Results HOXC6 was overexpressed in RCC and associated with poor prognosis. Overexpression of HOXC6 promoted the migration and invasion of CRC cells in vitro and in vivo. Increase in CCL2 expression by upregulation of HOXC6 could attract more infiltrating M2 macrophages. IL6 secreted by M2 macrophages could induce EMT in tumor cells by upregulating HOXC6 and activating the Wnt/β-catenin signaling pathway via inhibition of DKK1 secretion.

Conclusions Our study indicates that overexpression of HOXC6 induces EMT by regulating the DKK1/Wnt/β-catenin axis. The positive crosstalk between M2 macrophages and HOXC6 in tumors led to poor prognosis of RCC.

**IDDF2020-ABS-0090**
MALONDIALDEHYDE LEVELS IN THE TESTICULAR ORGAN OF HYPERLIPIDEMIC RAT (RATTUS NORVEGICUS) WITH QUAIL EGG YOLK DIET

1Muhammad Luthfi Adnan*, 1Dini Islamiana, 1Hilmi Ardian Sudarto, 1Miranti Dewi Pramantingtyas, 1Undergraduate Program of Medicine, Faculty of Medicine, Universitas Islam Indonesia, Indonesia; 2Deparment of Physiology, Faculty of Medicine, Universitas Islam Indonesia, Indonesia

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Background Hyperlipidemia can cause infertility due to cell damage to the testicular organs. Hyperlipidemia can induce lipid peroxidation, which results in the formation of malondialdehyde (MDA). Quail egg yolk is one of the many foods consumed by Asians that contain high lipid levels. The aim of this study was to determine the effect of quail egg yolk diet on MDA levels of testicular organ on the rat.

Methods The subjects are male Wistar (Rattus norvegicus) strain rats 2–3 months with body weight 200–300 grams divided into two groups (K+ and K−). Group of K+ were given quail egg yolk for two weeks with a dose of 5 ml while group of K− were only given fed ad libitum. All rats terminated to taken the testicular organ to measure the level of MDA. All data were statistically analyzed with one-way ANOVA. Values were considered significant at p<0.05.

Results Mean of MDA level (nmol/gram) in rats was 0.95 ± 0.75 in K− group and 8.64 ± 0.13 in K+ group. The one-way ANOVA test showed significant differences in activity between the group with p<0.001 and Post Hoc test p<0.001.

Conclusions Quail egg yolk significantly increases MDA Levels in the testicular organ of hyperlipidemic rats. Consumption of quail egg yolks can affect the cell activity of testicular organs.

**IDDF2020-ABS-0112**
GUT-SKIN AXIS: DECODING THE LINK BETWEEN THE GUT MICROBIOME AND HIVES

Learn-Han Lee*, Vengadesh Lethumanan, Loh Teng-Hern Tan, Hooi-Leng Ser, Jodi Woon-Fei Lian. Novel Bacteria and Drug Discovery Research Group (NBDD), Microbiome and Bioresource Research Strength (MBRS), Jeffrey Cheah School of Medicine and Health Sciences, Monash University Malaysia, Malaysia

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Background Urticaria (hives) is a rash affecting the skin and mucosa, often characterized by appearance wheals, angioedema, and pruritus. Chronic urticaria (CU) is classified by the occurrence of urticaria which exceeded six weeks (almost daily) without specific triggers and identifiable cause. CU is a common disease that has detrimental effects on quality of life. However, its aetiology remains unclear. There is increasing evidence that dysbiosis of the intestinal microbiota is associated with dermatologic conditions. The human gut microbiome has a significant role in the regulation of the immune system, which can be implicated in the development of immune-mediated diseases such as CU. This systematic review aims to investigate the relationship of gut bacteria and the development of CU.

Methods The systematic literature search was executed in accordance with the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) guideline. Research commenced using MeSH terms relevant to the topic, (‘gut’ OR ‘microbiome’ OR ‘microbiota’ OR ‘microflora’) AND (‘urticaria’ OR ‘hives’) were performed on four databases (PubMed, EMBASE, ProQuest, Scopus; publication date limit to 29 February 2020). Titles and abstracts of all retrieved intergenic non-protein coding RNA 01446 (LINC01446), a 3484 bp ncRNA, is known to locate at chromosome 7p12.1. However, its biological functions and specific action mechanism in gastric cancer (GC) are still unclear.

Methods LINC01446 expression levels in GC cells were detected by quantitative real-time PCR (qRT-PCR). LINC01446 siRNAs and overexpression vector were transfected into GC cells to down-regulate or up-regulate LINC00460 expression. The biological functions of LINC01446 were investigated in vitro and in vivo. Transcriptome RNA-seq, Western blot, RIP, RNA pull-down and ChIP assays were used to determine the mechanism of LINC01446 in regulating underlying targets.

Results In our study, LINC01446 was proved to be markedly up-regulated in GC tissues relative to and positively correlated with the poor survival of GC patients. The multivariate Cox regression model showed that LINC01446 functioned as an independent prognostic factor for the survival of GC patients. Functionally, LINC01446 facilitated the proliferation and metastasis of GC cells. Moreover, RNA-seq analysis demonstrated that LINC01446 knockdown primarily regulated the genes relating to the growth and migration of GC. Mechanistically, LINC01446 could widely interact with histone lysine-specific demethylase LSD1 and recruit LSD1 to the Ras-related dexamethasone-induced 1 (RASD1) promoter, thereby suppressing RASD1 transcription.

Conclusions Overall, these findings suggest that LINC01446/LSD1/RASD1 regulatory axis may provide bona fide targets for anti-GC therapies.

**IDDF2020-ABS-0100**
LONG INTERGENIC NON-PROTEIN CODING RNA 01446 FACILITATES THE PROLIFERATION AND METASTASIS OF GASTRIC CANCER CELLS THROUGH INTERACTING WITH THE HISTONE LYSINE-SPECIFIC DEMETHYLASE LSD1

1Webin Lian*, 2Hongchi Xu, 3Jianlin Ren, 2Changheng Yan, 2Yifan Lian. 1Quanzhou First Hospital Affiliated to Fujian Medical University, China; 2Zhongshan Hospital, Xiamen University, China

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Background Growing evidence illustrated that long non-coding RNAs (lncRNAs) exhibited widespread effects on the progression of human cancers via various mechanisms. Long
studies were screened and refined by the inclusion and exclusion criteria. Studies involving gut microbiome and CU were considered for inclusion. Irrelevant articles based on title/abstract level, case reports, conference abstracts and other studies with no experimental intervention (reviews, book, commentaries) were excluded.

Results Three studies were eligible for final qualitative analysis, with a total of 100 participants. Research findings have shown that CU patients have a significant decrease in abundance of Firmicutes (Lactobacillus; Faecalibacterium prausnitzii), Actinobacteria (Bifidobacterium), Bacteroidetes (Bacteroides fragilis, Bacteroides plebeius), whilst an increase in abundance of Proteobacteria. The research suggested that increased abundance of Proteobacteria might enhance the permeability of intestinal mucus inner layer and enable bacterial infiltration, causing inflammation of epithelium and impairment of gut barrier function which leads to the development of inflammatory skin diseases.

Conclusions As a summary, this outcome provides a preliminary understanding of microbial composition in CU patients (figure 1). This offers a new avenue of research for potential CU treatment via maintaining gut health.