

IFN- γ 基因多态性与HBV感染及原发性肝细胞癌易感性的研究

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Study on Susceptibility of HBV Infection and Primary Hepatocellular Carcinoma with Gene Polymorphism of IFN- γ

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摘要 目的

探讨细胞因子IFN- γ 基因-1615C/T和+5171A/G位点单核苷酸多态性在广西人群中的分布及其对原发性肝细胞癌(HCC)发生、乙型肝炎病毒(HBV)感染的影响。方法设计以医院为基础的病例对照研究,对375名HCC患者、377名HBV携带者和406健康对照进行频数匹配,采用TaqMan MGB实时荧光定量PCR技术对上述位点进行分型。应用Logistic回归模型分析基因型在三组中的分布差异及基因环境交互作用,并进行连锁不平衡和单倍型分析。结果-1615C/T和+5171A/G位点的基因多态性在三组中分布差异无统计学意义($P>0.05$)。Logistic回归分析结果显示,吸烟、饮酒和肝癌相关家族史与基因存在交互作用;饮酒联合-1615C/T位点突变型基因T能增加HBV感染风险($OR=1.72$, $95\%CI:1.11\sim3.26$);两个位点的突变型基因T和G联合肝癌相关家族史能增加HCC患病风险($OR:29.24, 52.03, 95\%CI:6.91\sim123.6, 7.02\sim385.4$)。IFN- γ 的-1615C/T和+5171A/G位点存在连锁不平衡($D' = 0.976, P=2.22\cdot 10^{-16}$),但单倍型分布在HCC组与总对照组(HBV携带者对照和健康对照)间无统计学差异。结论IFN- γ 的-1615C/T和+5171A/G位点的突变型基因可能不是广西人患HCC和感染HBV的直接危险因素,但环境危险因素对HCC发生和HBV感染有协同作用。

关键词: 原发性肝细胞癌; HBV感染; IFN- γ ; 基因多态性

Abstract: Objective

To explore the distribution of cytokines IFN- γ gene (-1615C/T and +5171A/G) single nucleotide polymorphisms in Guangxi people, and the impact of hepatitis B virus (HBV) infection and primary hepatocellular carcinoma (HCC) occurrence. Methods A case-control study based on hospital was carried out and all the objects were frequency matched by 375 HCC patients - 377 HBV carriers - 406 healthy control. TaqMan MGB Real-Time fluorescence quantitative PCR technology was applied to detect the SNPs of the two loci. The distribution of the genotype and the interaction of gene-environment in the three groups were analyzed by Logistic regression model. The linkage disequilibrium and haplotype of IFN- γ gene were analyzed. Results There was no significant statistically difference in the polymorphisms of -1615C/T and +5171A/G loci among the three groups ($P>0.05$). There were gene-environment interactions in smoking, alcohol consumption, liver cancer related family history with IFN- γ gene according to logistic regression analysis. Alcohol consumption combined -1615 locus mutant gene G increased HBV infection risk ($OR=1.72, 95\%CI:1.11\sim3.26$). The two loci mutant genes combined with liver cancer related family history also enhanced HCC risk ($OR:29.24, 52.03, 95\%CI:6.91\sim123.6, 7.02\sim385.4, respectively$). -1615C/T and +5171A/G sites on IFN- γ had linkage disequilibrium ($D' = 0.976, p=2.22\cdot 10^{-16}$), but the haplotypes between HCC groups and the total controls (HBV carriers and healthy control) had no significant statistically difference. Conclusion The mutant genes of -1615C/T and +5171A/G loci might not influence the occurrence of HCC and HBV infection directly in the population of Guangxi, however they enhanced the risk interacted with the environment risk factors.

Key words:

服务

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





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