Abstract IDDF2021-ABS-0123 Figure 1

cancer properties, including ectoine, desferrioxamine B/E, and antimycin. Subsequently, one of the predicted genes from the LAPs gene cluster was selected and de novo synthesized for expression study using *Escherichia coli* model. In the anti-MRSA test, the purified protein Protein135 showed a minimum inhibitory concentration (MIC) of 3 μ M against MRSA ATCC 33591 at 24-hour incubation. Intriguingly, Protein135 demonstrated higher growth inhibition than vancomycin, both at sub-MIC of 0.25 μ M, against MRSA ATCC 33591.

Conclusions Altogether, these findings highlight the importance of novel strain from the mangrove forest, particularly in searching for beneficial bioactive compounds. In addition, our preliminary study unveils the promising wound healing property of Protein135 in the MRSA infection murine model.

IDDF2021-ABS-0126

EXPLORING THE GUT MICROBIOTA VARIATION IN RESPONSE TO VIBRIO INFECTION

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10.1136/gutjnl-2021-IDDF.42

Background Our gut microbiota is composed of trillions of bacteria residing in the human gastrointestinal system. Any

Abstract IDDF2021-ABS-0126 Figure 1

disruptions to this bacterial community are often associated with various diseases' progression, including gastrointestinal infection. One significant diarrheal pathogen is *Vibrio* species, the critical etiological agent to cholera and gastrointestinal infection that affects millions of people annually. They invade the host's gastrointestinal system, releasing virulence toxins that aid in the colonization, dysbiosis of the gut flora, and cause diarrhea. The gut microbiota community is diverse, yet how this microbial diversity confers resistance or susceptibility to intestinal pathogens is not well explored. Hence, this systematic review aims to understand the gut microbiota variation in response to *Vibrio* infection.

Methods This systematic review was conducted based on the PRISMA guidelines. Predefined MeSH terms 'Vibrio' AND 'gut microbiota' OR 'microbiome' were used in three databases (Pubmed, Ovid Medline, ProQuest; from database inception to June 2021). Studies reporting on the gut microbiota variation and *Vibrio* were included. Studies without *Vibrio* and/or gut microbiome data were excluded, including conference proceedings, reviews, systematic reviews, meta-analyses, and commentaries.

Results Six studies according to the study's inclusion criteria out of 243 articles were selected for qualitative analysis. Three studies identified the gut microbiota composition in patients' samples with *Vibrio* species infection, while the remaining studies explored the modulation of gut microbiota in *Vibrio*-induced animal models. The human studies demonstrated variation in the gut microbiome of patients with *Vibrio* infection as compared to the healthy subjects. *Bacteroides*, *Lactobacillus*, *Bifidobacterium*, and *Akkermansia* are among the prominent taxa found to be significantly decreased in abundance but increase in the taxa of Firmicutes among the patients and animals, elevating the infection and causing diarrhea. An in vivo study also reported the abundance of *Ruminococcus/Blautia obeum*, that able to inhibit the *Vibrio cholerae* virulence in the gut (IDDF2021-ABS-0126 Figure 1. Illustration of gut microbiota variation in response to *Vibrio* infection).

Conclusions There is a potential connection between gut dysbiosis and *Vibrio* species infection, in which the gut microbiota possibly increase/decrease its abundance to inhibit the infection. These findings provide insights on gut microbiota response towards environmental pathogen and frame proper treatment strategies in clinical settings.

IDDF2021-ABS-0130 DYSBIOSIS OF GUT ARCHAEA IN OBESITY RECOVERED AFTER BARIATRIC SURGERY

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10.1136/gutjnl-2021-IDDF.43

Background Obesity is associated with altered gut microbiota. Archaeome is an important part in intestinal microbiota, however, the role of archaea in obesity remains unknown. We aimed to delineate the alterations of gut archaea in subjects with obesity and explore the effect of bariatric surgery-associated weight loss intervention in modulating the gut archaeal composition.

Methods A total of 231 obese subjects and 184 lean controls with shotgun metagenomic sequencing available were retracted from three public cohorts. Of which, 23 obese subjects undergone bariatric surgery were followed up for 3 months. Alterations of gut archaeome, bacteriome and archaeal-bacterial correlations were assessed in obese and lean subjects.

Results Fecal archaea microbiota of obese subjects showed a significantly lower Chao1 richness and Shannon diversity compared to lean controls ($p < 0.05$). Principal component analysis of archaeome revealed distinct clusters in obese subjects and lean controls ($p < 0.05$). Two archaeal orders including Methanobacteriales and Methanococcales were increased and three orders including Natrilbales, Methanocellales and Thermoproteales were decreased in obese subjects compared with lean controls. Top 20 differentially abundant archaeal genera discriminated obese subjects from lean controls with an area under the receiver operating characteristic curve (AUC) of 0.79 (95% CI 0.72-0.85) in the discovery cohort. It was further validated in two independent cohorts, with AUC of 0.83 (0.73-0.92) and 0.86 (0.77-0.94), respectively. Further prediction analysis by archaeal genera revealed a significantly lower probability of obesity in subjects after bariatric surgery than their baseline ($p < 0.05$). Moreover, the ecological analysis showed significantly lower trans-kingdom correlations between archaea and bacteria in obese subjects compared with lean controls, and these correlations were partly recovered in subjects receiving bariatric surgery.

Conclusions This study demonstrated for the first time that obesity is characterized by gut archaeome dysbiosis across multiple cohorts with different demographic characteristics. Weight loss by bariatric surgery-induced a significant change of gut archaeome, suggesting that gut archaeome may play an important role in treating obesity. Studies are needed to explore the potential benefit of reshaping gut archaeome to improve weight loss intervention efficacy.

IDDF2021-ABS-0132 MOVING BEYOND THE GASTROINTESTINAL TRACT: THE INVOLVEMENT OF GUT MICROBIOME IN ENDOMETRIOSIS

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10.1136/gutjnl-2021-IDDF.44

Background The importance of gut health and microbiome have been emphasized in women's health. Endometriosis is characterized by abnormal growth of endometrial cells that occurs outside of the uterus (IDDF2021-ABS-0132 Figure 1. Potential association of gut microbiome in endometriosis). Animal models of endometriosis display different gut microbiota compositions compared to healthy controls, and the oral administration of broad-spectrum antibiotics has been shown to reduce disease progression. The current study investigates the relationship between the gut microbiome and endometriosis in humans before exploring potential mechanisms involved.

Methods Searches were performed in three databases (PubMed, Ovid Medline, ScienceDirect; database inception to 31st May 2021 using 'gut', 'microbiome,' or 'microbiota' combined with 'endometriosis' following PRISMA guidelines). All