

7th International *C. difficile* Symposium Microbiome characterization of patients with *Clostridioides difficile* infection, recurrent *Clostridioides difficile* infection and colonized patients..

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Introduction and purpose

- The conditioning factor for CDI is the loss of microbiota diversity.
- Distinction between colonization and infection is sometimes difficult.
- Identifying risk factors or biological markers for R-CDI is important for proper treatment and prevention.
- The objective of our study was to characterize the microbiota profile of Healthy, CDI, R-CDI and colonized patients (CDC), in order to establish possible differences that could be used in CDI identification and risk stratification.

Methods

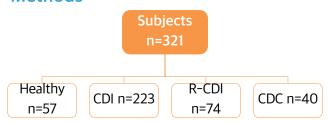


Figure 1. Number of healthy, *Clostridioides difficile* infection (CDI), recurrence of CDI (R-CDI) and colonized subjects (CDC).

- Clinical and fecal samples were collected after informed consent was obtained (Figure 1)
- ◆ The hypervariable V4 region of the 16s rRNA gene was sequenced on Illumina Miseq platform according to standard protocols.
- Data preprocessing, OTU clustering and taxonomic classification were done using MOTHUR software, RDP and SILVA database.
- Alpha diversity and statistical analysis were conducted by MOTHUR and R software.

Results

- Alpha-diversity, richness and evenness were significantly higher in the healthy group versus CDI and CDC (p<0.05;all) (Figure 2).
- Patients with previous CDI had lower diversity and richness(p<0.01;all).
- → Those patients with R-CDI had lower diversity and richness than CDI (p<0.01) (Figure 3).</p>



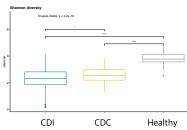




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Results

Figure 2. Alpha-diversity of healthy, *Clostridioides difficile* infection (CDI), and colonized subjects (CDC).



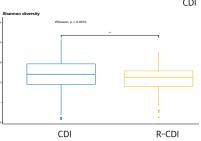


Figure 3. Alpha-diversity of Clostridioides difficile infection (CDI) and recurrence of CDI (R-CDI).

- * R-CDI had higher abundance of *Clostridiaceae*, *Fusobacterium* and *Akkermansia* (p<0.001;all), and lower levels of *Bacteroides*, *Collisnella*, *Prevotella*, *Alistipes*, *Parabacteroides*, *Blautia*, *Barnesiella*, *Butyricimonas*, and *Oscillibacter* (p<0.001;all) (Figure 4).
- Colonized patients had higher abundance of Blautia, and lower abundance of Bacteroides, Clostridiaceae, Clostridium_XIVa, Lactobacillus, Romboutsia, Prevotella, Staphylococcus, Streptococcus and Veillonella (p<0.001;all) (Figure 5)

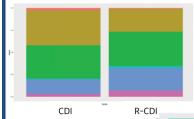


Figure 4. Microbiota profile of *Clostridioides difficile* infection (CDI) and recurrence of CDI (R-CDI).

Figure 5. Microbiota profile of healthy, *Clostridioides difficile* infection (CDI), and colonized subjects (CDC).

CDI CDC Healthy

Conclusions

- We found significant differences in the microbiota profile between CDI and CDC patients.
- Prior CDI episodes and R-CDI episodes had a cumulative impact in the microbiota.
- We identified several groups of microorganisms that may serve as microbiological markers of true CDI.