The closed eye harbours a unique microbiome in dry eye disease

Kent A. Willis1, Cameron K. Postnikoff2,3, Amelia Freeman4, Gabriel Rezonzew4, Kelly Nichols2, Amit Gaggar5 & Charitharth V. Lal4,5*

Dry eye affects millions of individuals. In experimental models, dry eye disease is associated with Th17-mediated inflammation of the ocular surface that may cause persistent damage to the corneal epithelium. However, the initiating and perpetuating factors associated with chronic inflammation of the ocular surface remain unclear. The ocular microbiota alters ocular surface inflammation and may influence dry eye disease development and progression. Here, we collected serial samples of tears on awakening from sleep, closed eye tears, during a randomized clinical trial of a non-pharmaceutical dry eye therapy and used 16S rRNA metabarcoding to characterize the microbiome. We show the closed dry eye microbiome is distinct from the healthy closed eye microbiome, and that the microbiome remains distinct despite daily saline eye wash upon awakening. The ocular microbiome was described only recently, and this report implicates a distinct microbiome in ocular disease development. Our findings suggest an interplay between microbial commensals and inflammation on the ocular surface. This information may inform future studies of the pathophysiological mechanisms of dry eye disease.

Results

Cohort. Based on the power and pairwise sample-size estimator for permutational multivariate analysis of variance (PERMANOVA) application Micropower14, a low abundance 16S rRNA dataset similar to previous ocular microbiome studies15 would require a minimal sample size of 30 to generate a discriminatory power of 0.8 with a significance level of 0.05. Therefore, we aimed for at least 35 subjects per study arm. Table 1 shows the demographics and clinical characteristics of the enrolled participants. Based on a post-enrolment exam and two clinical surveys, after informed consent, participants were classified as dry eye or controls (Supplemental S1). The resulting 36 control and 36 dry eye subjects were then stratified by clinical severity of eye disease.

1Division of Neonatology, Department of Pediatrics, College of Medicine, The University of Tennessee Health Science Center, Memphis, TN, USA. 2School of Optometry, University of Alabama At Birmingham, Birmingham, AL, USA. 3CooperVision, Inc, Pleasanton, CA, USA. 4Division of Neonatology, Department of Pediatrics, College of Medicine, University of Alabama At Birmingham, Birmingham, AL, USA. 5Program in Protease and Matrix Biology, Department of Pediatrics, College of Medicine, Women and Infants Center, University of Alabama At Birmingham, 176F Suite 9380, 619 South 19th Street, Birmingham, AL 35249-7335, USA. *email: clal@peds.uab.edu
The study arms were randomized to daily saline eye wash upon awakening or non-intervention. A baseline closed eye tear sample was collected at randomization and the final sample after one month, yielding 144 samples. We did not observe significant differences between the stratifications normal versus mild and moderate versus severe with respect to microbial ecology (Fig. 2), however clear separation was present between the mild and moderate samples, indicating this more granular stratification introduced a false dichotomy in the data. Therefore, we combined subjects stratified as none or mild into one cohort, now termed normal, and we combined subjects stratified as moderate or severe into another cohort, now termed dry eye (Fig. 3). We used this simplified stratification system for all subsequent analyses (Methods).

Sequencing data. To investigate the bacterial microbiome of the closed dry eye, we used metabarcoding of the 16S rRNA gene. Sequence reads were generated using the Illumina MiSeq system (57,022 ± 47,011 counts/sample, Methods). We assigned taxonomy using the Greengenes database15, producing 1,593 data points/sample. After excluding any reads aligned to chloroplasts or cyanobacteria, we examined the bacterial community composition using the remaining 1,195 data points/sample that were collectively assigned to 185 genera16. All tear samples had the standard > 1,000 aligned reads/sample, so we retained all samples for further analysis. We used a log2-transformation of cumulative sum scaling17 (log2 CSS) to normalize our dataset, consistently producing 8,700–10,000 reads per sample (Supplemental Fig. S2).

### Table 1. Participant demographics and clinical characteristics. *SD = standard deviation.

|                      | Normal eye | Dry eye | P value |
|----------------------|------------|---------|---------|
| Subjects, n          | 36         | 36      |         |
| Sex                  |            |         | 1.00    |
| Female, n (%)        | 23 (64)    | 23 (64) |         |
| Male, n (%)          | 13 (36)    | 13 (36) |         |
| Age, y ± SD          | 33 ± 11    | 52 ± 19 | < 0.0001|
| DEQ Score            | 1.2 ± 1.3  | 12 ± 3.3| < 0.0001|
| Phenol Red Thread Wetting length, mm ± SD | 22 ± 5.3 | 19 ± 8.6 | 0.08 |
| NIKBUT, s ± SD       | 15 ± 5.2   | 10.7 ± 5.2| 0.0007 |
| Corneal Staining     | 0.47 ± 0.8 | 3.6 ± 3 | < 0.0001|
| InflammaDry Positive, n (%) | 13 (36) | 20 (55) | 0.10 |
| Randomization        |            |         |         |
| Control, n (%)       | 16 (44)    | 20 (56) |         |
| Treatment, n (%)     | 20 (56)    | 16 (44) |         |

(Fig. 1 and Supplemental Table 1). The study arms were randomized to daily saline eye wash upon awakening or non-intervention. A baseline closed eye tear sample was collected at randomization and the final sample after one month, yielding 144 samples. We did not observe significant differences between the stratifications normal versus mild and moderate versus severe with respect to microbial ecology (Fig. 2), however clear separation was present between the mild and moderate samples, indicating this more granular stratification introduced a false dichotomy in the data. Therefore, we combined subjects stratified as none or mild into one cohort, now termed normal, and we combined subjects stratified as moderate or severe into another cohort, now termed dry eye (Fig. 3). We used this simplified stratification system for all subsequent analyses (Methods).
(OTUs), we used Spearman network analysis (Fig. 2d).

at the order level. To further investigate the importance of particular bacterial operational taxonomic units Table 3) among 49 genera with significant differences in relative abundance. Figure 2c shows relative abundance

P
operational taxonomic unit level. Positive correlations with a

sum scaling. Two-way ANOVA, *P = 7.5 × 10−5, evenness, P = 0.001. (c) Relative abundance of bacterial orders. Log2(CSS), log, transformation of cumulative-

Figure 2.
f
disease have different alpha diversity of the closed eye microbiome as quantified by the richness (ANOV A, 
P = 4.8, P = 7.5 × 10−5), evenness (ANOVA, f = 13, P = 2.1 × 10−12) and Shannon diversity (ANOVA, f = 12, 
P = 5.9 × 10−12) indices. (b) The beta diversity of the closed eye microbiome in individuals with dry eye disease is distinct by principal coordinate analysis (PCoA) of Bray–Curtis dissimilarity (R² = 0.21 P = 0.00033), redundancy analysis (RDA, variance = 95.25, f = 3.71, P = 0.001) and canonical correspondence analysis (CCA, chi² = 0.16 f = 3.64 P = 0.001). (c) Relative abundance of bacterial orders. Log2(CSS), log, transformation of cumulative-

The closed eye microbiome is distinct in dry eye disease. In the taxonomic analysis of the closed eye microbiome, the microbiomes of individuals with chronic dry eye disease and those without form distinct communities. Dry eye microbial communities are more diverse as quantified by richness (ANOVA, f = 4.8, P = 7.5 × 10−5), evenness (ANOVA, f = 13, P = 2.1 × 10−12) and Shannon diversity (ANOVA, f = 12, P = 5.9 × 10−12), Fig. 2a). As demonstrated by principal coordinate analysis (PCoA) of Bray–Curtis dissimilarity, these communities form unique clusters as quantified by PERMANOVA (R² = 0.21, P = 0.00033, Fig. 2b). Similarly, multivariate redundancy analysis (RDA, variance = 95.25, f = 3.71, P = 0.001) and canonical correspondence analysis (CCA, chi² = 0.16 f = 3.64, P = 0.001) showed these communities are distinct, and this factor accounts for the majority of variance in the data. As quantified by two-way ANOVA adjusted for false discovery rate (FDR), univariate analysis of the relative abundance of bacterial genera showed individuals with dry eye disease have differences in the relative abundance of 113 genera, with the most significant differences in the abundance of OPB56, Methylobacteriaceae, Bacteroidetes, Pseudomonas, and Meiothermus (all with FDR-adjusted P < 2 × 10−22, Supplemental Table 2). Similarly, mixed effects regression detected the most significant differences in the abundance of OPB56, Bacteroidetes, Pseudomonas, Meiothermus, and Methylobacteriaceae (all with FDR-adjusted P < 3.1 × 10−20, Supplemental Table 3) among 49 genera with significant differences in relative abundance. Figure 2c shows relative abundance at the order level. To further investigate the importance of particular bacterial operational taxonomic units (OTUs), we used Spearman network analysis (Fig. 2d).

The closed eye microbiome remains distinct despite daily eye wash. Daily eye wash on awakening with sterile saline is a proposed therapy for dry eye disease18. We tested if the microbial community composition could be normalized by the prescription of daily saline eye wash. The diversity of subjects with dry eye remained higher than that of normal individuals (richness, t-test, P = 7.5 × 10−5, evenness, t-test, P = 2.8 × 10−13 and Shannon diversity, t-test, P = 6.1 × 10−14, Fig. 3a). Multivariate analysis of these communities also remained distinct (PCoA, PERMANOVA, R² = 0.15, P = 0.00033, RDA, variance = 64.75, f = 5.73, P = 0.001 and CCA, chi² = 0.11 f = 5.62, P = 0.001, Fig. 3b). Univariate analysis detected the most significant differences in the abundance of MLE112, Lactobacillaceae, Streptococcus, Sphingobium, Caldicobacteriaceae and Anaerococcus (ANOVA, P < 0.01, Fig. 3c). Core microbiome analysis highlighted key differences in the distribution of unique genera, and network analysis demonstrated the importance of key taxa (Fig. 3d,e).

The closed eye microbiome in dry eye disease is distinct at baseline. To gain further insight into the microbial community composition of the dry eye, we narrowed our focus to examine the closed eye microbial composition at the time of randomization. As indicated by our more general analysis, both diversity (richness,
The distinct clustering of these communities on multivariate analysis also persisted (PCoA of Bray–Curtis dissimilarity (PERMANOVA, $R^2 = 0.15, P = 0.00033$), redundancy analysis (RDA, variance = 64.75, $f = 5.73, P = 0.001$) and canonical correspondence analysis (CCA, $\chi^2 = 0.11 f = 5.62 P = 0.001$). (c) Relative abundance of bacterial genera. Log2(CSS), log transformation of cumulative-sum scaling. Two-way ANOVA, $* P = 0.01$, $** P = 0.001$, $*** P = 0.0001$. (d) Core microbiome analysis showing differences in microbial colonization at the genus level. (e) Spearman network analysis at the operational taxonomic unit level. Positive correlations with a $P$ value $< 0.05$ are shown as an edge with the relative size determined by the importance of the taxa to the network.

Figure 3. Daily eye rinse does not alter the closed eye microbiome. (a) Despite daily eye washes, the alpha diversity of the closed eye microbiome remains similar to baseline in individuals with and without dry eye disease as quantified by the richness (ANOVA, $f = 9.2, P = 7.5 \times 10^{-5}$), evenness (ANOVA, $f = 28, P = 2.8 \times 10^{-14}$) and Shannon diversity indices (ANOVA, $f = 28, P = 6.1 \times 10^{-14}$). (b) The beta diversity of the closed eye microbiome remains distinct in individuals with and without dry eye disease by principal coordinate analysis (PCoA) of Bray–Curtis dissimilarity (PERMANOVA, $R^2 = 0.15 P = 0.00033$), redundancy analysis (RDA, variance = 64.75, $f = 5.73, P = 0.001$) and canonical correspondence analysis (CCA, $\chi^2 = 0.11 f = 5.62 P = 0.001$). (c) Relative abundance of bacterial genera. Log2(CSS), log transformation of cumulative-sum scaling. Two-way ANOVA, $* P = 0.01$, $** P = 0.001$, $*** P = 0.0001$. (d) Core microbiome analysis showing differences in microbial colonization at the genus level. (e) Spearman network analysis at the operational taxonomic unit level. Positive correlations with a $P$ value $< 0.05$ are shown as an edge with the relative size determined by the importance of the taxa to the network.

Machine learning accurately classifies dry eye samples at baseline. To more precisely identify unique features with the potential to function as biomarkers for patients with dry eye, we used linear discriminant analysis of effect size. We identified 5 genera that reliably identified individuals with dry eye (Pseudomonas, Methylobacteriaceae, Pseudomonas, Bradyrhizobium and Allobaculum) as the five most differently abundant genera (all with FDR-adjusted $P < 1.1 \times 10^{-11}$, Supplemental Fig. S5a and Supplemental Table 4). At the phylum level, Firmicutes and Bacteroidetes remained unchanged, while Verrucomicrobia (FDR $P = 4.6 \times 10^{-11}$) and Proteobacteria (FDR $P = 6.0 \times 10^{-11}$) showed the most significant differences. Core microbiome analysis identified 45 genera unique to dry eye with only 14 genera unique to the normal eye (Fig. 4d). Discriminant analysis of principal components at the order level revealed the abundances of orders OPB56 and Rhizobiales are important to discriminate the normal eye while the orders Halanaerobiales, Erysipelotrichales and Anaeroplasmatales are important to discriminate dry eye (Fig. 4e). Network analysis provided further evidence of these distinct communities (Fig. 4f).

The closed eye microbiome in dry eye disease diverges after one month. We examined the stability of the dry eye microbiome by focusing on samples collected one month later in the same individuals. The dry eye microbiome remained distinct from the normal microbiome and remained consistent with the community composition noted at baseline (Supplemental Figs. S2, S3). Only slight divergence was noted over the course of a month (Fig. 5). The diversity of dry eye remained higher than that of the normal eye (richness, $t$-test, $P = 5.2 \times 10^{-5}$, evenness, $t$-test, $P = 1.9 \times 10^{-11}$ and Shannon diversity, $t$-test, $P = 8.6 \times 10^{-11}$, Fig. 5a). The distinct clustering of these communities on multivariate analysis also persisted (PCoA of Bray–Curtis dis-
similarity, PERMANOVA, $R^2 = 1.52, P = 0.00033$, RDA, variance = 66.53, $f = 8.52, P = 0.001$ and canonical correspondence analysis (CCA, $\chi^2 = 0.10, f = 8.14, P = 0.001$). (c) Relative abundance of bacterial orders. Log$_2$(CSS), log, transformation of cumulative-sum scaling. Two-way ANOVA (**displaying only, $P < 0.001$). (d) Core microbiome analysis showing differences in microbial colonization at the genus level. (e) Discriminant analysis of principal components at the order level. (f) Spearman network analysis at the operational taxonomic unit level. Positive correlations with a $P$ value $< 0.05$ are shown as an edge with the relative size determined by the importance of the taxa to the network.

**Figure 4.** The closed eye microbiome in dry eye disease is distinct at baseline. (a) The alpha diversity of the tear microbiome remains distinct as quantified by the richness ($t$-test, $f = 6.1, P = 0.016$), evenness ($t$-test, $f = 24$, $P = 5.8 \times 10^{-5}$) and Shannon diversity ($t$-test, $f = 23, P = 8.1 \times 10^{-5}$) indices. (b) The beta diversity of the tear microbiome remains distinct as quantified by principal coordinate analysis (PCoA) of Bray–Curtis dissimilarity (PERMANOVA, $R^2 = 1.34, P = 0.00033$), redundancy analysis (RDA, variance = 53.95, $f = 7.12, P = 0.001$) and canonical correspondence analysis (CCA, $\chi^2 = 0.09, f = 7.12, P = 0.001$). (c) Relative abundance of bacterial orders. Log$_2$(CSS), log, transformation of cumulative-sum scaling. Two-way ANOVA (**displaying only, $P < 0.001$). (d) Core microbiome analysis showing differences in microbial colonization at the genus level. (e) Discriminant analysis of principal components at the order level. (f) Spearman network analysis at the operational taxonomic unit level. Positive correlations with a $P$ value $< 0.05$ are shown as an edge with the relative size determined by the importance of the taxa to the network.

similarly, PERMANOVA, $R^2 = 1.52, P = 0.00033$, RDA, variance = 66.53, $f = 8.52, P = 0.001$ and CCA, $\chi^2 = 0.10, f = 8.14, P = 0.001$ (Fig. 5b). ANOVA identified differentially abundant genera (Fig. 5c). Similarly, negative binomial regression noted the greatest differences in the abundance of the genera *Methylobacteriaceae*, *Pseudomonas*, *Azospirillum*, *Bradyrhizobium* and *Coriobacteriaceae* (all with FDR-adjusted $P < 7.0 \times 10^{-18}$, Supplemental Fig. S6a, Supplemental Table 4). However, core microbiome analysis detected 76 unique genera in dry eye, 69 shared genera and 24 genera unique to the normal eye (Fig. 5d). Similar to baseline, discriminant analysis of principal components at the order level identified OPB58 and Bacteroidetes as important discriminators of the normal eye. In individuals with dry eye, additional orders enabled further discrimination, with Flavobacterales, Alteromonadales and Actinomycetes, Anaeroplasmatales and Desulfuromonadales functioning as the primary discriminators instead of Halanaeobiales and Erysipelotrichales (Fig. 5d).

**Machine learning more accurately identifies dry eye after a month.** We used linear discriminant analysis of effect size to identify bacterial genera with the potential function as biomarkers and noted 10 genera unique to individuals with dry eye and 9 unique to individuals with a normal eye, supporting the limited divergence of these communities (Supplemental Fig. S6b). We also developed a support vector machine with leave-one-out cross-validation that identified dry eye samples with 97% accuracy. Variable importance analysis of a random forest classifier noted the prevalence of *Methylobacterium*, *Megasphaera*, *Parabacteroides*, S247, *Bifidobacterium*, *Streptococcus*, *Desulfovibrio* and *Acetobacter* as important identifiers (Importance $> 20$, Supplemental Fig. S6c).

**The closed eye microbiome at baseline is unaltered by the cellular concentration of the tear fraction.** To ensure our microbial ecological analysis was unaltered by the cellular content of the tear fraction, we compared the microbial composition of each cellular fraction collected at baseline. Three samples were excluded from this analysis due to incomplete data (n = 141). Community composition was similar between both high and low cellular fractions (Fig. 6).

**Discussion**
Direct crosstalk between resident microbes and host immune cells at the mucosal surface is a critical determinant of inflammatory diseases. Furthermore, Th17 and T regulatory (Treg) cells are implicated in the development of chronic dry eye disease and particularly attuned to the ecology of the resident microbiota. Therefore, the ocular surface microbiome may contribute to the pathogenesis of dry eye disease. Understanding how the ocular
Figure 5. The final closed eye microbiome in dry eye disease remains distinct. (a) The alpha diversity of the closed eye microbiome remains distinct after four weeks as quantified by the richness (t-test, f = 19, P = 5.2 × 10^{-5}), evenness (t-test, f = 65, P = 1.9 × 10^{-11}) and Shannon diversity (t-test, f = 60, P = 8.6 × 10^{-11}) indices. (b) The beta diversity of the closed eye microbiome remains distinct after four weeks as quantified by principal coordinate analysis (PCoA) of Bray–Curtis dissimilarity (PERMANOVA, R^2 = 1.52 P = 0.00033), redundancy analysis (RDA, variance = 66.53, f = 8.52, P = 0.001) and canonical correspondence analysis (CCA, chi^2 = 0.10, f = 8.14, P = 0.001). (c) Relative abundance of bacterial orders. Log(CSS), log transformation of cumulative-sum scaling. Two-way ANOVA, * P = 0.05, ** P = 0.01, *** P = 0.001. (d) Core microbiome analysis showing differences in microbial colonization at the order level. (e) Discriminant analysis of principal components at the order level. (f) Spearman network analysis at the operational taxonomic unit level. Positive correlations with a P value < 0.05 are shown as an edge with the relative size determined by the importance of the taxa to the network.

Figure 6. The closed eye microbiome is unaltered by the cellular fraction at baseline. (a) The alpha diversity is similar as quantified by the richness (t-test, f = 1.3, P = 0.26), evenness (t-test, f = 1.8, P = 0.19) or Shannon diversity (t-test, f = 1.9, P = 0.17) indices. (b) The beta diversity of the closed eye microbiome is unaltered by the cellular fraction as quantified by principal coordinate analysis (PCoA) of Bray–Curtis dissimilarity (PERMANOVA, R^2 = 0.00894 P = 0.124), redundancy analysis (RDA, variance = 4.82, f = 1.29, P = 0.152) and non-metric multidimensional scaling (NMDS, stress = 0.228). (c) Relative abundance of bacterial genera. Log(CSS), log transformation of cumulative-sum scaling. Two-way ANOVA, * P = 0.05, ** P = 0.01, *** P = 0.001. (d) Discriminant analysis of principal components. (e) Spearman network analysis at the operational taxonomic unit level. Positive correlations with a P value < 0.05 are shown as an edge with the relative size determined by the importance of the taxa to the network.
microbiome influences ocular surface disease will be important for therapeutics development. In this report, we present initial evidence of a distinct closed eye microbiome in dry eye.

In this study, human subjects collected samples of sterile saline rinse applied to the ocular surface upon awakening in the morning. Using saline eye wash to collect microbiome samples is novel and less invasive than swabs of the ocular surface; that has previously been used for the collection of tear leukocytes. In a series of pooled samples from both eyes, the closed eye tear microbiome in individuals with dry eye disease was distinct from that in healthy controls. Compared to swabs, where the fibre content and limited sample area artificially reduce the number of detectable microbes, saline wash may allow for a less biased sampling of the entire ocular surface, especially when pooled from both eyes. Furthermore, a general weakness of previous ocular microbiome studies is focusing on shifts in percent relative abundance to identify differences in bacterial taxa. A strength of our analysis is the rigorous log2 normalization of the cumulative sum scaling transformation performed prior to analysis. With this normalization, our data more accurately reflect true shifts in relative abundance. Using untransformed data, the most significant difference in our dataset was a nearly complete substitution of unnormalized relative abundance of Pseudomonas spp. and Staphylococcus spp. and the untransformed data, the most significant difference in our dataset was a nearly complete substitution of unnormalized relative abundance of Staphylococcus spp. and Pseudomonas. A more rigorous normalization performed on our dataset confirmed the reduction in Pseudomonas spp. but not in Staphylococcus spp. In addition, 48 other genera may warrant further investigation.

Previous research has focused on the open eye microbiome, while we focused on the closed eye microbiome. Every night during sleep, inflammatory species move into the closed eye tear film, perhaps in response to the immune system of the closed eye cannot regulate the ocular surface microbiome during sleep. In the closed eye, preliminary evidence suggests higher numbers of neutrophils with an enhanced degranulation response to lysozyme and antimicrobial compounds in tears. In dry eye disease, our results suggest increased microbial diversity is a hallmark of disease. Consistent with the behaviour of the ocular microbiome in other pathological conditions, this finding may represent breakdown of the microbe-oriented homeostatic mechanisms of the host. This finding may also suggest increased diversity results from the incorporation of pathobionts into this community. Alternatively, the altered ecology may result from changes within the ocular surface of the dry eye that support establishment of a wider diversity of microbes in this environment. These additional species may not be harmful but instead highlight the dynamic biome within the human eye. The evidence may also suggest the immune system of the closed eye cannot regulate the ocular surface microbiome during sleep. In the closed eye, preliminary evidence suggests higher numbers of neutrophils with an enhanced degranulation response in dry eye disease, indicating a dysregulated immune response. This dysregulation may permit increased microbial diversity. The reverse could also be true: enhanced microbial diversity may induce a greater inflammatory response.

This dataset consistently demonstrates the microbial communities in dry eye disease are distinct and remain distinct even with daily saline eye wash. We asked subjects in the treatment arm to self-administer saline eye wash immediately upon awakening. We hypothesized this time may represent the point of maximum difference since microbes likely accumulate on the ocular surface overnight. However, we did not observe reduced bacterial diversity with saline eyewash, with daily saline eye wash becoming more similar to control eye communities with treatment. Instead, the communities slightly diverged: the microbial communities in dry eye remained different from those in healthy eyes. Thus, the mechanisms of microbial accumulation on the ocular surface may be more complex than moisture content or tear clearance. Intrinsic immune or mechanical alterations in dry eye disease may foster differences in commensal abundance, potentially explaining how altered bacterial communities on the ocular surface contribute to dry eye disease.

Our study has a few potential limitations. Our assessment of the microbiome of the ocular surface is based upon 97% homologous OTU clustering using Illumina MiSeq for the bacterial 16S rRNA gene and as such as some intrinsic limitations. The first, that based on this technology, only relative abundance of the specific OTUs can be determined. In future studies it may be informative to utilize shotgun metagenomic sequencing or spike-in methods with standardized microbial communities to have a better understanding of absolute microbial abundance. In addition, patients with dry eye were older than their disease-free counter parts. This may warrant further exploration in future studies.
More research is needed to reveal whether an altered microbiome drives the development of dry eye disease. The ocular microbiome potentially hosts a distinct core microbiome that may be perturbed in eye disease. Our study provides novel insight into the closed eye microbiome in dry eye disease. Th17 cells are sensitive to changes in the microbiome in experimental dry eye disease. Therefore, increased microbial abundance in dry eye may promote the development of dry eye disease by altering T cell subpopulations on the ocular surface. More research is needed to reveal whether an altered microbiome drives the development of dry eye disease.

Methods

Participants. This study was performed in accordance with the Declaration of Helsinki under the supervision of the Institutional Review Board of the University of Alabama at Birmingham (UAB), Birmingham, AL, USA. We performed a secondary analysis of the closed eye microbiome from a prospective, randomized controlled trial of the efficacy of daily eye wash to ameliorate inflammation in individuals with dry eye disease (ClinicalTrials.gov NCT03332342). Informed consent was obtained from all participants prior to enrolment. Participants were selected based on the results of an optometric exam and two clinical surveys, the Dry Eye Questionnaire 5 (DEQ5) and the Ocular Surface Disease Index (OSDI). Exclusion criteria included pregnancy, contact lens use within the past three months, glaucoma treatment, current tobacco use, systemic dry-eye-associated inflammatory disease, the use of any form of anti-inflammatory therapy for dry eye treatment in the last three months, other antimicrobial or anti-inflammatory medications in the last month (Supplemental Fig. S1). Subjects were asked to refrain from any artificial tear use for the duration of the study. Stratification into normal and dry eye was driven primarily by symptoms, as measured by the DEQ5 score, with further stratification among both groups determined by phenol red thread wetting length, non-invasive keratograph tear break-up time (NIK-BUT) and the InflammaDry test (Quidel; San Diego, CA, USA. Supplemental Table 1). Dry eye subjects needed to have a DEQ5 score of greater than or equal to 6, and present with a positive dry eye result with the phenol red thread (< 10 mm), NIKBUT (< 10 s), or InflammaDry (positive). Dry eye subjects also had to endorse the sensation of dry eye or have a previous diagnosis of dry eye disease. Equal-weight randomization to either daily preservative-free sterile saline eye wash or control was performed using REDCap (Research Electronic Data Capture; Nashville, TN, USA). Subjects were trained on the first visit to perform a daily eye wash. Briefly, as described previously, subjects were provided one 50 mL sterile centrifuge tube, and two 10 mL sterile polypropylene syringes (BD Biosciences; San Jose, CA, USA) with 5 mL of sterile saline per syringe. Immediately after awakening, subjects were instructed to wash their eyes by dispensing the saline gently across the ocular surface, with the runoff collected in a sterile Eppendorf tube. Runoff from both eyes was collected to produce a pooled sample of both eyes collected into the same tube. Subjects in the treatment group completed daily therapy for approximately 28 days.

Sample collection. We utilized the power and pairwise sample-size estimator for PERMANOVA application Micropower, as implemented at https://fedematt.shinyapps.io/shinyMB/, to estimate the minimum number of samples to produce a power of 0.8 to detect a difference with a significance level of 0.05 via PERMANOVA. We processed the collected eyewash samples immediately after they were dropped off by study subjects. Pooled tear collections were centrifuged at 270× g, and the supernatant was aliquoted into 1.5 mL microcentrifuge tubes and stored at −80 °C. If more than one million cells were recovered in the pooled tear collection, supernatant aliquots were spiked with cells from the precipitate to increase the potential yield of microbiome analysis by normalizing the cellular fraction.

Illumina MiSeq metabarcoding. In the Microbiome Core at UAB, microbial genomic DNA was isolated and PCR was used with unique bar-coded primers to amplify the V4 region of the 16S rRNA gene to create a 16S amplicon library of the samples. The PCR products were sequenced using the NextGen Illumina MiSeq System (Illumina Inc.; San Diego, CA, USA). The sequence data covered the 16S rRNA V4 region with a PCR product length of ~255 bases and 250 base paired-end reads. The raw dataset is available at the NCBI Sequence Read Archive: https://www.ncbi.nlm.nih.gov/bioproject/597168.

Bioinformatics. Sequences were grouped into OTUs using QIIME v1.9.41. We assigned taxonomy using the Ribosomal Database Project classifier (threshold 0.8) against the Greengenes database. We then imported these processed sequence reads into Calypso v8.84 for further processing and data analysis. After excluding any OTUs assigned to chloroplasts or cyanobacteria, we a priori determined to exclude any samples with < 1,000 reads/sample from downstream analysis. We then utilized cumulative sum scaling to normalize the relative
abundance and performed a log$_2$-transformation of the resulting data, centring the taxonomic counts to 0 and scaling to a range of –2 to 2 with a variance of 1. For alpha diversity analysis, samples were rarefied to a depth of 7,804 reads/sample. Alpha diversity was then quantified using the richness, evenness and Shannon diversity indices, testing for significant differences with ANOVA adjusted for FDR. For beta diversity, we visualized the data using PCoA, RDA, CCA or NMDS. We used PERMANOVA to determine significant differences in beta diversity. Relative abundance was quantified using ANOVA adjusted by FDR for multiple groups or negative binomial regression (DESeq2 function) for binary comparisons. For Spearman network analysis, we displayed positive correlations with an FDR-adjusted $P<0.05$ as an edge. Relative size of the point represents the importance of the taxa to the resulting network. In addition, we used core microbiome analysis (of the 300 most abundant taxa), linear discriminant analysis of effect size, and discriminant analysis of principal components to select significant features. For analyses with two groups, we developed machine learning classifiers using support vector machine and random forest algorithms.

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**Author contributions**

CKP, AG and CVL conceptualized the study. CKP, KN facilitated the performance of the randomized clinical trial and oversaw the sample collection. GR performed the DNA extraction. KAW designed and performed the biostatistical analyses, drafted the manuscript and prepared the figures. CVL directed the performance of the study and provided funding. KAW, CKP, AF, GR, KN, AG and CVL edited the manuscript and approved the final version.

**Competing financial interests**

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**Additional information**

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**Correspondence** and requests for materials should be addressed to C.V.L.

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