised C-reactive protein levels are associated with advanced periodontal disease. *J Clin Periodontol*. 2008;35(9). doi:10.1111/j.1600-051X.2008.01288.x
3. Paraskevas S, Huizinga JD, Loos BG. A systematic review and meta-analyses on C-reactive protein in relation to periodontitis. *J Clin Periodontol*. 2008;35(4). doi:10.1111/j.1600-051X.2007.01173.x
4. Wyss-Coray T, Rogers J. Inflammation in Alzheimer disease-A brief review of the basic science and clinical literature. *Cold Spring Harb Perspect Med*. 2012;2(1). doi:10.1101/cshperspect.a006346
5. Pasquale LR, Hyman L, Wiggs JL, et al. Prospective Study of Oral Health and Risk of Primary Open-Angle Glaucoma in Men: Data from the Health Professionals Follow-up Study. In: *Ophthalmology*. Vol 123. ; 2016. doi:10.1016/j.ophtha.2016.07.014
6. Sun KT, Shen TC, Chen SC, et al. Periodontitis and the subsequent risk of glaucoma: results from the real-world practice. *Sci Rep*. 2020;10(1). doi:10.1038/s41598-020-74589-6
7. Byun SH, Yoo DM, Chang M, Choi HG, Hong SJ. Relationship between periodontitis and glaucoma: A cross-sectional study. *J Ophthalmol*. 2020;2020. doi:10.1155/2020/5384602
8. Astafurov K, Elhawy E, Ren L, et al. Oral microbiome link to neurodegeneration in glaucoma. *PLoS One*. 2014;9(9). doi:10.1371/journal.pone.0104416
9. Polla D, Astafurov K, Hawy E, Hyman L, Hou W, Danias J. A Pilot Study to Evaluate the Oral Microbiome and Dental Health in Primary Open-Angle Glaucoma. *J Glaucoma*. 2017;26(4). doi:10.1097/IJG.0000000000000465
10. Wang Y, Hou XW, Liang G, Pan CW. Metabolomics in glaucoma: A systematic review. *Invest Ophthalmol Vis Sci*. 2021;62(6). doi:10.1167/IOVS.62.6.9
11. Baima G, Iaderosa G, Citterio F, et al. Salivary metabolomics for the diagnosis of periodontal diseases: a systematic review with methodological quality assessment. *Metabolomics*. 2021;17(1). doi:10.1007/s11306-020-01754-3

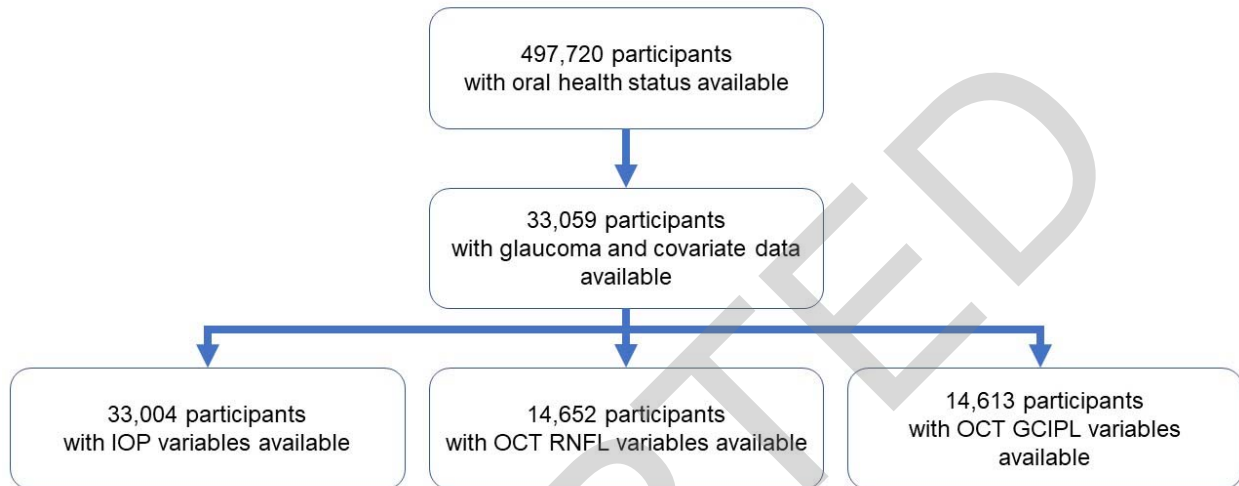
12. Goulão B, MacLennan GS, Ramsay CR. Have you had bleeding from your gums? Self-report to identify giNGival inflammation (The SING diagnostic accuracy and diagnostic model development study). *J Clin Periodontol*. 2021;48(7). doi:10.1111/jcpe.13455
13. Abbood HM, Hinz J, Cherukara G, Macfarlane T V. Validity of Self-Reported Periodontal Disease: A Systematic Review and Meta-Analysis. *J Periodontol*. 2016;87(12). doi:10.1902/jop.2016.160196
14. Pau A, Croucher R, Marcenes W, Leung T. Development and validation of a dental pain-screening questionnaire. *Pain*. 2005;119(1-3). doi:10.1016/j.pain.2005.09.016
15. Pau A, Viswanath KP, Croucher R. Validation of a dental pain screening questionnaire in a semi-urban hospital setting in South India. *Int Dent J*. 2010;60(2). doi:10.1922/IDJ-2332Pau09
16. Palmqvist S, Söderfeldt B, Arnbjerg D. Self-assessment of dental conditions: validity of a questionnaire. *Community Dent Oral Epidemiol*. 1991;19(5). doi:10.1111/j.1600-0528.1991.tb00160.x
17. Gilbert GH, Duncan RP, Kulley AM. Validity of self-reported tooth counts during a telephone screening interview. *J Public Health Dent*. 1997;57(3). doi:10.1111/j.1752-7325.1997.tb02970.x
18. Morris R, Carstairs V. Which deprivation? a comparison of selected deprivation indexes. *J Public Health (Bangkok)*. 1991;13(4). doi:10.1093/oxfordjournals.pubmed.a042650
19. MacGregor S, Ong JS, An J, et al. Genome-wide association study of intraocular pressure uncovers new pathways to glaucoma. *Nat Genet*. 2018;50(8). doi:10.1038/s41588-018-0176-y
20. Khawaja AP, Cooke Bailey JN, Wareham NJ, et al. Genome-wide analyses identify 68 new loci associated with intraocular pressure and improve risk prediction for primary open-angle glaucoma. *Nat Genet*. 2018;50(6). doi:10.1038/s41588-018-0126-8
21. Hysi PG, Cheng CY, Springelkamp H, et al. Genome-wide analysis of multi-ancestry cohorts identifies new loci influencing intraocular pressure and susceptibility to glaucoma. *Nat Genet*. 2014;46(10). doi:10.1038/ng.3087
22. Tewarie P, Balk L, Costello F, et al. The OSCAR-IB consensus criteria for retinal OCT quality assessment. *PLoS One*. 2012;7(4). doi:10.1371/journal.pone.0034823
23. Khawaja AP, Chua S, Hysi P, et al. Comparison of Associations with Different Macular Inner Retinal Thickness Parameters in a Large Cohort: The UK Biobank. *Ophthalmology*. 2020;127(1). doi:10.1016/j.ophtha.2019.08.015

24. Bycroft C, Freeman C, Petkova D, et al. The UK Biobank resource with deep phenotyping and genomic data. *Nature*. 2018;562(7726). doi:10.1038/s41586-018-0579-z
25. Craig JE, Han X, Qassim A, et al. Multitrait analysis of glaucoma identifies new risk loci and enables polygenic prediction of disease susceptibility and progression. *Nat Genet*. 2020;52(2). doi:10.1038/s41588-019-0556-y
26. Chan MPY, Grossi CM, Khawaja AP, et al. Associations with intraocular pressure in a large cohort: Results from the UK Biobank. *Ophthalmology*. 2016;123(4). doi:10.1016/j.ophtha.2015.11.031
27. 20. NHS Digital. Adult Dental Health Survey 2009 - Summary report and thematic series . The Health and Social Care Information Centre.
28. Lehrer S, Rheinstein PH, Schmeidler J. A Component or Multiple Components of Bleeding Gums May Ameliorate Both Glaucoma and Alzheimer's Disease. *Cureus*. Published online 2022. doi:10.7759/cureus.21004
29. National Dental Epidemiology Programme for England. *Oral Health Survey of Adults Attending General Dental Practices 2018.*; 2018.
30. Tomar SL, Asma S. Smoking-attributable periodontitis in the United States: findings from NHANES III. National Health and Nutrition Examination Survey. *J Periodontol*. 2000;71(5).
31. Leite FRM, Nascimento GG, Scheutz F, López R. Effect of Smoking on Periodontitis: A Systematic Review and Meta-regression. *Am J Prev Med*. 2018;54(6). doi:10.1016/j.amepre.2018.02.014
32. Calsina G, Ramón JM, Echeverría JJ. Effects of smoking on periodontal tissues. *J Clin Periodontol*. 2002;29(8). doi:10.1034/j.1600-051X.2002.290815.x
33. Salvi GE, Carollo-Bittel B, Lang NP. Effects of diabetes mellitus on periodontal and peri-implant conditions: Update on associations and risks. In: *Journal of Clinical Periodontology*. Vol 35. ; 2008. doi:10.1111/j.1600-051X.2008.01282.x
34. Chávarry NGM, Vettore MV, Sansone C, Sheiham A. The relationship between diabetes mellitus and destructive periodontal disease: a meta-analysis. *Oral Health Prev Dent*. 2009;7(2). doi:10.3290/j.ohpd.a15518
35. Khader YS, Dauod AS, El-Qaderi SS, Alkafajei A, Batayha WQ. Periodontal status of diabetics compared with nondiabetics: A meta-analysis. *J Diabetes Complications*. 2006;20(1). doi:10.1016/j.jdiacomp.2005.05.006

36. Simpson TC, Weldon JC, Worthington H V., et al. Treatment of periodontal disease for glycaemic control in people with diabetes mellitus. *Cochrane Database of Systematic Reviews*. 2015;2015(11). doi:10.1002/14651858.CD004714.pub3

ACCEPTED

**Figure 1. Flowchart of participant inclusion.** Out of 497,720 participants with oral health status available, a fraction of patients had glaucoma, IOP, RNFL, GCIPL, and covariable data available for analysis.





**Table 1. Demographic variables among subjects with and without oral health conditions in the UK Biobank**

	<b>Without oral health conditions (n=326,905)</b>	<b>With oral health conditions* (n=170,815)</b>	<i>P-value</i>
Age (years)**	55.9 ± 8.1	57.8 ± 8.0	<0.0001
Sex (% male)***	46.1% (150,784)	44.6% (76,126)	<0.0001
Ethnicity			<0.0001
Caucasian	95.3% (310,196)	93.3% (158,816)	
Asian	1.8% (5,720)	2.4% (4,050)	
Black	1.3% (4,293)	2.2% (3,709)	
Other	1.7% (5,412)	2.2% (3,661)	
Townsend deprivation index	-1.5 ± 3.0	-0.8 ± 3.3	<0.0001
Diabetes	4.6% (14,934)	7.4% (12,511)	<0.0001
Alcohol			<0.0001
Never	17.6% (57,448)	23.7% (40,399)	
1-3 drinks x per month	11.0% (36,016)	11.4% (19,408)	
1-2 drinks x per week	26.1% (85,210)	25.3% (43,071)	
3-4 drinks x per week	24.2% (79,090)	20.8% (35,478)	
Daily or almost daily	21.0% (68,744)	18.9% (32,256)	
Smoker			<0.0001
Never	58.6% (191,025)	47.3% (80,498)	
Former	32.3% (105,025)	39.3% (66,794)	
Current	9.2% (29,681)	13.4% (22,741)	
Systolic blood pressure (mmHg)	137.3 ± 18.5	138.8 ± 18.9	<0.0001
Body mass index kilograms/meter <sup>2</sup>	27.1 ± 4.4	27.9 ± 4.7	<0.0001
Energy intake (kJ/day)	8806.5 ± 3191.7	8868.0 ± 3311.1	0.23
Physical activity METS -minutes/week	2638.3 ± 2659.2	2674.2 ± 2817.6	<0.0001
Systemic beta blocker use	5.5% (17,919)	8.0% (13,719)	<0.0001

\*Defined as bleeding gums, painful gums, toothache, loose teeth, and/or denture wear at the time of the survey; Townsend deprivation index is a measure of material deprivation based on residential address with a higher score corresponding to higher levels of poverty (including

measures for employment, lack of car or home ownership, and household crowding); METS represent metabolic equivalents

\*\* All continuous variables presented as means  $\pm$  standard deviations

\*\*\* All categorical variables presented as percentage (n).

ACCEPTED

**Table 2. Adjusted odds ratio of glaucoma prevalence in association with oral health in the UK Biobank (N=33,059)**

	<b>Odds ratio (95% confidence interval)</b>	<b>P-value</b>
Any oral health problem*	1.04 (0.95, 1.14)	0.39
Age (years)	1.10 (1.08, 1.11)	<0.0001
Gender		0.0005
Female	Reference	--
Male	1.33 (1.13, 1.55)	<0.0001
Ethnicity		0.02
Caucasian	Reference	--
Asian	1.80 (1.19, 2.72)	0.005
Black	1.52 (0.97, 2.38)	0.07
Other	1.19 (0.69, 2.06)	0.53
Townsend deprivation index	1.02 (0.99, 1.05)	0.13
Diabetes	1.56 (1.21, 2.02)	0.0007
Alcohol		0.23
Never	Reference	--
1-3 drinks x per month	0.73 (0.53, 1.00)	0.05
1-2 drinks x per week	0.99 (0.78, 1.24)	0.91
3-4 drinks x per week	0.92 (0.73, 1.17)	0.51
Daily or almost daily	1.02 (0.81, 1.29)	0.85
Smoker		0.73
Never	Reference	--
Former	1.05 (0.89, 1.23)	0.57
Current	0.94 (0.68, 1.28)	0.67
Systolic blood pressure (mmHg)	1.00 (0.99, 1.00)	0.84
Body mass index kilogram/meter <sup>2</sup>	1.01, (0.99, 1.03)	0.13
Energy intake (kJ/day)	1.00 (1.00, 1.00)	0.73
Physical activity METS - minutes/week	1.00 (1.00, 1.00)	0.56
Systemic beta blocker use	1.00 (0.75, 1.33)	0.98
Spherical equivalent (diopters)	0.97 (0.95, 0.99)	0.01
MTAG_PRS	2.48 (2.34, 2.63)	<0.0001
MTAG_PRS * Any dental problem	2.48 (2.18, 2.82)	0.84**

\*Defined as history of painful or bleeding gums, toothache, loose teeth, or denture wear at the time of survey; \*\*represents a p-for-interaction.

MTAG\_PRS = multitrait glaucoma polygenic risk score; Townsend deprivation index is a measure of material deprivation based on residential address with a higher score corresponding to higher levels of poverty; METS represent metabolic equivalents

**Table 3. Association between oral health and intraocular pressure (IOP) in the UK Biobank (n=33,004)**

	<b>Difference in IOP, mmHg (95% confidence interval)</b>	<b>P-value</b>
Any oral health problem*	-0.08 (-0.15, -0.009)	0.03
Age (years)	0.05 (0.04, 0.05)	<0.0001
Gender		<0.0001
Female	Reference	--
Male	0.34 (0.27, 0.41)	<0.0001
Ethnicity		0.29
Caucasian	Reference	--
Asian	0.06 (-0.15, 0.2826)	0.567
Black	0.10 (-0.11, 0.32)	0.35
Other	-0.19 (-0.37, 0.07)	0.07
Townsend deprivation index	0.003 (-0.42, 0.04)	0.10
Diabetes	0.18 (0.03, 0.34)	0.02
Alcohol		<0.0001
Never	Reference	--
1-3 x per month	0.04 (-0.04, 0.12)	0.56
1-2 x per week	0.13 (0.06, 0.19)	0.34
3-4 x per week	0.23 (0.16, 0.30)	<0.0001
Daily or almost daily	0.36 (0.29, 0.43)	<0.0001
Smoker		0.006
Never	Reference	--
Former	-0.02 (-0.06, 0.03)	0.51
Current	-0.21 (-0.28, -0.13)	<0.0001
Systolic blood pressure	0.03 (0.03, 0.04)	<0.0001
Body mass index kilogram/meter <sup>2</sup>	-0.005 (-0.01, 0.002)	0.23
Energy intake (kJ/day)	0.000003 (-0.000007, 0.00001)	0.46
Physical activity METS -minutes/week	0.000002 (-0.00001, 0.00001)	0.73
Systemic beta blocker use	-0.53 (-0.62, -0.43)	<0.0001
Spherical equivalent (diopters)	-0.13 (-0.13, -0.12)	<0.0001
IOP_PRS	0.72 (0.70, 0.75)	<0.0001

IOP_PRS * Any dental problem	-0.02 (-0.07, 0.02)	0.37**
------------------------------	---------------------	--------

\*Defined as history of painful or bleeding gums, toothache, loose teeth, or denture wear at the time of survey. IOP\_PRS = intraocular pressure polygenic risk score; \*\* this p-value represents a p-for-interaction. Townsend deprivation index is a measure of material deprivation based on residential address with a higher score corresponding to higher levels of poverty.

ACCEPTED

**Table 4. Association between oral health and mRNFL thickness in the UK Biobank (n=14,652)**

	<b>Difference in thickness, microns (95% confidence interval)</b>	<b>P-value</b>
Any oral health problem*	-0.14 (-0.27, -0.009)	0.04
Age (years)	-0.04 (-0.27, -0.009)	0.04
Gender		<0.0001
Female	Reference	--
Male	-0.62 (-0.75, -0.50)	<0.0001
Ethnicity		<0.0001
Caucasian	Reference	--
Asian	-0.95 (-1.39, -0.52)	<0.0001
Black	-1.30 (-1.72, -0.88)	<0.0001
Other	-0.33 (-0.75, 0.08)	0.12
Townsend deprivation index	0.007 (-0.01, 0.03)	0.53
Diabetes	-0.71 (-1.00, -0.42)	<0.0001
Alcohol		<0.0001
Never	Reference	--
1-3 x per month	-0.02 (-0.25, 0.21)	0.88
1-2 x per week	-0.01 (-0.20, 0.18)	0.89
3-4 x per week	-0.21 (-0.40, -0.01)	0.04
Daily or almost daily	-0.34 (-0.61, -0.21)	<0.0001
Smoker		0.48
Never	Reference	--
Former	0.04 (-0.09, 0.17)	0.58
Current	0.03 (-0.19, 0.25)	0.80
Systolic blood pressure	-0.004 (-0.008, -0.0006)	0.02
Body mass index kilogram/meter <sup>2</sup>	-0.04 (-0.05, -0.02)	<0.0001
Energy intake (kJ/day)	0.00002 (-0.000002, 0.00004)	0.07
Physical activity METS -minutes/week	0.000005 (-0.00001, 0.00003)	0.66
Systemic beta blocker use	-0.21 (-0.49, 0.06)	0.13
Spherical equivalent (diopters)	-0.34 (-0.37, -0.31)	<0.0001
MTAG_PRS	-0.02 (-0.05=9, 0.05)	0.51
MTAG_PRS * Any dental problem	0.09 (-0.04, 0.21)	0.17**

Downloaded from <http://journals.lww.com/glaucorajournal> by BhdMfsePfkav1zeoum1tQIN4a+kjLHEZ9bsIho4XM on 04/14/2024

\*Defined as history of painful or bleeding gums, toothache, loose teeth, or denture wear at the time of survey; mRNFL=macula retinal nerve fiber layer; MTAG\_PRS = multitrait glaucoma polygenic risk score. \*\*This p-value represents a p-for-interaction. Townsend deprivation index is a measure of material deprivation based on residential address with a higher score corresponding to higher levels of poverty.

ACCEPTED

**Table 5. Association between oral health and mGCIPL thickness in the UK Biobank (n=14,613)**

	<b>Estimate, microns (95% confidence interval)</b>	<b>P-value</b>
Any oral health problem*	0.004 (-0.17, 0.17)	0.96
Age (years)	-0.15 (-0.16, -0.14)	<0.0001
Gender		0.55
Female	Reference	--
Male	-0.05 (-0.22, 0.12)	0.55
Ethnicity		<0.0001
Caucasian	Reference	--
Asian	-1.42 (-2.00, -0.85)	<0.0001
Black	-0.22 (-0.78, 0.33)	0.44
Other	0.53 (-0.02, 1.08)	0.06
Townsend deprivation index	-0.03 (-0.06, -0.001)	0.04
Diabetes	-0.73 (-1.11, -0.34)	0.0002
Alcohol		<0.0001
Never	Reference	--
1-3 x per month	0.09 (-0.22, 0.39)	0.57
1-2 x per week	0.11 (-0.15, 0.36)	0.40
3-4 x per week	-0.05 (-0.31, 0.20)	0.68
Daily or almost daily	-0.51 (-0.78, -0.25)	0.0001
Smoker		0.004
Never	Reference	--
Former	0.14 (-0.03, 0.31)	0.12
Current	0.17 (-0.12, 0.47)	0.25
Systolic blood pressure	-0.005 (-0.009, 0.0002)	0.06
Body mass index kilogram/meter <sup>2</sup>	-0.03 (-0.05, -0.02)	0.0004
Energy intake (kJ/day)	0.0002 (-0.00001, 0.00004)	0.27
Physical activity METS -minutes/week	0.00002 (-0.00001, 0.00005)	0.22
Systemic beta blocker use	-0.40 (-0.77, -0.03)	0.03
MTAG_PRS	-0.16 (-0.25, -0.07)	0.0006
MTAG_PRS * Any dental problem	0.11 (-0.05, 0.27)	0.19**



\*Defined as history of painful or bleeding gums, toothache, loose teeth, or denture wear at the time of survey; mGCIPL=macula ganglion cell inner plexiform layer; \*\* This p value represents a p-for-interaction.

ACCEPTED