ORIGINAL RESEARCH Characteristics of Oral-Gut Microbiota in Model Rats with CUMS-Induced Depression

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Purpose: The diversity and composition of the oral and gut microbiota of depressed rats were analyzed to explore the microbiological etiology of major depressive disorder (MDD).

Methods: The depressed rat model was established by inducing chronic unpredictable mild stress (CUMS). After the establishment of the model, body weight measurements and behavioral tests were conducted. The diversity and composition of oral and gut microbiota were analyzed using 16SrRNA sequencing.

Results: There were significant differences in the alpha and beta diversity of the oral microbiota of rats in the CUMS and control groups. The top three most abundant genera in the oral microbiota were Rothia, Psychrobacter, and Streptococcus. Linear discriminant analysis effect size (LEfSe) analysis showed that the abundance of Rothia decreased and that of Psychrotrophs increased in the CUMS group, and the differences were statistically significant. The top three most abundant genera in the gut microbiota were Lactobacillus, Ruminococcus and Oscillospira. LEfSe analysis showed that the abundance of Ruminococcus decreased in the CUMS group, and the difference was statistically significant. Spearman correlation analysis was performed to analyze the differential microbiota and depression-like behavior, which showed that differential microbiota significantly correlated with body weight, total distance traveled, average speed, and number of rearing. Spearman correlation analysis of oral and gut differential microbiota demonstrated a strong positive correlation between Facklamia in the oral cavity and Enterococcus, Streptococcus in the intestine (r=0.64-0.73, P<0.01); along with a strong negative correlation between Desulfovibrio in the oral cavity and Enterococcus, Turicibacter in the intestine (r=-0.51--0.72, P<0.05).

Conclusion: Significant differences were observed in the diversity and composition of oral and gut microbiota between the CUMS depression model and control groups. Modulating the oral and gut microbiota may have positive effects on MDD. Keywords: major depressive disorder, oral microbiota, gut microbiota, 16SrRNA sequencing

Introduction

The human microbiota is a diverse, complex, and symbiotic aggregate of microorganisms residing at various body sites like skin, oral cavity, gastrointestinal tract, respiratory tract, and urogenital tract.¹ The oral cavity and gut are the two largest microbial habitats in the human body. An increasing number of studies have shown that the microbiota is involved in various physiological processes such as metabolism, neurotransmission, blood circulation, and immune response. The human microbiota reportedly plays a major role in maintaining the overall health. Moreover, microbial dysbiosis has been linked to obesity, malnutrition, neurological disorders, behavioral disorders, and cancer.²

Major depressive disorder (MDD) is a common mental illness characterized by significant and persistent depression, loss of interest, and anhedonia.³ Recent epidemiological surveys have shown that more than 300 million people worldwide suffer from MDD,⁴ and approximately 1 million people die from MDD every year. Depressive disorders seriously affect the physical and mental health of patients and are a major burden on the global society.^{5,6} Previous studies have reported that the occurrence and development of MDD are associated with multiple factors such as

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neurotransmitters, endocrine, imaging, and immunity. However, the pathogenesis of MDD still remains unclear. In recent years, research on microbiota has brought a new perspective to the study of mechanisms of depressive disorders. There is substantial evidence that gut microbiota play an important role in the occurrence and development of MDD through the gut-brain axis,^{7–9} and that imbalance in gut microbiota can result in the occurrence of MDD.^{10,11} Additionally, previous studies have shown that probiotics can alleviate MDD by modulating gut microbiota.^{12,13}

The oral cavity is the second largest microbiome after the gut tract. Studies have elucidated a relationship between oral microorganisms and diseases such as diabetes,¹⁴ premature birth,¹⁵ cardiovascular diseases,¹⁶ and Alzheimer's disease.¹⁷ However, only a few studies have investigated the role of oral microbiota in MDD. Studies have shown that oral administration of *Porphyromonas gingivalis* can change the composition of gut microbiota by reducing the gut barrier function and modulating the gut immune system.¹⁸ Accumulating evidence shows that oral microorganisms can change the gut microbiota by invading the intestine and resulting in microecological imbalance.¹⁹ Therefore, delving into the study of oral microbiota is also a potential research direction and would be of great value in elucidating the etiology of MDD.

The rodent model is a key tool for studying the biological mechanisms of mental illness. Its advantage is that it can nullify the influence of sex, age, diet, and genetics. Moreover, it maintains a highly controlled environment and reduces the variation among individuals in terms of oral and gut microbiota.²⁰ Chronic unpredictable mild stress (CUMS) is a rodent model of depression. It is currently the most commonly used, reliable and effective model.²¹ In this study, a CUMS depression rat model was constructed, and 16SrRNA sequencing method was used to compare the compositional differences in the oral and gut microbiota of rats between the CUMS and control groups to provide a certain reference for elucidating the microbiological etiology of MDD.

Materials and Methods

Animals

A total of 20 male adult Sprague-Dawley (SD) rats (6 weeks old), weighing 180 ± 20 g, were purchased from Beijing Vitong Lever Experimental Animal Technology Co., Ltd. [license number: SCXK (Beijing) 2021–0006]. Rats were raised in SPF-level animal rooms under controlled temperature (22 ± 2 °C) and humidity (50%–60%), with a 12:12-h light: dark cycle. Real-time ventilation was used to keep the air fresh.

Animal Grouping and CUMS Procedure

The rats were randomly divided into two groups (n = 10/group): control and CUMS. Rats were individually housed in cages, and the CUMS model was established in accordance with the methods described in a previous study.^{21,22} The rats in the model group were exposed to CUMS to induce depressive-like symptoms. Stimulation methods included cold stimulation, heat stimulation, binding, noise stimulation, foreign body stimulation, tail clamping stimulation, dark stimulation, light stimulation, strobe light stimulation, moist padding, tilting the cage, fasting and food prohibition. One to two stimulation methods were randomly selected every day for a total of 28 days, and the methods were not repeated for three consecutive days.

Body Weight and Behavioral Tests

Body weight(g) of the 20 rats was measured before and after modeling. The rats were subjected to sucrose preference test (SPT) and open field test (OFT) to evaluate depressive-like behaviors.

SPT

The SPT is mainly used to evaluate anhedonia in rats.²³ Reduced sucrose preference is generally considered an indicator of anhedonia, which is a core symptom of depression. Prior to the test, all the rats were trained to consume 1% sucrose solution for two days, and on the third day, an SPT test was conducted. For the test, the rats were given free access to one bottle with 100 mL of pure water and another bottle with 100 mL of sucrose solution. The positions of the two water bottles were alternated every 12 h to avoid position preference. At the beginning and the end of the 24 h test, the solution

weights were measured, and the sucrose preference percentage was calculated using the following formula: Sucrose preference rate (%) = sucrose consumption/water + solution consumption \times 100%.

OFT

The OFT was used to assess the locomotor activity and exploratory behavior of the rats.²³ The OFT-100 open field experiment system consists of a box measuring 100 cm \times 100 cm \times 40 cm in dimension. Each rat was individually placed at the center of the box and observed for 5 min with an infrared camera. The total distance traveled(cm), average speed (cm/s), and number of rearing of each rat were recorded.

Sample Collection

After conducting the behavioral tests, the rats were anesthetized, and oral samples were collected using cotton swabs from the back of the tongue to the palate, buccal mucosa, upper and lower vestibules, and floor of the mouth for 30s. Fresh fecal samples from the cecum of each rat were collected using sterile cotton swabs, placed in a sterile sampling tube, and quickly transferred to a -80° C freezer.

16S rRNA Gene Amplicon Sequencing

Total genomic DNA samples were extracted using the OMEGA Soil DNA Kit (M5635-02), following the manufacturer's instructions and stored at -20 °C before performing any further analysis. The quantity and quality of extracted DNAs were measured using a NanoDrop NC2000 spectrophotometer and agarose gel electrophoresis, respectively. PCR amplification of the bacterial 16S rRNA genes V3–V4 region was performed using the forward primer 338F (5'-ACTCCTACGGGAGGCAGCA-3') and the reverse primer 806R (5'-GGACTACHVGGGTWTCTAAT-3'). The PCR components comprised: 5 μ L buffer (5×), 0.25 μ L Fast pfu DNA Polymerase (5U/ μ L), 2 μ L (2.5 mM) dNTPs, 1 μ L (10 uM) of each Forward and Reverse primer, 1 μ L DNA Template, and 14.75 μ L ddH2O. Thermal cycling consisted of initial denaturation at 98 °C for 5 min, followed by 25 cycles consisting of denaturation at 98 °C for 30s, annealing at 53 °C for 30s, and extension at 72 °C for 45s, with a final extension of 5 min at 72 °C.

Bioinformatics Analysis

Operational taxonomic unit (OTU) clustering was performed using QIIME 2 software (V2019.1),²⁴ and the OTU representative sequences were compared with the Greengenes database for species information annotation. Alpha and beta diversity metrics were calculated using QIIME 2 (V2019.1). Beta diversity analysis was performed using principal coordinate analysis (PCoA) based on Jaccard distance. Linear discriminant analysis effect size (LEfSe) was used to identify the differential genera between two groups. Spearman correlation analysis was used to assess the correlation, and data are shown as a heat map using the GenesCloud program (https://www.genescloud.cn/chart/CorHeatmap).

Statistical Analysis

Statistical analysis and graphing of behavioral data were performed using GraphPad Prism 8.0. When the data satisfied the normal distribution and the variances were homogeneous, an unpaired *t* -test was used. When the data did not satisfy the normal distribution, non-parametric test was used. Their correlation was evaluated using Spearman correlation analysis, and P value <0.05 was considered statistically significant.

Results

Body Weight and Behavioral Tests

There was no significant difference in terms of body weight between the control and CUMS groups before modeling (P=0.0882; Figure 1A). After modeling, rats in the CUMS group showed significant weight gain compared to rats in the control group (P<0.0001; Figure 1B). The preference coefficient for sugar water in SPT significantly decreased in the CUMS group (P=0.0454; Figure 1C) in comparison to the control group. In the context of OFT, the total distance



Figure I Comparison of body weight and behavioral tests between in the two groups. (A) Baseline body weight of rats; (B) Body weight of rats after modeling; (C) Sugar water preference coefficient of rats; (D)Total distance of rats; (E) Average speed of rats; (F) Number of rearing of rats; (n=10), (*P<0.05, **P<0.01, ****P<0.0001).

(P = 0.0014; Figure 1D), average speed (P < 0.0001; Figure 1E), and number of rearing (P = 0.0034; Figure 1F) of the CUMS group was found to be significantly decreased than that of the control group.

The above results show that the CUMS group showed obvious depression-like symptoms, mainly anhedonia, reduced locomotor and exploratory behaviors, indicating that our depression model was successfully established.

Analysis of Alpha and Beta Diversity of Oral and Gut Microbiota in the Two Groups

In the alpha diversity analysis, the Chao1 index and Observed species analysis of the oral microbiota showed that there were significant differences in these two indexes between the control and CUMS groups (Chao1: P < 0.001, Observed Species: P < 0.01). However, no significant differences were observed in the gut microbiota (Figure 2).

In the beta diversity analysis, there was a significant separation in the oral and gut microbiota between the CUMS and control groups, indicating the rich diversity of microbiota composition (Figure 3).

Analysis of Oral and Gut Microbiota Composition in the Two Groups

The top three most abundant bacterial genus within the oral cavity included *Rothia* (control group 34.06% vs CUMS group 21.28%), *Psychrobacter* (control group 14.78% vs CUMS group 34.02%), and *Streptococcus* (control group 14.43% vs CUMS group 9.79%) (Figure 4). LEfSe analysis showed that the abundance of *Rothia* decreased and that of Psychrotrophs increased in the CUMS group, and the difference was statistically significant.



Figure 2 Difference of alpha diversity in the two groups. (A) Differences in alpha diversity of oral flora in the two groups. (B) Differences in alpha diversity of gut flora in the two groups. (O stands for oral flora, G stands for gut flora; Chao I and Observed species represent the bacterial richness; the larger the value is, the higher the bacterial richness), (n=10), (**P<0.01,***P<0.001).



Figure 3 Difference of beta diversity in the two groups using principal coordinate analysis (PCoA) based on Jaccard distance. The closer the projection distance of the two points on the coordinate axis, the more similar the community composition. (O stands for oral flora, G stands for gut flora), (n=10).

The top three most abundant bacterial genus in the gut microbiota included Lactobacillus (control group 26.62% vs CUMS group 20.27%), *Ruminococcus* (control group 2.64% vs CUMS group 2.51%) and *Oscillospira* (control group 2.29% vs CUMS group 1.72%) (Figure 4). LEfSe analysis showed that the abundance of *Ruminococcus* reduced in the CUMS group, and the difference was statistically significant.



Figure 4 The composition and relative abundance of oral and gut flora at the genus level in the two groups. (O stands for oral flora, G stands for gut flora).

LEfSe Analysis of Oral and Gut Microbiota in the Two Groups

At the genus levels, *Psychrobacter, Aerococcus, Desulfovibrio, Bacteroides, Lactococcus, Isobaculum, Akkermansia*, and *Arthrobacter* showed higher abundance in the oral microbiota of the CUMS group, while *Rothia, Facklamia, Paracoccus, Stenotrophomonas*, and *Vibrio* showed lower abundance (Figure 5A).

In the gut microbiota of the CUMS group, *Butyricicoccus* and rc4_4 showed higher abundance, while *Ruminococcus*, *Enterococcus*, *Streptococcus* and *Turicibacter* showed lower abundance (Figure 5B). All of the above differences were found to be statistically significant (P < 0.05).

Correlation Analysis Between Oral and Gut Differential Microbiota and Depressive-Like Behavior

Spearman correlation analysis was performed on the differential microbiota and depressive-like behavior. The results showed that *Rothia, Facklamia, Vibrio, Enterococcus*, and *Streptococcus* positively correlated with body weight and



Figure 5 Taxonomic biomarkers found by LEfSe in the two groups. (A) Taxonomic biomarkers found by LEfSe in oral flora. (B) Taxonomic biomarkers found by LEfSe in gut flora. (O stands for oral flora, G stands for gut flora).

negatively with total distance traveled, average speed, and number of rearing. *Psychrobacter* and *Aerococcus* positively correlated with the total distance traveled, average speed, and number of rearing, while negatively with body weight. *Desulfovibrio*, *Bacteroides* and *Lactococcus* positively correlated with body weight (P < 0.05) (Figure 6). The Spearman correlation coefficient and P value are shown in Table 1.

Correlation Analysis Between Oral and Gut Differential Microbiota

Spearman correlation analysis of oral and gut differential microbiota demonstrated a strong positive correlation between *Facklamia* in the oral cavity and *Enterococcus*, *Streptococcus* in the intestine (r = 0.64-0.73, P < 0.01); and a strong negative correlation between *Desulfovibrio* in the oral cavity and *Enterococcus*, *Turicibacter* in the intestine (r=-0.51--0.72, P < 0.05) (Figure 7). The Spearman correlation coefficient and P value are listed in Table 1.



Figure 6 Heat map of correlation between oral-gut differential flora and depressive-like behavior. (Red indicates positive correlation and blue indicates negative correlation), (*P<0.05, **P<0.01, ***P<0.001).

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	Rothia	Fackla mia	Vibrionace ae_Vibrio	Psychro bacter	Aeroco ccus	Desulfovi brio	Lactococ cus	Enteroco ccaceae	Streptoco ccus	Turici bacter	rc4_4	Body Weight	Sucrose Preference	Total Distance	Mean Speed	Number of Rearing
Rothia	1															
Facklamia	0.829**	1														
Vibrionaceae_Vibrio	0.630**	0.713**	1													
Psychrobacter	-0.869**	-0.711**	-0.507*	1												
Aerococcus	-0.684**	-0.824**	-0.724**	0.547*	1											
Desulfovibrio	-0.544*	-0.464*	-0.536*	0.162	0.589**	1										
Lactococcus	-0.378	-0.441	-0.450*	0.058	0.423	0.495*	1									
Enterococcaceae	0.558*	0.643**	0.709***	-0.438	-0.705***	-0.508*	-0.43	1								
Streptococcus	0.489*	0.732***	0.448*	-0.543*	-0.672**	-0.107	-0.295	0.620**	T							
Turicibacter	0.405	0.423	0.574**	-0.229	-0.677**	-0.719***	-0.139	0.452*	0.171	T						
rc4_4	-0.206	-0.346	-0.499*	0.244	0.171	0.071	0.317	-0.493*	-0.271	0.006	T					
Body weight	0.659**	0.702***	0.602**	-0.445*	-0.695***	-0.609**	-0.659**	0.708***	0.517*	0.321	-0.394	I				
Sucrose preference	-0.108	-0.143	-0.471*	0.063	0.423	0.337	0.263	-0.344	-0.07 I	-0.39	0.29	-0.295	I			
Total distance	-0.638**	-0.669**	-0.711***	0.552*	0.636**	0.36	0.325	-0.610**	-0.623**	-0.32I	0.323	-0.608**	0.209	1		
Mean speed	-0.638**	-0.669**	-0.711***	0.552*	0.636**	0.36	0.325	-0.610**	-0.623**	-0.321	0.323	-0.608**	0.209	1.000**	1	
Number of rearing	-0.599***	-0.688**	-0.635**	0.713***	0.609**	0.15	0.022	-0.448*	-0.561*	-0.367	0.256	-0.452*	0.232	0.682**	0.682**	I

 Table I The Spearman Correlation Coefficient Value and P value

Notes: *P<0.05, **P<0.01, ***P<0.001.



Figure 7 Heat map of correlation between oral differential flora and gut differential flora. (Red indicates positive correlation and blue indicates negative correlation), (*P<0.05, **P<0.01, ***P<0.001).

Discussion

Characteristics of Gut Microbiota in Model Rats with CUMS-Induced Depression

Microbiota play a vital role in human health. Recent studies have shown that gut microbiota can affect the host brain function and behavior through the gut-brain axis.^{7,8} In the present study, we found that the relative abundance of *Ruminococcus, Enterococcus, Streptococcus* and *Turicibacter* in the gut microbiota of the CUMS group was reduced. Cheung et al reported that the relative abundance of *Anaerostipes, Blotella, Clostridium, Clostridium, Parabacteroidetes, Coralibacter*, and *Streptococcus* increased in patients with MDD, while that of *Bifidobacterium, Microbacterium, Faecalibacterium* and *Ruminococcus* decreased.⁷ Sanada et al conducted a meta-analysis of 10 studies and showed that the relative abundance of *Faecalibacterium, Ruminococcus, Bifidobacterium, and Escherichia* reduced in patients with MDD.²⁵ The results of the two abovementioned studies are partly similar to our findings. Studies have also shown that *Candida, Streptococcus, Escherichia*, and *Enterococcus* can produce 5-hydroxytryptamine (5-HT), which is an inhibitory neurotransmitter and the reduction of which can lead to the occurrence of MDD.²⁶ Gut microbiota and their products can directly affect the brain nervous system through the enteric nervous system and blood flow, and also indirectly mediate the occurrence of MDD through systemic inflammation.²⁷

Characteristics of Oral Microbiota in Model Rats with CUMS-Induced Depression

The oral cavity harbors the second largest microbiome after the gut tract and is close to the brain, making it more likely to cause damage to the cranial nerves. Wingfield et al examined the composition of the salivary microbiota of patients with MDD and found that the abundance of *Rothia, Haemophilus, Prevotella, Treponema*, and *Neisseria* was significantly depleted in comparison to the control group.²⁸ This study found significant differences in the α and β diversity of oral microbiota between the control and CUMS group. The CUMS group had the highest content of *Rothia* in the oral microbiota. *Rothia* belongs to the phylum *Firmicutes*, class *Clostridiales*, order *Clostridiales*, and family *Lachnospiraceae*. It is a Gram-positive, obligate anaerobic bacterium and a natural colonizer of the oral cavity. The *Rothia* genus comprises several species, including *R. dentocariosa, R. mucilaginosa* and *R. aeria. Rothia* can lead to bloodstream infections, endocarditis or other infections in immunocompromised patients. Ranga et al found that the *R. mucilaginosa* in the oral cavity produces enterobactin, which inhibits the growth of cariogenic *Streptococcus* mutans and pathogenic *Staphylococcus aureus* by chelating iron ions, which can cause oral microbiota imbalance.²⁹ *Rothia* can

also colonize the gastrointestinal tract and produce short-chain fatty acids. Studies have confirmed that *Rothia* in the intestine plays an important role as an anti-inflammatory bacterium in MDD.³⁰⁻³³

Interactions Between Oral and Gut Microbiota

In addition, we also found a correlation between the differential oral and gut microbiota, suggesting that the oral-gut microbiota axis may be involved in the pathogenesis of MDD. Previous studies have shown that there is a significant correlation between the microbial species present in the saliva and feces, and both have the same dynamics.^{34,35} Valles-Colomer et al studied more than 9700 human metagenomes and determined the human-to-human transmission of oral and gut microbiota, suggesting that the oral-gut microbiota axis plays an important role in the related diseases.³⁶ Jun Qian et al found that periodontitis salivary microbiota can exacerbate depressive-like behavior by directly affecting the host gut microbiota,³⁷ through oral-gut microbiota axis. Although the mouth and intestines are anatomically connected by the gastrointestinal tract, in health, physical and chemical barriers (such as stomach acid and bile acids) separate the mouth from the intestines. However, impairment of the oral-gut barrier can lead to translocation and communication between the organs. Oral microbiota can translocate to the gut, and gut microbiota and spread to the oral cavity. This two-way interaction can change the ecosystem in both environments. Therefore, oral microbiota and their products can affect the brain function directly via the facial nervous system and bloodstream, or indirectly by being involved in gut dysbiosis and systemic inflammation.

Conclusion

In order to eliminate the influence of factors such as human sex, age, diet, and genetics, this study was the first to explore the characteristics of oral and gut microbiota of depressed rats exposed to CUMS, which was innovative. However, this analysis was limited to a cross sectional and did not take into account any changes in oral-gut microbiota dysbiosis and MDD over time. We found that the oral and gut microbiota of CUMS group were disordered, and the disordered microbiota may be the cause of MDD. In addition, we also found differences between the oral and gut microbiota, but there was also a degree of connection. Both may be involved in the occurrence of MDD, and modulating oral and gut microbiota may provide new options for treatment of MDD.

Ethics Statement

This study complies with the Chinese National Laboratory Animal-Guidelines for Ethical Review of Animal Welfare (GB/T35892-2018) and was approved by the Medical Ethics Committee of First Hospital of Shanxi Medical University. The approval number is DWYJ-2023-148.

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Disclosure

All authors declare that they have no conflicts of interest for this work.

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