

Intestinal microbiome dysbiosis increases *Mycobacteria* pulmonary colonization in mice by regulating the *Nos2*-associated pathways

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Abstract Increasing researches reveal gut microbiota was associated with the development of tuberculosis (TB). How to prevent or reduce *Mycobacterium tuberculosis* colonization in the lungs is a key measure to prevent TB. However, the data on gut microbiota preventing *Mycobacterium* colonization in the lungs were scarce. Here, we established the clindamycin-inducing intestinal microbiome dysbiosis and fecal microbial transplantation models in mice to identify gut microbiota's effect on *Mycobacterium*'s colonization in the mouse lungs and explore its potential mechanisms. The results showed that clindamycin treatment altered the diversity and composition of the intestinal bacterial and fungal microbiome, weakened the trans-kingdom network interactions between bacteria and fungi, and induced gut microbiome dysbiosis in the mice. Gut microbiota dysbiosis increases intestinal permeability and enhances the susceptibility of *Mycobacterium* colonization in the lungs of mice. The potential mechanisms were gut microbiota dysbiosis altered the lung transcriptome and increased *Nos2* expression through the 'gut-lung axis'. *Nos2* high expression disrupts the intracellular antimicrobial and anti-inflammatory environment by increasing the concentration of nitric oxide, decreasing the levels of reactive oxygen species and *Defb1* in the cells, and promoting *Mycobacteria* colonization in the lungs of mice. The present study raises a potential strategy for reducing the risks of *Mycobacteria* infections and transmission by regulating the gut microbiome balance.

eLife Assessment

This **important** study shows the effect of gut dysbiosis on the colonization of mycobacteria in the lung. The data with comprehensive analysis of gene expression profiles in the lung with dysbiotic mice is **compelling** and goes beyond the current state of the art. However, the mechanistic insight, where the lung epithelial cell line was used, and the experiments with Mtb infection are currently **incomplete**.

Introduction

Tuberculosis (TB) is a chronic infectious disease caused by *Mycobacterium tuberculosis* (Mtb), with approximately 10 million people being infected each year and 1.6 million deaths in 2021 (Furin et al., 2019; Bagcchi, 2023). The epidemic of TB has brought heavy economic and social burdens to countries around the world. According to the World Health Organization (WHO) Global TB Report

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2022, the global spending on essential TB services for TB diagnosis, treatment, and prevention has exceeded \$6.4 billion (*Global tuberculosis report 2023, 2023*). Besides, drug-resistant forms of Mtb are currently on course to be the world's deadliest pathogens, responsible for a quarter of deaths due to antimicrobial resistance (*Furin et al., 2019*). The prevalence of drug-resistant Mtb is also a major threat to the control of TB. Hence, it has been an urgent task for researchers to develop more effective measures for the prevention and treatment of TB.

Recent advances in gut microbiota explorations have led to improved knowledge of the 'gut-lung axis'. Some studies have shown that intestinal microbiota dysbiosis may affect the occurrence and development of respiratory diseases including asthma, chronic obstructive pulmonary disease, respiratory pathogens infection, etc. by the 'gut-lung axis' (*Keely et al., 2012; Barcik et al., 2020; Wang et al., 2023*). The documents about the relationship between gut microbiota and TB have increased in recent years, and increasing evidence supports the presence of intestinal microbiota dysbiosis in TB patients. A case-control study of pulmonary TB in pediatric patients showed that the abundance of *Prevotellaceae* and *Enterococcaceae* was increased, and the abundance of *Oscillospiraceae* and *Bifidobacteriaceae* was decreased in TB patients compared with healthy control subjects (*Li et al., 2019*). Maji et al. have found that the levels of *Prevotella* decreased and *Bacteroides* increased in TB patients (*Maji et al., 2018*). Recently, our teams' study in active TB patients showed the abundance of *Bacteroidetes* significantly altered compared with health subjects (*Han et al., 2024*). Another study also showed the phylum of *Bacteroidetes* altered in the feces samples of patients with recurrent TB (*Luo et al., 2017*). All these researches indicated that Mtb infection can lead to the alteration of intestinal microbiota composition.

However, the causal relations between Mtb infection and intestinal microbiota dysbiosis have not been fully elucidated. Some factors including alcohol, smoking, HIV infection, and diabetes have been proven to cause gut microbiota dysbiosis, and these also are important risk factors for TB (*Naidoo et al., 2019*). A recent study has shown that the mice fed with a high-fat diet-induced gut microbiota dysbiosis increased the risk of developing active TB in mice (*Arias et al., 2019*). Another study found *Firmicutes/Bacteroidetes* ratio increased in a murine model of type 2 diabetes induced by an energy-dense diet, and increased the susceptibility to Mtb infection (*Sathkumara et al., 2021*). These findings support a viewpoint that the changes in the gut microbiome are a contributing factor in Mtb infection pathogenesis and its clinical presentation. Many documents supposed that gut microbiota dysbiosis could modulate immunity and inflammation reaction at distal sites such as the lungs, reduce colonization resistance by external pathogens, and promote TB susceptibility (*Naidoo et al., 2019; Osei Sekyere et al., 2020*). Hence, modulation of the gut microbiota and balance of the gut-lung axis was a potential avenue for TB prevention and management.

Based on our previous studies and other reports, we found the effect of Mtb infection on the gut *Bacteroidetes* was most significant. Hence, in the present study, clindamycin (CL), an antibiotic that selectively disorders anaerobic *Bacteroidetes* (*You et al., 2019*), was used to establish a mouse model of intestinal microbiota dysbiosis. We uncovered the important role of gut microbiome dysbiosis affecting *Mycobacteria* colonization in the mouse lungs using the technology of fecal microbiota transplantation (FMT). Furthermore, based on the 'gut-lung axis' theory, we performed a transcriptomic analysis of the mice lungs after *Mycobacteria* infection. We aim to explore the potential role and mechanism of gut microbiome dysbiosis enhancing *Mycobacteria* colonization in the lungs of mice.

Results

Clindamycin increases *Mycobacterium smegmatis* colonization in the lungs of mice

According to **Figure 1a** procedure, we established a clindamycin-treated mice model to assess the effects of clindamycin-associated gut dysbiosis on *Mycobacterium smegmatis* (MS) colonization in the lungs of mice. The results showed that the mice with clindamycin treated presented a significantly higher colonization of MS in the lungs than that of control mice (**Figure 1b, Figure 1—source data 1**). The size of the cecum of CL-treated mice was markedly more dilated compared with the control groups (**Figure 1c, Figure 1—source data 1**). To assess the effect of CL on gut mucosal damage and permeability, the intestinal fatty acid binding protein (i-FABP, a marker of enterocyte death) and lipopolysaccharides (LPS, an endotoxin, as a marker of gut permeability) in serum were investigated

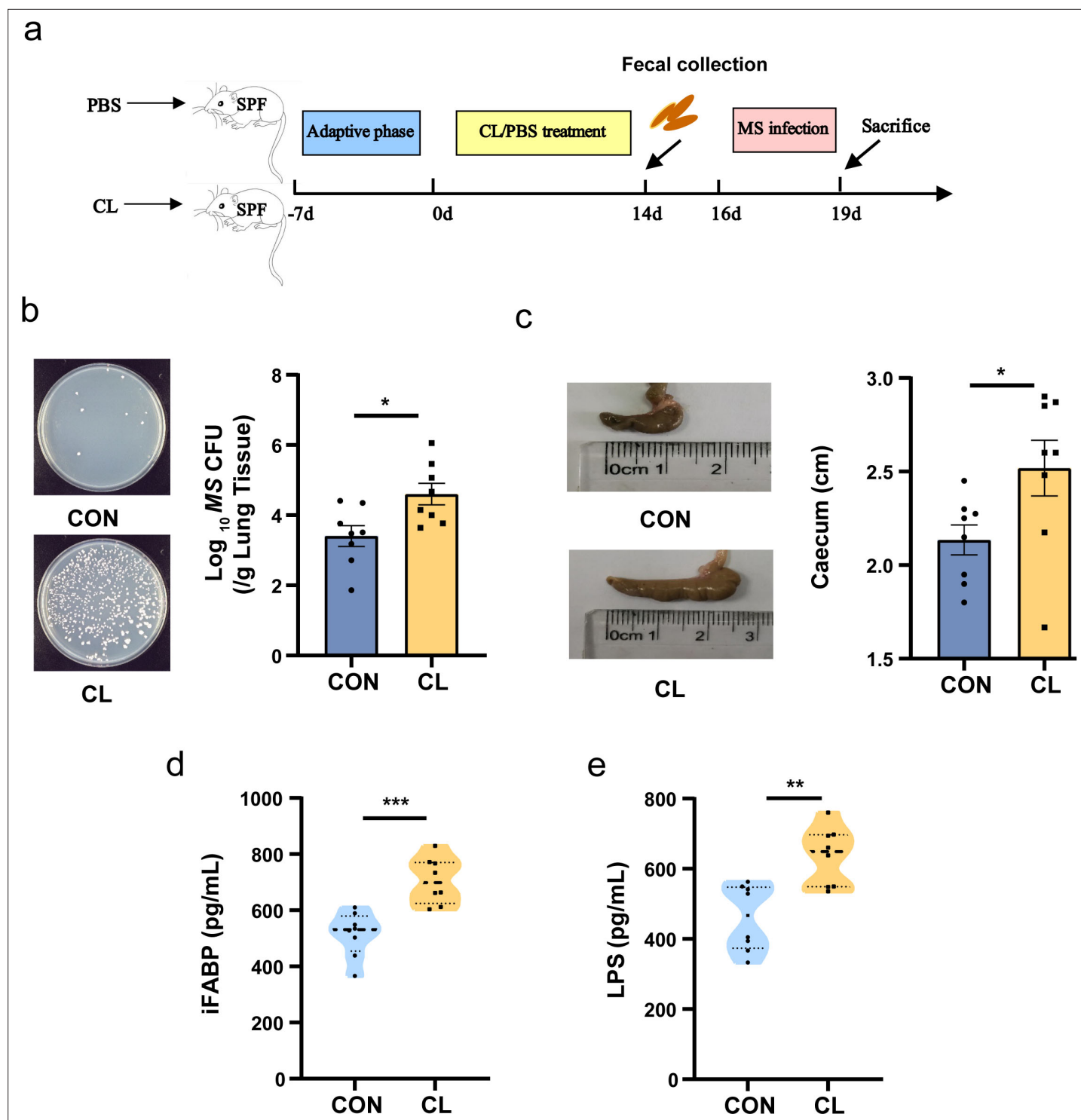


Figure 1. Clindamycin increases MS pulmonary colonization in mice. **(a)** The experimental procedures of clindamycin-inducing gut microbiota dysbiosis model in mice. **(b)** The colonization of MS in the lungs of mice after infection at 72 hr. **(c)** The length of cecum after clindamycin treatment. **(d)** The level of iFABP in serum. **(e)** LPS concentration in serum. CL: clindamycin treatment group ($n = 8$), CON: control group ($n = 8$). MS: *Mycobacterium smegmatis*, iFABP: intestinal fatty acid binding protein, LPS: lipopolysaccharides. * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$.

The online version of this article includes the following source data and figure supplement(s) for figure 1:

Source data 1. Effect of clindamycin on MS pulmonary colonization and intestinal permeability of mice.

Figure supplement 1. Pathological sections of intestinal and lung tissue of mice with CL treatment.

with enzyme-linked immunosorbent assay (ELISA) technology. The results showed that the levels of i-FABP and LPS were significantly higher in the CL group than in the control group (CON) (**Figure 1d, e, Figure 1—source data 1**). Pathological sections of intestinal tissue showed the intestinal epithelial tissues of mice with no obvious alteration after clindamycin treatment vs the CON group (**Figure 1—figure supplement 1**). H&E staining of paraffin-embedded lung sections revealed that there was more diffused inflammation and inflammatory cell infiltration in the CL-treated mice compared with that of the CON group animals after infecting with MS (**Figure 1—figure supplement 1**). All these results indicated that CL treatment damages the enterocyte, increases gut permeability, promotes the fermentation of cecum contents, and increases MS colonization in the lungs of mice.

Altered the diversity and composition of the gut microbiome in mice

To assess the effect of CL treatment on the gut microbiome of mice, we investigated the alteration of gut microbiota (bacterial) and mycobiota (fungi) by 16S rRNA and ITS2 amplicon sequencing, respectively. For the gut bacterial analysis, the α -diversity was significantly decreased in the CL group compared with the CON group (**Figure 2a, Figure 2—source data 1**). The β -diversity based on the weighted UniFrac distance showed that the CL group samples were separated from the CON group ($R = 0.8488$, $p = 0.001$) (**Figure 2b**). It indicated that the intestinal bacterial microbial community structure was a significant difference between the CL-treated mice and the control mice. At the phylum level, we observed a remarkable decrease of *Bacteroidota* and a significant increase of *Firmicutes* and *Proteobacteria* in the CL group compared to the control group (**Figure 2c, Supplementary file 1**). At the genus level, the CL group had a significantly higher relative abundance of *Bacteroides*, *Lactobacillus*, *Escherichia-Shigella*, and *Faecalibaculum*, and a lower abundance of *Dubosiella*, *Alloprevotella*, *Akkermansia*, and *Bifidobacterium* compared with that in the CON group (**Figure 2d, Supplementary file 1**).

For the fungal mycobiota analysis, the Chao1 and inverse Simpson indices of the CL group was significantly increased compared to that of the CON group (**Figure 2a**). The principal coordinate analysis (PCoA) showed that the samples were significantly distinguished between two groups ($R = 0.6038$, $p = 0.001$) (**Figure 2b**), which suggested that the community of intestinal mycobiota in the two groups is different. At the phylum level, the fungal microbiota was dominated by the Ascomycota and Basidiomycota in two groups, and Ascomycota showed no obvious differences between the two groups. However, the relative abundance of Basidiomycota significantly increases in the CL group compared with the CON group (**Figure 2c, Supplementary file 1**). The LEfSe analysis at the genus level showed that the relative abundance of *Aspergillus*, *Penicillium*, *Cladosporium*, and *Alternaria* was enriched in the CL group vs the CON group (**Figure 2d, Supplementary file 1**). These results are consistent with the Wilcoxon rank-sum test analysis (**Figure 2—figure supplement 1**). Taken together, all these data indicated that CL treatment altered the diversity and composition of the intestinal microbiome (including bacterial and fungi) in the mice and induced gut microbiome dysbiosis.

Clindamycin weakens the trans-kingdom networks construction of gut bacterial and fungi

To evaluate the effect of clindamycin treatment on gut microbiome balance, we analyzed the fungi-to-bacteria species ratio with ITS/16S. The results show that the ITS/16S ratio of the CL group was significantly increased compared with the CON group (**Figure 3a, Figure 3—source data 1**). We also assessed the alterations in the ratio of the dominant phyla in the gut microbiome. The results showed that the ratio of *Firmicutes/Bacteroidota* in the CL group was increased, and *Ascomycota/Basidiomycota* was significantly decreased compared with that of the CON group (**Figure 3b, c, Figure 3—source data 1**). These results revealed that clindamycin treatment not only disturbed the intestinal equilibrium between bacteria and fungi but also destroyed the balance among the dominant bacteria (or fungi). Then, to assess the interplay between bacteria and fungi, we performed the trans-kingdom networks analysis at the genus level. The results showed that the trans-kingdom networks were altered in the CL group vs that in the CON group. In the control group, the bacteria and fungi were closely related to each other, gathering in a cluster and forming a more complex network (**Figure 3d**). There were 154 nodes including 45 fungi and 109 bacteria, and 810 edges; the relative connectedness was 5.26 (**Table 1**). However, the complexity of the trans-kingdom network in the CL group was dramatically decreased, and the interplay between bacteria and fungi was also

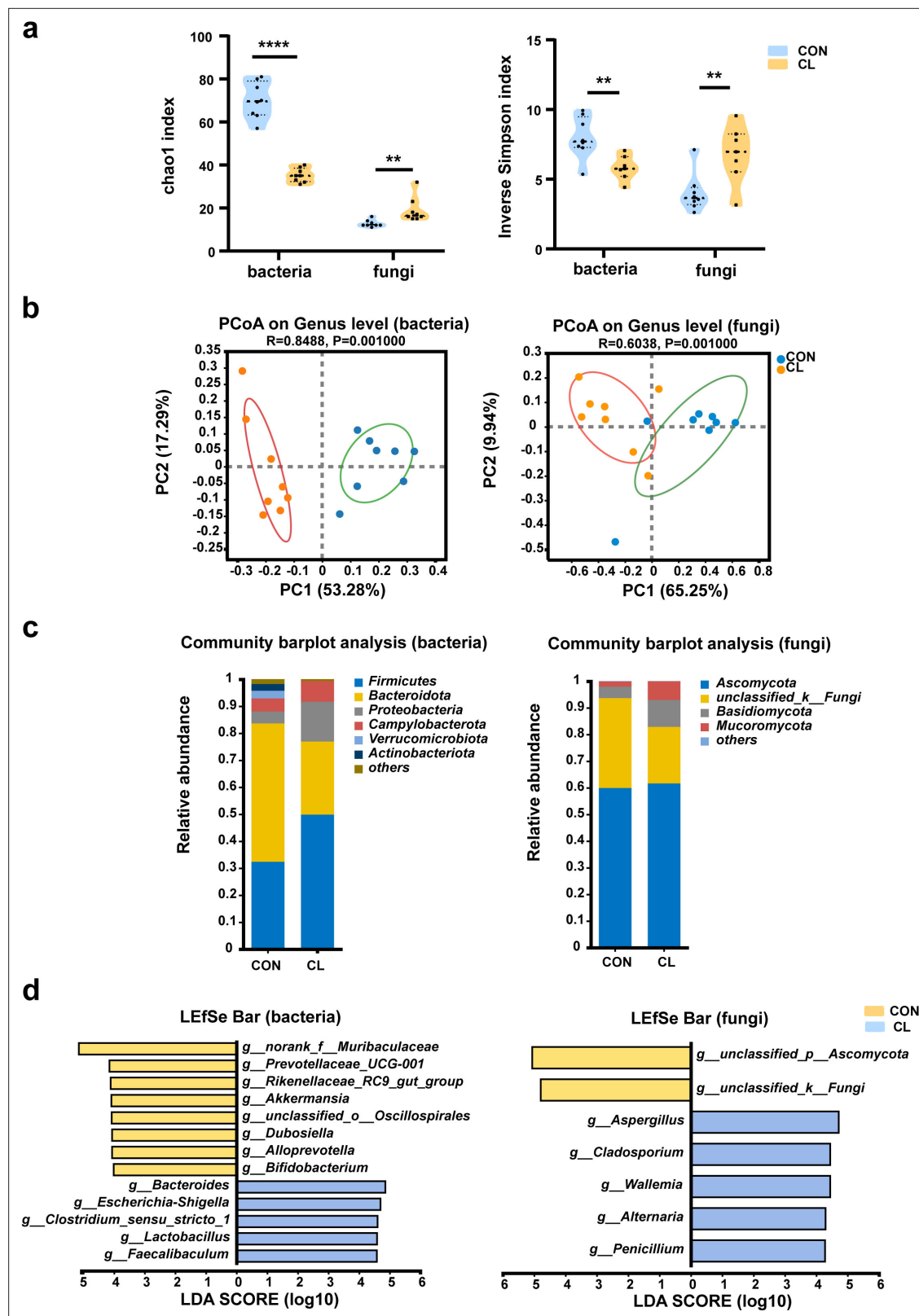


Figure 2. Clindamycin altered the diversity and composition of the gut microbiome in mice. (a) Chao1 and inverse Simpson indices of gut microbiota. (b) Principal coordinate analysis (PCoA) of gut microbiota based on weighted Unifrac distance. (c) The composition distribution of gut microbiome (bacteria and fungi) at the phylum level. (d) The linear discriminant analysis effect size (LEfSe) analysis of the differentially abundant gut microbiota

Figure 2 continued on next page

Figure 2 continued

between the CL and CON groups at the genera level (linear discriminant analysis [LDA] > 4, $p < 0.05$). CL: clindamycin treatment group ($n = 8$), CON: control group ($n = 8$). ** $p < 0.01$, **** $p < 0.0001$.

The online version of this article includes the following source data and figure supplement(s) for figure 2:

Source data 1. Alpha diversity index of gut microbiome.

Figure supplement 1. The Wilcoxon rank-sum test of gut fungal relative abundance difference between the CL and CON groups at the genera level.

weakened (**Figure 3e**). A total of 110 nodes including 62 fungi and 48 bacteria gathered into clusters in this network. The edges decreased to 357 and the relative connectedness was 3.25 in the CL group network (**Table 1**). We also found that the interplay ratio between fungi in the CL group was 52.10%, which was much higher than that of the CON group (7.53%) (**Table 1**). All these results indicated that clindamycin treatment not only altered the proportion of bacteria and fungi, but also weakened the trans-kingdom networks, changed the interactions between bacteria and fungi, and resulted in gut microbiome dysbiosis.

Fecal microbial transfer from clindamycin-treated mice donors to antibiotics-treated conventional mice enhances MS colonization

We performed an FMT model to further verify the relationship of gut microbiota dysbiosis with the susceptibility to MS infection in the mice's lungs. The experimental design for the FMT is shown in **Figure 4a**. First, we analyzed the α -diversity of gut bacterial and fungal microbiota after FMT with the Chao1 and inverse Simpson indices. The results show that the α -diversity of gut bacterial microbiota was significantly decreased, and the α -diversity of gut fungal microbiota was significantly increased in the CL-FMT group compared with that in the CON-FMT group (**Figure 4—figure supplement 1**, **Figure 4—figure supplement 1—source data 1**). Then we performed PCoA analysis to distinguish the gut microbiome alteration between CL- and CON-recipient mice. The results show a significant separation of gut bacterial and fungal microbiota between CT- and AD-recipient samples ($p = 0.007$ and $p = 0.042$, **Figure 4b**). The fungal-to-bacterial species ratio was significantly increased in the CL- vs the CON-recipient groups ($p < 0.05$, **Figure 4c**, **Figure 4—source data 1**). To calculate the changes of the dominant phyla in the gut microbiome, the results show that *Firmicutes/Bacteroidota* in the CL-FMT group was decreased, and *Ascomycota/Basidiomycota* was no obvious different compared with that of the CON-FMT group (**Figure 4d, e**, **Figure 4—source data 1**). The trans-kingdom network analyses between bacteria and fungi showed the complexity of the microbiome network was significantly reduced and the interactions between bacteria and fungi were also weakened in the CL-FMT vs the CON-FMT groups (**Figure 4f**). The above results showed that the trends of the gut microbiome in recipient mice were consistent with those in the donor mice. Then, the Venn diagram was used to assess the gut microbiome profile of FMT mice at the genus level. The results showed that 85.11% (40/47) of bacterial genera and 52.38% (33/63) of fungi genera present in the CL inocula were successfully transferred to the CL-recipient mice, and 91.45% (107/117) of bacteria genera and 56.36% (31/55) of fungi genera in the CON inocula were also successfully transferred to the CON-recipient mice, respectively (**Figure 4g**). These results indicated that the FMT model was successfully established, and the CL-recipient mice showed similar characteristics of gut microbiota dysbiosis with the CL group.

Then, we assessed the susceptibility of MS in the lungs of FMT mice, the results showed that the burden of MS in the lungs of CL-recipient mice was significantly rise than that in the CON-recipient mice after 72 hr of MS infection (**Figure 4h**, **Figure 4—source data 1**). However, the size of the cecum showed no significant differences between the two groups (**Figure 4—figure supplement 2**, **Figure 4—figure supplement 1—source data 1**). The level of iFABP and LPS also significantly increased in the CL-recipient mice compared with the CON-recipient mice after FMT (**Figure 4—figure supplement 2**, **Figure 4—figure supplement 1—source data 1**). The pathological sections of intestinal tissue showed the intestinal epithelial tissues of mice have no significant damage after FMT (**Figure 4—figure supplement 2**, **Figure 4—figure supplement 1—source data 1**). Altogether, our data suggested that gut microbiota dysbiosis increases intestinal permeability and enhances the susceptibility of MS colonization in the lungs of mice.

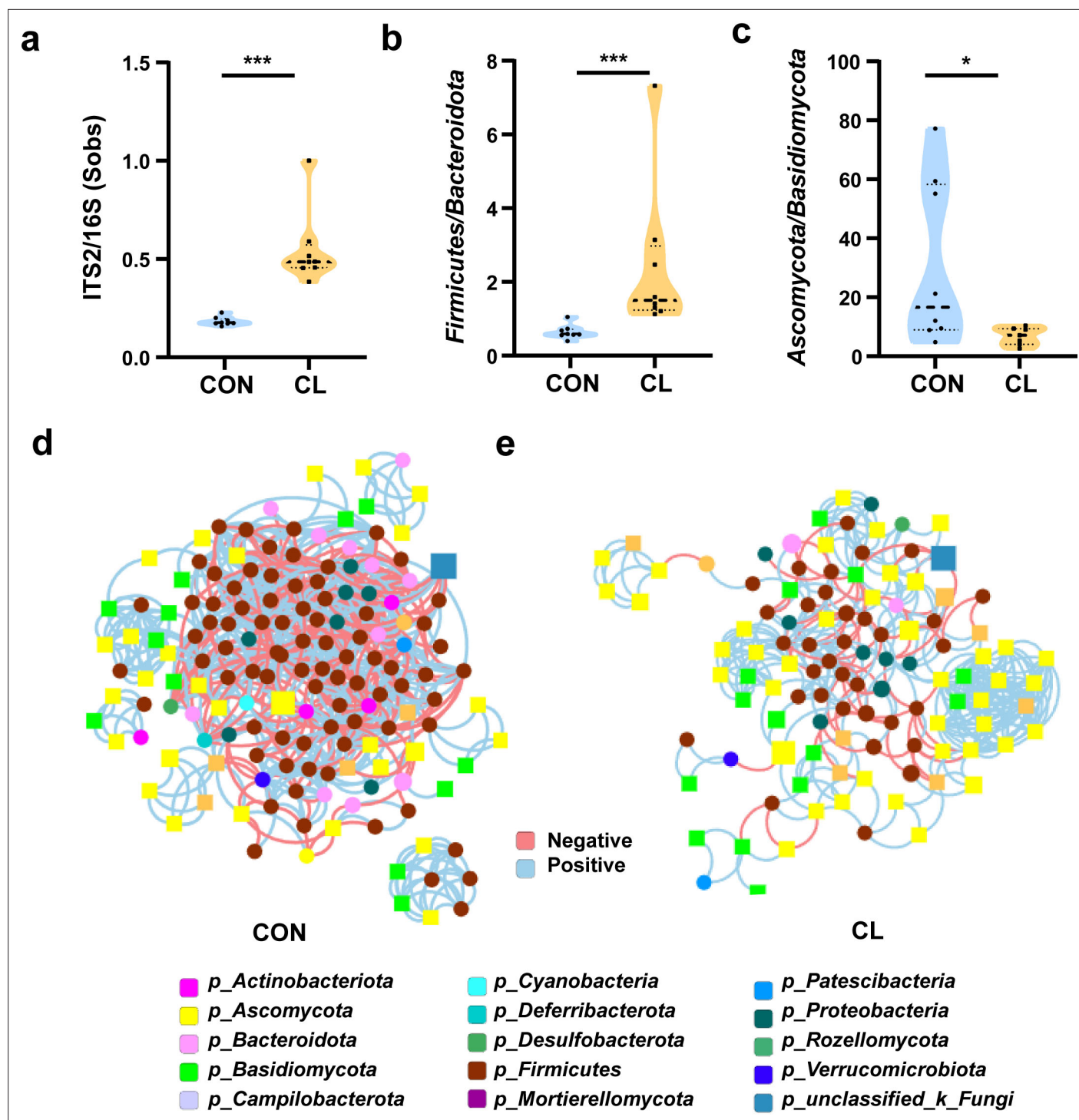


Figure 3. Clindamycin weakens the trans-kingdom network construction of gut bacteria and fungi. (a) The ratio of ITS2/16S at the genus level. (b) The relative abundant ratio of Firmicutes/Bacteroidota. (c) The relative abundant ratio of Ascomycota/Basidiomycota. (d) The trans-kingdom correlation networks of the CON at the genus level. (e) The trans-kingdom correlation networks of the CL at the genus level. CL: clindamycin treatment group ($n = 8$), CON: control group ($n = 8$). * $p < 0.05$; *** $p < 0.001$.

The online version of this article includes the following source data for figure 3:

Source data 1. Effects of clindamycin on the gut bacteria and fungi microbiota in mice.

Table 1. Parameters of the trans-kingdom abundance correlation networks.

	CON group	CL group
Nodes (n) (fungi/bacteria)	154 (45/109)	110 (62/48)
Edges (n)	810	357
Relative connectedness*	5.26	3.25
Bacteria–bacteria (%)	78.77	22.69
Bacteria–fungi (%)	13.70	25.21
Fungi–fungi (%)	7.53	52.10

*The relative connectedness of the network was calculated as the ratio of edges (the number of significant interactions) and nodes (the number of genera).

The gut microbiota dysbiosis altered the lung transcriptome and increased *Nos2* expression

To further explore the potential mechanisms by which intestinal microbiota dysbiosis affects MS colonization in the lungs of mice, we performed a transcriptome analysis of the mice’s lung tissue. The results showed that there were 1191 upregulated differentially expressed genes (DEGs) and 1013 downregulated DEGs in the CL vs the CON groups (false discovery rate [FDR] <0.05, and fold change [FC] >1, **Figure 5—figure supplement 1**). Compared with the CON-FMT group, 274 DEGs were upregulated and 32 DEGs were downregulated in the CL-FMT group (**Figure 5—figure supplement 1**). Then, we screened the overlapping DEGs between the two comparison sets including CL vs CON and CL-MFT vs CL-FMT. The Venn diagram showed that 93 upregulating DEGs and 5 downregulating DEGs were commonly expressed among these groups (**Figure 5a**). Subsequently, Gene Ontology (GO) and Kyoto Encyclopedia of Genes and Genomes (KEGG) pathway enrichment analyses were performed to clarify the function of these 98 DEGs. The top 30 biological processes enriched by GO showed that CL and CL-FMT groups mainly affected the immune response and inflammatory response, including ‘defense response’, ‘response to bacterium’, ‘cellular response to interleukin-1’, and ‘cellular response to lipopolysaccharide’ (**Figure 5b**). The enriched molecular functions of these DEGs were some cytokine and protease involving the immune defense response, including ‘cytokine activity’, ‘chemokine activity’, ‘nitric-oxide synthase binding’, and ‘CXCR chemokine receptor binding’ (**Figure 5b**).

The top 20 KEGG pathways enrichment analyses at classification level 1 indicated that these DEGs are mainly involved in environmental information processing, human diseases, and organismal systems (**Figure 5c** and **Supplementary file 2**). At classification level 2 of the KEGG analysis, we found six pathways relating to infectious disease, and seven pathways involving the immune system among the enrichment top 20 pathways (**Figure 5c** and **Supplementary file 2**). Interestingly, among the six pathways of infectious disease, there were three pathways involving bacterial infectious disease, including TB, legionellosis, and pertussis (**Figure 5c**). In addition, the pathway of ‘graft-versus-host disease’ is also enriched in these DEGs (**Figure 5c**).

For the future screening of the potential regulatory genes correlated with TB, nine KEGG pathways related to human disease were selected for interaction network analysis with DEGs. The results showed that eight genes were strongly associated with TB (**Figure 5d**). Among those genes, seven genes including *Nos2*, *Cd14*, *Tnf*, *Cd74*, *Clec4e*, *Ctsd*, and *Il6* are upregulating expression and the *Cd209a* gene is the downregulating expression in the CL and CL-FMT groups (**Figure 5—figure supplement 2**, **Figure 5—figure supplement 1—source data 1**). It is worth noting that the *Nos2* gene was an enriched expression in both GO and KEGG analyses. Then, we performed an RT-qPCR to assess the validity of the transcriptome. The results revealed that the genes of *Nos2*, *Ctsd*, *Cd74*, and *Tnf* were increased in CL-FMT and CL groups compared with that in control groups (**Figure 5e**, **Figure 5—source data 1**), which indicated that the RNA-seq data is reliable. In summary, our data revealed that gut microbiota dysbiosis significantly altered the transcriptomic profiling in the lung tissue of mice, and increased the expression of *Nos2* genes.

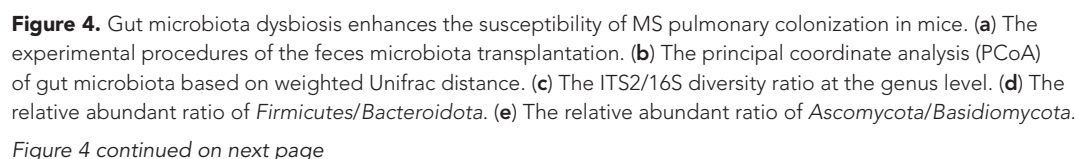


Figure 4 continued

(f) The trans-kingdom correlation networks of CON-FMT and CL-FMT groups at the genus level. (g) The Venn diagram of gut bacteria and gut fungi in different groups at the genus level. (h) The load of MS in the lungs of mice after infection at 72 hr. FMT: fecal microbiota transplantation, CON-FMT: the fecal microbiota of the control group was transplanted ($n = 6$), CL-FMT: the fecal microbiota of the clindamycin treatment group was transplanted ($n = 6$). MS, *Mycobacterium smegmatis*. * $p < 0.05$.

The online version of this article includes the following source data and figure supplement(s) for figure 4:

Source data 1. Effects of FMT on the gut bacteria and fungi microbiota in mice.

Figure supplement 1. The α -diversity analysis of gut microbiome after fecal microbiota transplantation (FMT).

Figure supplement 1—source data 1. Effects of FMT on the alpha diversity of gut microbiota and intestinal permeability in mice.

Figure supplement 2. The effect on gut mucosal damage and permeability after fecal microbiota transplantation (FMT).

Nos2 regulates the expression of nitric oxide, reactive oxygen species, and Defb1

Based on the above transcriptomic results, we speculate that the upregulating expression of *Nos2* induced by gut microbiota dysbiosis may play an important role in MS pulmonary colonization in mice. Hence, an overexpression *Nos2* vector, *Nos2*-pcDNA3.1, was constructed and transfected into A549 cells. Then the cells were infected with MS to verify the effect of *Nos2* overexpression on MS colonization ability. The results showed that the expression level of *Nos2* was significantly raised in A549 cells after transfection with the *Nos2*-pcDNA3.1 plasmid (**Figure 6a, Figure 6—source data 1**). MS colonization density was significantly increased in A549 cells with *Nos2*-pcDNA3.1 plasmid after infection MS 6, 12, and 24 hr (**Figure 6b, Figure 6—source data 1**), which suggested that *Nos2* overexpression increases the infection susceptibility of MS to A549 cells.

Subsequently, we explored the potential molecular mechanism by which *Nos2* overexpression increased MS colonization ability. *Nos2* is a key enzyme required for nitric oxide (NO) synthesis, so we detected the level of NO in A549 cells. The results showed that the NO level significantly increased in the A549 cells with transfection of the *Nos2*-pcDNA3.1 plasmid (**Figure 6c, Figure 6—source data 1**). Reactive oxygen species (ROS) play an important role in pathogens invade and colonize host organs (André et al., 2022). To explore whether *Nos2* regulates ROS production in A549 cells, ROS concentration in cells was detected by the fluorescent probe 2',7'-dichlorodihydrofluorescein diacetate (DCFH-DA). As fluorescence microscopy shown in **Figure 6d**, ROS concentration in A549 cells transfected with *Nos2*-pcDNA3.1 was significantly lower than that in the control A549 and A549 cells transfected with pcDNA3.1 empty vector (**Figure 6d, Figure 6—source data 1**). In addition, we also detected the expression of human β -defensin-1 (*Defb1*), an important antimicrobial peptide (Zhu et al., 2003), by RT-qPCR. The results showed that *Nos2* overexpression significantly reduced the level of *Defb1* in A549 cells (**Figure 6e, Figure 6—source data 1**). Taken together, our results revealed that *Nos2* overexpression disrupts the intracellular antimicrobial and anti-inflammatory environment by increasing the concentration of NO, decreasing the levels of ROS and *Defb1*, and resulting in the enhancement of MS pulmonary colonization capacity in mice.

Discussion

TB remains a major health challenge globally. How to effectively limit or reduce *Mtb* colonization in the host is a potential strategy to prevent this disease transmission and development. Recent studies have shown that the gut microbiome can affect distant organs via the 'gut-lung axis', and gut microbiota dysbiosis is a potential factor inducing respiratory diseases including TB (Waldman and Balskus, 2018; Ma et al., 2022). Previous studies have shown that the gut microbiota was significantly different in TB patients compared with healthy humans and revealed gut microbiota dysbiosis, especially *Bacteroidota*, and *Firmicutes*, is strongly associated with the development of TB (Maji et al., 2018; Luo et al., 2017). However, how the gut microbiota affects TB development by the 'gut-lung axis' remains unclear. In the present study, we established a mouse model of gut microbiota dysbiosis and an FMT model using clindamycin treatment and revealed the potential mechanisms of intestinal microbiota

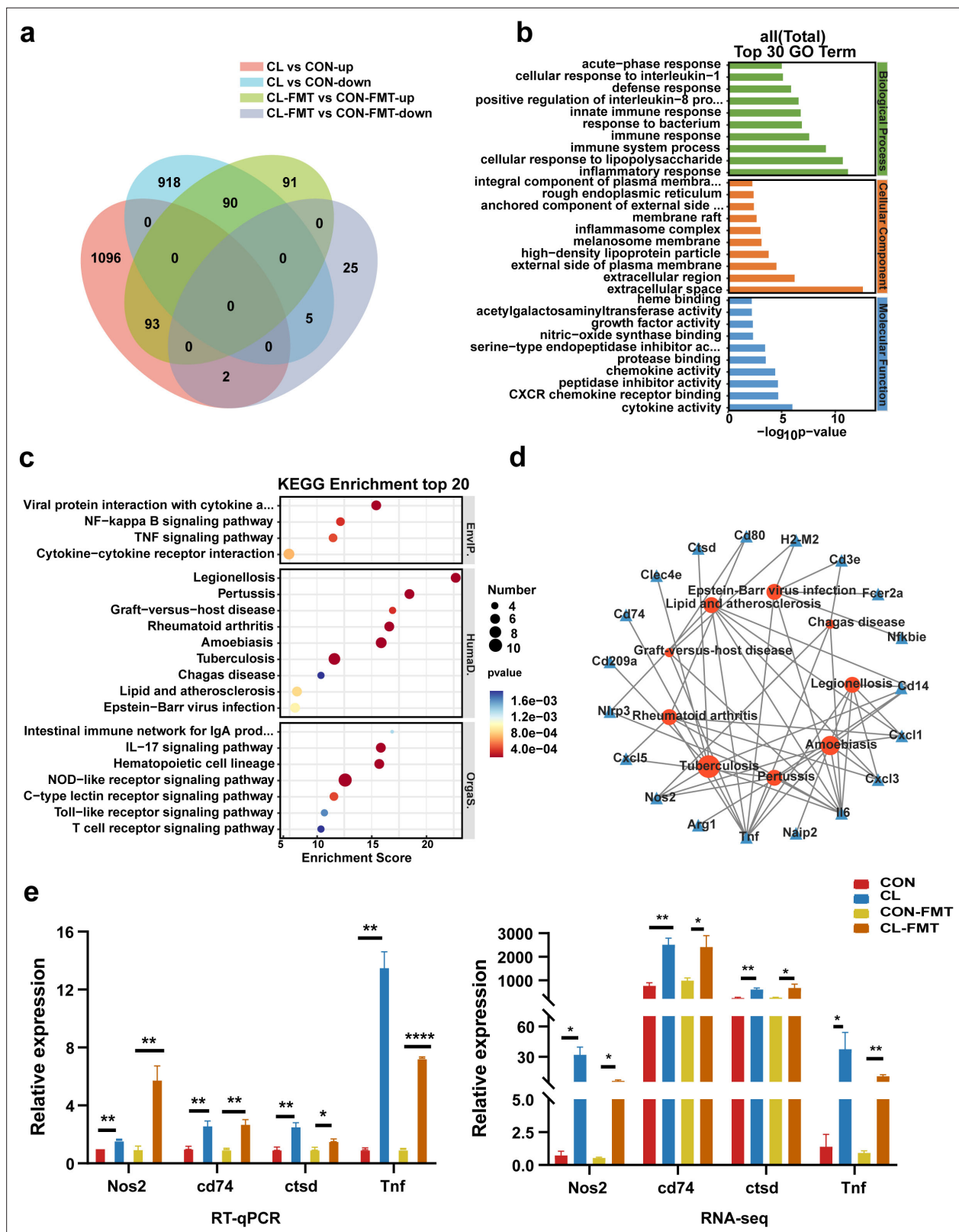


Figure 5. Gut microbiota dysbiosis altered the lung transcriptome of mice. (a) The Venn diagram of DEGs in the different groups. (b) The top 30 Gene Ontology (GO) enrichment analyses of DEGs. (c) The top 20 Kyoto Encyclopedia of Genes and Genomes (KEGG) enrichment analyses of DEGs. (d) Interaction network analysis of selected DEGs and significant KEGG pathways related to human diseases. (e) The expression levels of *Nos2*, *Ctsd*, *Cd74*, and *Tnf* with RT-qPCR and RNA-seq. DEGs: differentially expressed genes, CL: clindamycin treatment group, CON: control group; CON-FMT: the

Figure 5 continued on next page

Figure 5 continued

fecal microbiota of the control group was transplanted, CL-FMT: the fecal microbiota of the clindamycin treatment group was transplanted. * $p < 0.05$, ** $p < 0.01$, **** $p < 0.0001$.

The online version of this article includes the following source data and figure supplement(s) for figure 5:

Source data 1. The results of RT-qPCR and RNA-seq of *Nos2*, *Ctsd*, *Cd74*, and *Tnf*.

Figure supplement 1. The volcano plot of changes of differentially expressed genes.

Figure supplement 1—source data 1. RT-qPCR results of DEGs in the different groups.

Figure supplement 2. The expression levels of differentially expressed genes (DEGs).

dysbiosis promoting MS pulmonary colonization in the mouse by upregulating *Nos2* gene-associated pathways.

Antibiotics can induce intestinal microbiota dysbiosis, which in turn affects host immunity and leads to increased susceptibility and deterioration of a variety of diseases (Fishbein et al., 2023). Previous research has verified that gut *Bacteroidota* dysbiosis is strongly associated with the development of TB (Du et al., 2022; Huang et al., 2019; Yunusbaeva et al., 2024). Hence, we select clindamycin, an antibiotic that selectively disrupts anaerobic *Bacteroidota* (You et al., 2019), to treat the mice in this study. Our results show that the abundance of *Bacteroidota* was significantly reduced and the abundance of *Firmicutes* was increased in the CL group compared with the CON group. This result

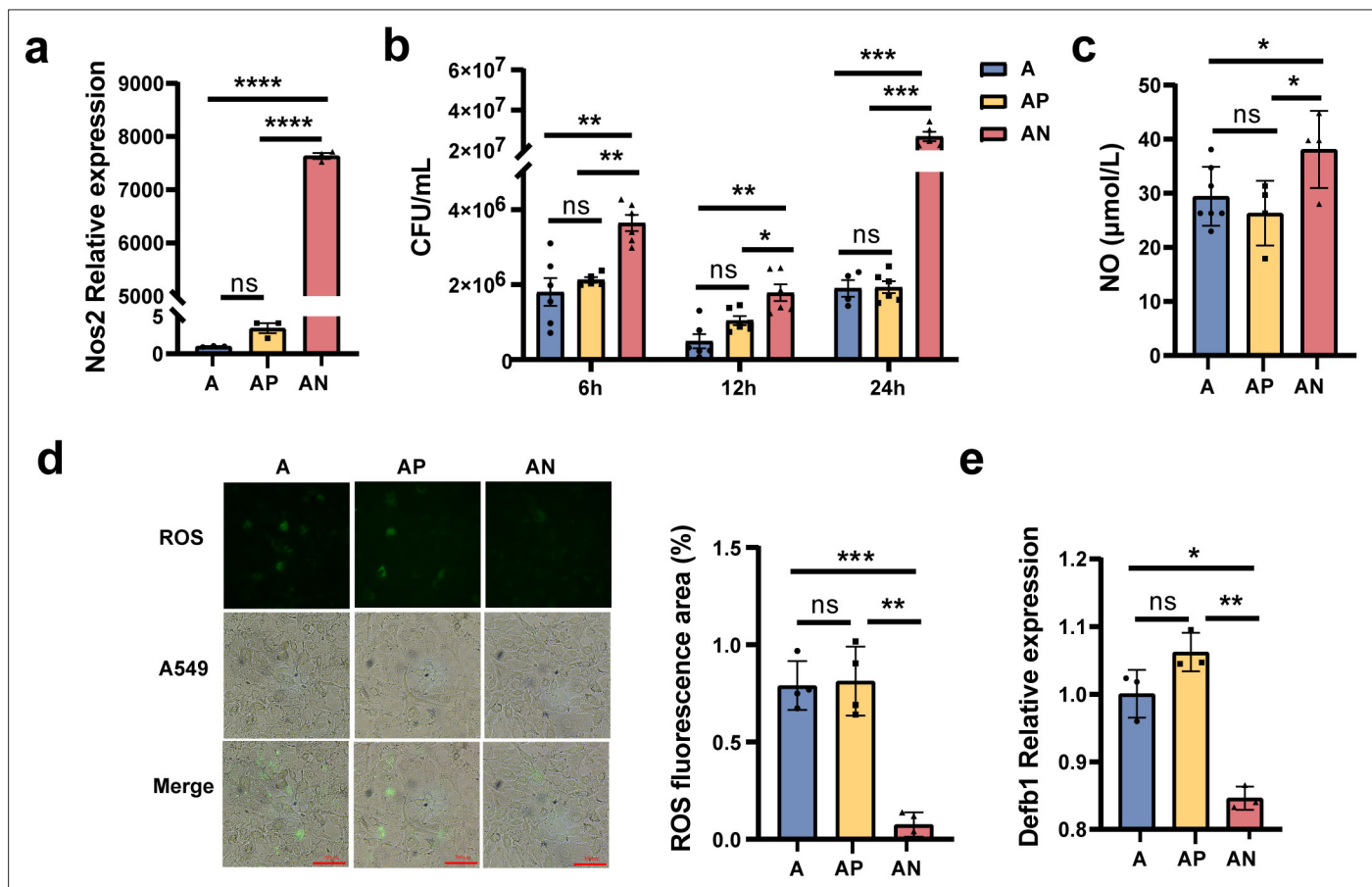


Figure 6. *Nos2* regulates the expression of nitric oxide (NO), reactive oxygen species (ROS), and *Defb1*. (a) The expression levels of the *Nos2* in the A549 cells. (b) The load of *Mycobacterium smegmatis* (MS) in the A549 cells at different times. (c) The NO concentration in the A549 cells. (d) The concentration of ROS in the A549 cells. (e) The expression levels of *Defb1* in A549. A: A549 cells control, AP: transfected A549 cells with blank pcDNA3.1 plasmid, AN: transfected A549 cells with *Nos2*-pcDNA3.1 plasmid. * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$, **** $p < 0.0001$.

The online version of this article includes the following source data for figure 6:

Source data 1. Effect of *Nos2* on the expression of nitric oxide (NO), reactive oxygen species (ROS), *Defb1* and the colonization of MS in cells.

is consistent with the previous study which reported that CL is effective in clearing *Bacteroidota* (You et al., 2019). *Bacteroidota* is a consortium of many commensal bacterial species responsible for major fermentation processes, glycolipid production, and promoting systemic Th1 immune responses (Mazmanian et al., 2005).

The fungi are also important components of the gut microbiome. The gut fungi mycobiota dysbiosis is related to many diseases (Azzam et al., 2020). In the present study, we found that the gut fungi mycobiota balance of mice was disrupted by CL, especially, *Aspergillus* and *Cladosporium* significant increase in the CL vs the CON groups. *Aspergillus* and *Cladosporium* are opportunistic pathogens that usually cause lung infections in immunocompromised patients (Wheeler et al., 2017; Sandoval-Denis et al., 2015). Bacteria and fungi harbor together in the gastrointestinal tract and occupy the same ecological niche. They interact with each other and develop complex ecological networks through these interactions. Homeostasis of microbial ecological networks limits pathogen invasion and infection (Lozupone et al., 2012). The ratio of ITS2/16S rRNA can reflect the diversity and composition structural alterations of the fungi–bacterial microbiota (Lemoinne et al., 2020). In general, antibiotics are directed against bacteria and do not affect fungi. However, our data revealed that CL treatment significantly altered the diversity and composition structure of the fungi–bacterial microbiota, and reduced the trans-kingdom network complexity and interaction between bacteria and fungi. These suggested that the alterations of gut bacteria can affect the composition and diversity of fungi. Previous documents had also reported that any changes in gut bacteria would inadvertently cause alterations in the gut fungal community, and targeted fungal interventions would also cause changes in the gut bacterial community (Haak et al., 2021; Sovran et al., 2018; Charlet et al., 2018; van Tilburg Bernardes et al., 2020). Hence, gut microbiota should be taken as a whole rather than a single genus to explain the relationship between gut microbiota and associated diseases.

Since most of the microbes in the gut are non-culturable, the FMT is considered to be an ideal model to study the function of gut microbiota. The success or not of FMT is influenced by many factors, such as host intestinal microbiota, immunity, and genetic factors (Porcari et al., 2023). During the FMT, not all microbiota in the donor feces have the same colonization ability in the receptors. Some research has revealed that the colonization success rate of *Bacteroidetes* is higher than that of *Firmicutes* (Hintze et al., 2014). In the present study, we found that the ratio of *Firmicutes*/*Bacteroidetes* in the CL treatment group increased compared to the control group (Figure 3b). However, after FMT, the ratio of *Firmicutes*/*Bacteroidetes* decreased in the CL-FMT group vs the CON-FMT group (Figure 4d). We speculated that the possible reason for this difference was that the colonization of *Firmicutes* decreased in the CL-FMT receptor group after transplantation. In contrast, the colonization of *Bacteroides* increased, resulting in a decrease in the proportion of *Firmicutes*/*Bacteroides* in the CL-FMT group. However, we considered the gut microbiota as a whole in the analysis of the experimental results. After FMT, we found that 85.11% of bacterial genera and 52.38% of fungi genera present in the CL inocula were successfully transferred to the CL-recipient mice, and 91.45% of bacteria genera and 56.36% of fungi genera in the CON inocula were also successfully transferred to the CON-recipient mice, respectively (Figure 4g). The trans-kingdom network analyses between bacteria and fungi showed that the trends of the gut microbiome in recipient mice were consistent with those in the donor mice. Therefore, the inconsistent results between Figures 3b and 4d will not affect this study's findings, and the FMT model established in this study is successful.

The FMT results revealed that gut commensal bacteria dysbiosis can promote MS colonization in the lungs of mice, but the underlying mechanisms remain not fully understood. Recent studies have suggested that changes in the 'gut–lung axis' are a contributing factor in the pathogenesis and clinical manifestations of Mtb infection (Yang et al., 2022; Alvarado-Peña et al., 2023.) What is the substance mediating the interaction between the lung and the gut? There have been a variety of results in the different studies. Yang et al. have found that the gut microbiota via modulation of lncRNA-mediated protective immunity against Mtb, and they found *Bacteroides fragilis* directly upregulated expression of lncRNA and promoted anti-TB immunity (Yang et al., 2022). However, another clinical study on TB found that the expression level of iNOS in the plasma of new-onset pulmonary TB patients was significantly higher than that of healthy humans (Chinta et al., 2016). In the present study, we found that *Nos2* expression significantly rises in the CL and CL-FMT groups. Hence, we speculated that the gut microbiota may increase MS colonization in mouse lungs by upregulation of *Nos2* gene expression and associated pathways. Lung epithelial cells, as first responders during Mtb infection, have been

shown to play an important role in TB pathogenesis (Scordo et al., 2016; Ryndak and Laal, 2019). Our results also confirmed that overexpression of *Nos2* in A549 cells (a human alveolar epithelial cell) can enhance MS colonization in these cells.

Nos2, also termed *iNOS*, is a homodimeric enzyme that is induced by immune stimulation in an independent of intracellular Ca^{2+} manner and plays an important role in infection, inflammation, immune regulation, and the control of intracellular bacterial pathogen infection (Chakravorty and Hensel, 2003). Compared with *Nos1* and *Nos3*, *Nos2* has the highest efficiency for production NO (Fischer et al., 2002). NO has direct antibacterial effects and immune-modulating function to intracellular pathogens and was considered a key molecule in controlling pathogens infections (Chakravorty and Hensel, 2003). It has been verified that reactive nitrogen intermediates produced by murine cells could inhibit Mtb infection (Scanga et al., 2001). The use of *Nos2* knock-out mice and NO synthase inhibitors is impaired in the control of Mtb growth (MacMicking et al., 1997; Chan et al., 1995). However, our data found that the high concentration of NO increased the colonization of MS in A549 cells. This result seems to contradict that NO has antibacterial activity. Some studies have also shown that NO can promote bacterial growth. Gusarov et al. reported that NO increases the resistance of bacteria to a broad spectrum of antibiotics, enabling the bacteria to survive and share habitats with antibiotic-producing microorganisms (Gusarov et al., 2009). Cole et al. showed that host cell-derived NO promoted the escape of listerolysin-dependent bacteria from phagocytic vacuoles into the cytoplasm by inhibiting the proton-pumping activity of V-ATPase and delaying phagolysosome fusion (Cole et al., 2012; Bogdan, 2012). Another study has shown that the maximal level of NO produced by human macrophages was not bactericidal or bacteriostatic to Mtb, and the number of viable mycobacteria was increased in macrophages that produced NO, and this is correlated with the expression of nitrate reductase (Jung et al., 2013). Therefore, we speculate that intracellular NO synthesized by *Nos2* has dual roles, including bactericidal at high concentrations and promoting bacterial growth at low concentrations.

ROS are commonly present in various habitats occupied by living organisms and the production of ROS appears as a very ancient host strategy for coping with pathogens (Rhen, 2019; Roy et al., 2017). ROS has shown potent antimicrobial activity against many pathogens including bacteria, fungi, and viruses. The accumulation of ROS is required for killing or inhibiting bacteria (Hong et al., 2019). NO interferes with antimicrobial killing by suppressing ROS accumulation (Gusarov et al., 2009). Hence, we assess the level of ROS in the overexpression of *Nos2* plasmids A549 cells. The results show that the ROS concentration in A549 cells transfected with *Nos2*-pcDNA3.1 was significantly lower than that in the control A549 and A549 cells transfected with pcDNA3.1 empty vector. This suggested that the promotion of bacterial proliferation by *Nos2* is dependent on lower ROS levels. The crosstalk between *Nos2* and ROS has been investigated in other research. Zheng et al. reported that the lower ROS level induced by *Nos2a* significantly inhibited *E. piscicida* proliferation and infection in zebrafish (Zheng et al., 2023). Another study has also shown that *Nos2*-produced ROS have an important role in maintaining homeostasis of the gut microbiota and defense against bacterial translocation (Matziouridou et al., 2018). Therefore, regulating the *Nos2*-ROS pathway may enhance the host's defense ability against pathogens.

β -defensin-1 is an antimicrobial peptide, which is mainly produced by various epithelial cells and is an important part of the innate immune response (Flores Saiffe Farias et al., 2015). *Defb1* is also expressed in the respiratory epithelium cells and protects the airways from respiratory pathogens (Diamond et al., 2008). Previous research has found that *Defb1* has 98% killing activity against active Mtb H37Rv (Álvarez et al., 2018), and also has an effective antibacterial effect against dormant mycobacteria (Sharma et al., 2017). Consistent with the above results, our data revealed that there was a strong negative correlation between *Defb1* gene expression and MS colonization in A549 cells. The present study was mainly performed at the gene level and lacked verification at the protein level. We will verify the role of the *Nos2*-*Defb1* pathway on MS colonization at the protein levels in future studies.

Altogether, the crosstalk between the gut microbiome and the lungs through the 'gut-lung axis' is complex and may involve multiple mechanisms, including immune response, metabolic disorders, cytokines, inflammation, etc. However, in the present study, we found that gut microbiome dysbiosis induced by clindamycin disturbs the gut equilibrium between bacteria and fungi, alters the lung transcriptome, and increases *Nos2* expression. Then, the microbiota dysbiosis could enhance the

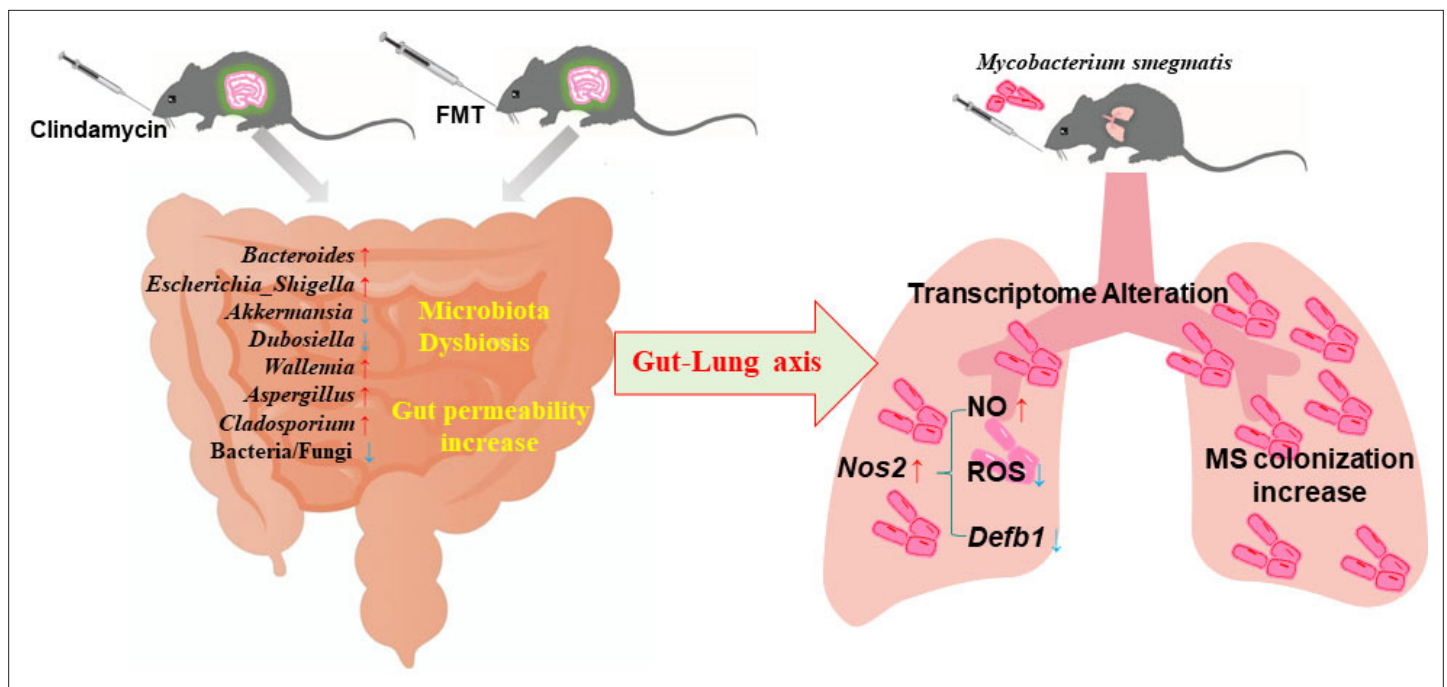


Figure 7. Mechanisms of the intestinal microbiome dysbiosis effect on the colonization of MS in the mouse lungs. Gut microbiome dysbiosis and gut permeability-increasing disrupt the lung transcriptome, and increase *Nos2* expression through the ‘gut–lung axis’. *Nos2* high expression weakens the intracellular antimicrobial and anti-inflammatory environment by increasing the concentration of NO, decreasing the levels of ROS and *Defb1* in the cells, and promoting MS colonization in the lungs of mice. MS: *Mycobacterium smegmatis*.

pulmonary colonization of MS in mice by regulating the *Nos2*–NO, *Nos2*–ROS, and *Nos2*–*Defb1* pathways (Figure 7). In the future, we need more experiments to explore the relationship between the intestinal microbiome and the *Nos2*-associated pathways in the lung and to further explore new targets for the prevention and treatment of TB.

Study limitations

This study has some limitations that need to be mentioned. First, due to the lack of experimental conditions in our laboratory that meet biosafety standards, we did not choose a wild-type Mtb strain to develop a mouse infection model, which may lead to the failure of the establishment of TB granuloma and other TB pathology icons and these also weaken the clinical significance of this study. Second, alveolar epithelial cells are one of the early contacting cells in Mtb infection, we only select alveolar epithelial cells (A549) to explore the colonization mechanism of intestinal microbiota affecting MS in vivo. This study lacks the choice of Mtb infection target cells–alveolar macrophages as the research object. Hence, future studies need to choose wild-type Mtb to establish animal infection models and alveolar macrophages for in vitro experiments to explore the regulation function of *Nos2* expression on NO, ROS, and *Defb1*, which will resolve these limitations of the study.

Conclusions

In conclusion, the present study reveals that (1) clindamycin treatment induces gut microbiome dysbiosis, disturbs the gut equilibrium between bacteria and fungi, reduces the interactions among bacterial–fungal trans-kingdom, and increases intestinal permeability. (2) The intestinal microbiome dysbiosis alters the lung transcriptome, increases *Nos2* expression, and enhances MS colonization in the lungs of mice. (3) Intestinal microbiota could promote the pulmonary colonization of MS in mice by regulating the expression of NO, ROS, and *Defb1* through *Nos2*-associated pathways, and changing the intracellular antimicrobial and anti-inflammatory environment. Hence, regulating the gut microbiome balance may be a potential strategy for reducing the risks of Mtb infections and transmission.

Materials and methods

Key resources table

Reagent type (species) or resource	Designation	Source or reference	Identifiers	Additional information
Strain, strain background (<i>Mycobacterium smegmatis</i>)	<i>Mycobacterium smegmatis</i> ; MS	ATCC	ATCC 19420 Cat#700084	
Cell line (<i>Homo sapiens</i>)	Human alveolar basal epithelial cell (A549)	Servieebio Biochemical Technology Co., Ltd	RRID:CVCL_0023	
Transfected construct (mouse)	pcDNA3.1-Nos2	Youbio Biochemical Technology Co., Ltd	NCBI GenBank accession No: NM_010927	
Commercial assay or kit	All-in-One Script RT premix	Kryptoner Mei Biotechnology Co., Ltd	#MR0502-2	
Commercial assay or kit	TB Green Premix Ex Taq II	TaKaRa Biotechnology Co., Ltd	#RR820A	
Commercial assay or kit	Nitric Oxide (NO) assay kit	Nanjing Jiancheng Biotechnology Research Institute Co., Ltd	#A012-1	
Commercial assay or kit	Reactive oxygen species (ROS) detection kit	Beyotime Biotechnology Co., Ltd	#S0033S	
Commercial assay or kit	The commercial ELISA kits of iFABP	Shanghai Enzyme-linked Biotechnology Co., Ltd	#TW9968	
Commercial assay or kit	The commercial ELISA kits of LPS	Shanghai Enzyme-linked Biotechnology Co., Ltd	#TW12543	
Chemical compound, drug	Clindamycin	Dalian Meilun Biotechnology Co., Ltd	CAS: 18323-44-9	1 mg/ml
Chemical compound, drug	Ampicillin	Shanghai Maclin Biochemical Technology Co., Ltd	CAS: 69-52-3	1 mg/ml
Chemical compound, drug	Metronidazole	Shanghai Maclin Biochemical Technology Co., Ltd	CAS: 443-48-1	1 mg/ml
Chemical compound, drug	Neomycin	Shanghai Maclin Biochemical Technology Co., Ltd	CAS: 1405-10-3	1 mg/ml
Chemical compound, drug	Vancomycin	Shanghai Maclin Biochemical Technology Co., Ltd	CAS: 1404-90-6	0.05 mg/ml
Other	Middlebrook 7H9/7H10	Qingdao Haibo Biotechnology Co., Ltd	#HB6270/HB9128	<i>Mycobacterium</i> culture media

Mouse husbandry and antibiotic treatment

Specific pathogen-free (SPF) C57BL/6 mice (6–8 weeks) were purchased from SiBeiFu (Beijing) Biotechnology Co., Ltd (No. SCXK (Beijing) 2019-0010). The mice were reared in an animal laboratory with a temperature of 22 ± 2°C and a relative humidity of 50 ± 5%, for 1 week before the model started. After adaptive feeding, mice were randomly divided into two groups, including a control group (CON) and a clindamycin group (CL), each group had 10 mice. During the experimental procedure, the CON group was fed with phosphate buffer saline (PBS). Whereas the CL group received 10 mg/kg clindamycin by oral gavage once a day for 14 days. Each oral gavage treatment did not exceed 200 µl, and treatment was stopped 2 days before the *mycobacteria* infection. All animal experiments were conducted according to the Declaration of Helsinki and were approved by the Animal Care and Use Ethics Committee of Xinxiang Medical University (No. EC-023-098).

Bacterial strains and infection

Due to the strong infectivity of Mtb, MS, a model strain for the study of TB, was used in the present experiments. MS was grown in Middlebrook 7H9 liquid medium supplemented with glycerin 0.5% and Tween 80 0.05%. After 18–24 hr of cultivation in a shaking incubator at 37°C, MS were centrifuged at 5000 rpm for 5 min, and the resulting pellet was suspended in sterile PBS to the concentration of 2 × 10⁷ CFU per ml. **Figure 1a** shows the experiment design procedure. For infection, mice were anesthetized by injection of 4% chloral hydrate (7 µl/g) and infected intranasally with 10⁷ CFU of MS. The mice were euthanized after infection 72 hr, and the left lung of each mouse was extracted and homogenized in PBS with 0.1% Tween 20. Tenfold serial dilutions were made in PBS with 0.1% Tween 20 and plated on Middlebrook 7H10 Agar plates, and colonies of MS were counted after 5 days of incubation at 37°C.

Fecal sample DNA extraction, 16S rRNA, and ITS sequencing

Fresh stool samples from each mouse were collected aseptically and the total genomic DNA from fecal samples was extracted using the Quick-DNA Kit for feces (QIAGEN, Germany) according to the manufacturer's instructions. The quality and concentration of DNA were determined using NanoDrop ND-2000 spectrophotometer (Thermo Scientific Inc, USA) and 1.0% agarose gel electrophoresis. Each DNA sample amplifies the hypervariable V3–V4 regions of the bacterial 16S rRNA genes using the primers 338F (5'-ACTCCTACGGAGGCAGCAGCA-3') and 806R (5'-GGACTACHVGGTWTAAAT-3'). The ITS2 regions of the fungi were amplified using primers ITS3F (5'-GCATCGATGAAGAACGCAGC-3') and ITS4R (5'-TCCTCCGCTTATTGATATGC-3'). The PCR products were purified with the AxyPrep DNA Gel Extraction Kit (Axygen Biosciences, Union City, CA, USA) and quantified using Quantus Fluorometer (Promega, USA). Purified amplicons were pooled in equimolar amounts, and paired-end sequenced on an Illumina MiSeq PE300 platform (Illumina, San Diego, USA) according to the standard protocols by Majorbio Bio-Pharm Technology Co. Ltd (Shanghai, China).

Serum iFABP and LPS measurement

The blood samples of all mice were collected via Eyeball, and the serum was separated by centrifugation at $1000 \times g$ for 20 min. Then, ELISA technology was used to measure the iFABP and LPS, according to the manufacturer's instructions. The commercial ELISA kits were obtained from Shanghai Enzyme-linked Biotechnology Co., Ltd (Shanghai, China) including iFABP (catalog number TW9968) and LPS (catalog number TW12543).

Fecal microbial transplantation

To establish the gut microbiota depletion mouse model, SPF mice were treated with a cocktail of antibiotics (1 mg/ml ampicillin, 1 mg/ml metronidazole, 1 mg/ml neomycin, and 0.5 mg/ml vancomycin) by oral gavage daily for 14 days. All mice were randomly separated into two groups, the CON-recipient group ($n = 10$) and the CL-recipient group ($n = 10$). Each group of mice was randomly housed in two cages with five mice in each cage. The fecal microbial transplantation procedure is shown in **Figure 4a**. Fecal samples from mice of the CON and CL groups were collected in sterile containers. Then 1 g fecal sample was suspended in 5 ml sterile PBS, followed by the vortexes, sedimentation, and filtrate with a 100- μ m cell strainer. The suspension of the same group was mixed as microbiota donors and immediately administered to the mice by oral gavage. 200 μ l of the supernatant containing fecal microbiota from either CON or CL donors was transferred to microbiota-depleted mice by oral gavage every day, for 14 days. All recipient mice were infected intranasally with 10^7 CFU MS after stopping FMT 2 days, and 3 days later of the infection, all mice were sacrificed, and the colonic contents and lung tissue samples were collected for further analysis.

Lung histological assessment

After the MS-infected mice were sacrificed, the lung tissues were perfused with sterile PBS and fixed in 4% paraformaldehyde for 3 days, followed by paraffin embedding. For histopathological analysis, 5 μ m sections were cut and stained using a standard H&E protocol. Motic EasyScan whole-slide scanner was used for scanning histological sections and images were analyzed using Matic DSAassistant Lite.

Lung tissue RNA extraction and RNA sequencing

The lung tissue from each mouse was separated aseptically and quickly stored in liquid nitrogen for subsequent RNA extraction. Total RNA was extracted from lung tissue using TRIzol reagent (Dingguo Changsheng Biotechnology Co., Ltd, Beijing, China) according to the manufacturer's protocol. RNA quantity and quality were determined using the NanoDrop 2000 Spectrophotometer (Thermo Scientific, USA). The Agilent 2100 Bioanalyzer (Agilent Technologies, Santa Clara, CA, USA) was used to assess RNA integrity. RNA-seq libraries were prepared using VAHTS Universal V6 RNA-seq Library Prep Kit following the manufacturer's recommendations. Then the paired-end RNA-seq libraries were sequenced with the Illumina HiSeq X Ten platform (2 $\times$ 150 bp read length) by OE Biotech Co., Ltd (Shanghai, China). After the quality control, clean reads performed bioinformatics analysis.

Quantification of gene expression using RT-qPCR

Total RNA was extracted from mouse lungs or A549 cells using the Trigol (Dingguo Changsheng Biotechnology Co., Ltd, Beijing, China) according to the manufacturer's protocol, and was reversed

transcribed using the kit from All-in-One Script RTpremix (Kryptoner Mei Co., Ltd, Zhengzhou, China). Then RT-qPCR was performed with TB Green Premix Ex Taq II (TaKaRa Biotechnology, China) to evaluate the amount of mRNA expression according to the manufacturer's recommendations. Subsequently, PCR products were detected on a sequence detection system. The primer sequences of RT-qPCR are listed in **Supplementary file 2** in this study. The relative gene expression levels were calculated using the $2^{-\Delta\Delta C_t}$ method.

Construction of *Nos2* overexpression plasmid and transfection into A549 cells

Human alveolar basal epithelial cell lines (A549) were purchased from Servieebio Biochemical Technology Co., Ltd (Wuhan), and cell lines have been authenticated with STR Short Tandem Repeat (STR) profiling. All the cell culture supernatants were found to be negative for mycoplasma contamination. To enhance *Nos2* expression levels in A549 cells, the recombinant plasmid *Nos2*-pcDNA3.1 was constructed. Based on the nucleotide sequences of *Nos2* (GenBank accession No: NM_010927), the forward and reverse primers *Nos2*F/*Nos2*R (**Supplementary file 2**) were used for cloning the open reading frame and inserted into pcDNA3.1 expression vector. The insert orientation was confirmed by separate XhoI and BamHI digests followed by agarose gel electrophoresis. The *Nos2*-pcDNA3.1 plasmid was mixed with lipo 8000 (Beyotime, Shanghai, China) and then transfected into A549 cells via electroporation. Twenty-four hours post-transfection, transfection efficiency was determined using RT-qPCR to ensure *Nos2* overexpression in A549 cells.

MS infects in A549 cells

A549, a human alveolar basal epithelial cell, was used to assess the infection capacity of MS. A549 cells were inoculated into 24-well plates with a density of 2×10^5 cells per well, and were cultured in RPMI1640 medium (Gibco Laboratories, USA) supplemented with 10% fetal bovine serum (Sangon Biotech Co., Ltd, Shanghai, China) and 100 U/ml penicillin/streptomycin (Beyotime, Shanghai, China) at 37°C. *Nos2*-pcDNA3.1 was transfected into A549 cells when the cell density reached 70–80%. After 24 hr, the supernatant was replaced with fresh medium containing MS and cultured for 6, 12, and 24 hr. Then the wells were washed three times with PBS to remove unattached bacteria. Subsequently, A549 cells were lysed with 1 ml 0.1% Triton X-100 (Shanghai Beyotime Biotechnology Co., Ltd, Shanghai, China) per well to prepare appropriate dilutions, which were plated on MiddleBrook7H10 Agar plates cultured for standard colony counts.

Detection of NO, ROS, and *Defb1*

Generation of NO in the A549 cells was detected by Nitric Oxide (NO) assay kit (Nanjing Jiancheng Biotechnology Research Institute Co., Ltd, Nanjing, China). Briefly, Cells were seeded into 24-well plates at a density of 2×10^5 cells per well, and *Nos2*-pcDNA3.1 was transfected into A549 cells when the cell density reached 70–80%. After 24 hr, the cells were harvested by trypsinization and centrifuged at 1000 rpm for 10 min. Then NO concentration was detected with the spectrophotometric method according to the manufacturer's protocols. ROS levels were measured using the DCFH-DA. Briefly, after 24 hr post-transfection of *Nos2*-pcDNA3.1, DCFH-DA was added to the culture medium at a final concentration of 10 μ mol/ml, and incubated for 1 hr at 37°C. The cells were washed twice with PBS and then stained with 10 μ mol/ml DCFH-DA at 25°C for 30 min in the dark room. Images were acquired by a confocal microscope (Nikon, Japan). For ROS quantification, A549 cells were collected, rinsed twice with PBS, and suspended in 10 μ mol/ml DCFH-DA for 30 min. After incubation, fluorescence was detected at 485 nm (excitation) and 530 nm (emission) using a microplate reader. All these analyses were conducted in three replicates. The expression of *Defb1* was detected using RT-qPCR, and the primer sequences of RT-qPCR are listed in **Supplementary file 2**.

Bioinformatics analysis

For microbiome analysis, the bioinformatics data were analyzed using the Majorbio Cloud platform (<https://cloud.majorbio.com>). The α -diversity at the genus level was assessed according to the Chao1 and inverse Simpson indices. The β -diversity was calculated by PCoA. A permutational analysis of variance was performed to assess the variation in the taxonomic structure of microbiota communities between groups. The Wilcoxon rank-sum test was used to assess the different structures of microbiota

communities between groups. LEfSe analyses were performed to compare different biomarkers between groups. The trans-kingdom network figures were built using the package igraph (version 1.2.6). For transcriptomic analysis, the bioinformatics data were analyzed using the OE Cloud platform (<https://www.oebiotech.com>). Differential expression analysis was performed using the DESeq2 q value <0.05 and foldchange >2 was set as the threshold for significantly DEGs. Venn diagram and volcano plot of DEGs were performed to explore gene expression patterns. The functional enrichment of the above DEGs was conducted using the GO database (<http://www.geneontology.org/>) and the KEGG (<http://www.genome.jp/kegg/>).

Statistical analysis

All statistics were performed using GraphPad Prism 8.0. If the data followed a normal distribution, unpaired Student’s t-tests were used to compare various parameters between the two groups. If the data did not follow a normal distribution, a non-parametric Mann–Whitney U test, and Wilcoxon rank-sum test were used to compare the results. One-way analysis of variance was used for three or more groups of data. The graphs were made with GraphPad Prism 8.0 or R package (version 3.6.2). p-values of <0.05 were set as a threshold for statistical significance.

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Additional information

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Author contributions

MeiQing Han, Investigation, Methodology, Writing – original draft; Xia Wang, Conceptualization, Resources; Lin Su, Shiqi Pan, Ningning Liu, Investigation, Methodology; Duan Li, Methodology; Liang Liu, Software; JunWei Cui, Software, Supervision; Huajie Zhao, Funding acquisition, Writing – review and editing; Fan Yang, Conceptualization, Funding acquisition, Writing – review and editing

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Ethics

All animal experiments were conducted according to the Declaration of Helsinki and were approved by the Animal Care and Use Ethics Committee of Xinxiang Medical University (No. EC-023-098).

Peer review material

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Author response <https://doi.org/10.7554/eLife.99282.3.sa2>

Additional files

Supplementary files

- Supplementary file 1. The relative abundance of gut bacteria and fungi at the phylum and genera levels. (a) The relative abundance of bacteria in CL and CON groups at phylum levels. (b) The relative abundance of bacteria in CL and CON groups at general level. (c) The relative abundance of fungi in CL and CON groups at the phylum level. (d) The relative abundance of fungi in CL and CON groups at the genera level. The Wilcoxon rank-sum test was performed between the two groups. CL: CL treatment group, CON: control group.
- Supplementary file 2. The Kyoto Encyclopedia of Genes and Genomes (KEGG) enrichment pathways analysis of the lung transcriptome data and the primers of RT-qPCR. (a) The primary enrichment pathways of differentially expressed genes (DEGs) in KEGG analysis at classification level 1. (b) The primary enrichment pathways of DEGs in KEGG analysis at classification level 2. (c) The primers of RT-qPCR.
- MDAR checklist

Data availability

16s RNA / ITS and transcriptome sequencing data have been deposited in NCBI under the accession codes PRJNA1091926 and PRJNA1099882. All data generated or analyzed during this study are included in the manuscript and supporting files. Source data files have been provided for the figures and figures supplements.

The following datasets were generated:

Author(s)	Year	Dataset title	Dataset URL	Database and Identifier
Han M, Wang X, Su L, Yang F	2024	SrRNA gene sequencing and ITS gene sequencing of mice	https://www.ncbi.nlm.nih.gov/bioproject/PRJNA1091926	NCBI BioProject, PRJNA1091926
Han M, Wang X, Su L, Yang F	2024	RNA-Seq of mice	https://www.ncbi.nlm.nih.gov/bioproject/PRJNA1099882	NCBI BioProject, PRJNA1099882

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