Research progress of intestinal microecology in the occurrence and development of precancerous lesions of liver

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Abstract. Intestinal microecology refers to the interaction between the host and the microorganisms in the human intestinal tract, which is composed mainly of intestinal flora. Intestinal microflora affects the physiological and pathological changes of the host through metabolic activity and host interaction. Precancerous lesion of liver is a potential benign liver disease, which may lead to malignant transformation of liver. It is the intermediate stage from benign lesion to malignant transformation. Recent studies have shown that intestinal microecology is closely related to the occurrence of precancerous lesions of the liver. This study expounds the interaction of the bridge between intestine and liver, the gut-liver axis, the intestinal microecology and the precancerous lesions of liver, hoping to provide a new idea for clinical prevention and treatment of precancerous lesions of liver.

1 Introduction

Intestinal microecology is a complex and diverse ecosystem, which has a specific symbiotic relationship with the host. At present, intestinal microecology plays an important role in the physiological and pathological process of the host, which is closely related to the pathogenesis of liver diseases[1]. The gut microecology includes the gut microbiome, which normally maintains a dynamic equilibrium with the host, but is disrupted by changes in the host and environment, leading to disease[2].

Hepatocellular carcinoma (HCC) is the most common primary liver cancer and the sixth most common malignant tumor in the world, with high malignancy and the third highest mortality rate in the world[3]. Hepatitis—liver fibrosis—liver cirrhosis—liver precancerous lesions—liver cancer is the five stages of liver disease development. In the liver cirrhosis, the liver cancer malignant transformation process, will experience a quite long time, namely the liver cancer precancerous lesion stage[4]. Because the early symptoms of liver cancer are not obvious and easy to be ignored, and the cure rate of advanced liver cancer is low, the cure difficulty is big and the treatment plan is complex, so it is more meaningful to prevent the occurrence and development of precancerous lesions than to treat liver cancer. Precancerous lesion of liver is a potential benign liver lesion with the possibility of malignant transformation of liver. It is the intermediate stage from benign lesion to malignant transformation. Precancerous lesions of liver mainly include liver cell dysplasia, hepatic adenomatous hyperplasia, liver fibrosis, liver cirrhosis and other diseases. It is influenced by hepatitis, alcoholic liver disease, nonalcoholic fatty liver disease, cirrhosis and many other factors. In recent years, with the further study of the intestinal liver axis, it has been found that the intestinal microecological imbalance and the progression of chronic liver disease can affect each other[5]. This article discusses the relationship between Intestinal microecology and the occurrence and development of precancerous lesions of liver, and provides more directions for the prevention and treatment of precancerous lesions of liver.

2 Gut microecology

The intestinal microecology is a kind of ecosystem which is composed of the interaction between the microorganism and the host. Its main functions include participating in metabolic activities, nutritional effects, immunity and protection of receptors from foreign microorganisms[6]. Dysbacteriosis can lead to inflammation and fibrosis of the liver[6,7]. Probing into the composition of intestinal microflora is helpful to understand the relationship between intestinal microecology and host diseases.

3 Composition of intestinal flora

The gut microbiome consists of bacteria, archaea, and eukaryotes that congregate in the gastrointestinal tract and have evolved with their hosts over thousands of years to form complex and mutually beneficial interactions[8,9]. According to statistics, there are more than 1014 kinds of microorganisms in the digestive tract, among which there are 10 times more bacterial cells and
Although the gut microbiome was previously thought to consist of between 500 and 1,000 species of microbes, a recent large-scale study estimated that the gut microbiome consists of more than 3,500 species of bacteria. Intestinal flora can be divided into three categories: beneficial bacteria, pathogenic bacteria and conditional pathogenic bacteria. Beneficial bacteria such as Bifidobacterium, lactobacillus, etc.; Pathogenic bacteria such as Staphylococcus, proteus, clostridium, pseudomonas Aeruginosa and opportunistic pathogens such as Enterococcus, Escherichia coli, bacteroides. The intestinal microflora can be divided into five groups: bacteroidetes, fircmutes, actinomycetes, proteobacteria and verruculose microphytes. The number of bacteroidetes and fircmutes accounts for more than 90% of healthy people's intestinal tract, the quantity of bacteroidetes and firmicutes was less.

4 Metabolites of Intestinal Flora

The metabolic products of the intestinal flora are short-chain fatty acid and secondary bile, in which short-chain fatty acid is the major. Short-chain fatty acid is produced by intestinal bacteria that ferment polysaccharides that the body can not digest. Short-chain fatty acid regulates various aspects of the host by inhibiting histone deacetylase and activating specific G protein coupled receptors, such as intestinal motility, inflammation, glucose homeostasis, and energy. Short-chain fatty acid consists mainly of acetate, propionate and Butyrate, which are important energy sources for intestinal epithelial cells to maintain the intestinal barrier, and propionate, which converts and inhibits cholesterol synthesis in the liver. Intestinal dysbacteriosis is caused by overgrowth of bacteria and abnormal accumulation of metabolic products, which leads to the destruction of intestinal barrier and liver damage caused by large numbers of bacteria entering the liver. Studies have shown that exogenous acetate and propionate can prevent hepatic steatosis, inhibit hepatic fat synthesis, reduce liver damage caused by hepatic steatosis and prevent precancerous liver lesions. Short-chain fatty acid can also increase the risk of hepatocellular carcinoma by inducing cholestasis.

5 Intestinal hepatic axis

The liver is an organ with a dual blood supply. It supplies oxygenated blood to the liver from the celiac artery and receives blood from the intestine to the hepatic portal vein. About 75% of the liver's blood comes from the hepatic vein. Therefore, there is a close relationship between the intestine and the liver, forming the intestine-liver axis. The enteric-hepatic axis refers to the mutual regulation of the intestinal mucosa and the liver through the portal circulation. The relationship between intestinal flora and liver is regulated and stabilized by the interaction of metabolism, immunity and neuroendocrine. Intestinal barrier is a barrier formed by connecting adjacent cells through tight junction protein, which is a natural barrier against bacteria and their metabolites. In the absence of the host's gut microbiome, the antigens (derived from pathogenic microorganisms) pass through these tight junctions, which are recognized by the dendritic cell and activate the adaptive immune system, resulting in increased intestinal permeability. Small doses of pathogen-related molecules such as Lipopolysaccharide (Lps), peptidoglycan and flagellin enter the liver via Toll-like receptor 4(Tlr4) and cause liver fibrosis.

6 Intestinal microecology and precancerous lesions of liver

Changes of intestinal microflora in patients with precancerous lesions of liver. When the liver becomes diseased, the dynamic equilibrium between the host and the gut ecology is broken down, leading to a disruption of the gut flora. Liver fibrosis and cirrhosis may lead to bacterial translocation through different mechanisms, including overgrowth of intestinal bacteria, derangement of intestinal lumen factors, increase of intestinal permeability and decrease of immune function. According to research, liver fibrosis and cirrhosis are closely related to intestinal microecological imbalance. In cirrhosis, a decrease in bile acid secretion and portal hypertension overgrowth of the intestinal flora. The study showed that the intestinal flora of patients with cirrhosis showed higher levels of pathogenic bacteria such as enterobacteriaceae and streptococci, as well as lower levels of beneficial bacteria such as bifidobacterium and lactobacillus, compared with healthy controls. Quantitative metagenomic analysis of intestinal microflora in patients with liver.

Intestinal microecological imbalance promotes the occurrence and development of precancerous lesions of liver

Intestinal microecological imbalance refers to increased permeability of intestinal mucosa, a large number of bacterial translocation. Bacterial translocation refers to the migration of live bacteria or bacterial endotoxin (lipopolysaccharide, peptidoglycan, lipopeptide) from the intestinal cavity to the mesenteric lymph nodes and other extraintestinal sites. When bacteria overgrow, intestinal permeability increases and a lot of bacterial translocation, especially of lipopolysaccharide (gram-negative), reaches the liver through the hepatic portal vein, it binds to toll-like receptor 4 in various liver cells, such as kupffer cells (KCs) and hepatic stellate cells (HSCs), causing chronic hepatitis, cirrhosis and liver cancer. Therefore, the activation of LPS-TLR4 pathway is very important in the development of liver precancerous lesions. When activated by LPS-TLR4, KCs in liver cells secrete TNF-α, IL-6 and other inflammatory mediators, which cause oxidative stress and inflammatory reaction in liver, damage cell DNA and induce gene mutation. At the same time, when HSCs are activated by the LPS-TLR4 pathway.
pathway, they can synthesize a large number of extracellular matrix proteins, which can promote liver fibrosis, and further promote the development of liver cirrhosis. After activation, HSCs also secrete angiogenesis factors, IPS induced the migration of mesenchymal endothelial cells and the formation of blood vessels, which is a key step in the development of hepatocellular carcinoma. Therefore, the composition and function changes of intestinal microecology may be closely related to the development of liver fibrosis and cirrhosis, thus promoting the development of precancerous liver lesions.

7 Conclusion

Intestinal microecological imbalance is closely related to the occurrence and development of precancerous lesions of liver. Precancerous lesions of the liver in patients with intestinal pathogenic bacteria significantly increased and beneficial bacteria significantly reduced, at the same time, the imbalance of intestinal microecology leads to the overgrowth of intestinal bacteria, the increase of intestinal permeability and bacterial translocation, which induce a lot of lipopolysaccharide to enter the liver through the portal vein. Therefore, the composition of intestinal flora, metabolic products, prevent bacterial translocation, block the signal pathway may delay or prevent the development of precancerous liver lesions. At present, there are few studies on the relationship between intestinal microecology and precancerous lesions of liver. Therefore, it is necessary to explore the relationship between intestinal microecology and liver precancerous lesion to provide a new therapeutic idea for liver precancerous lesion.

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