论著

恶性疟原虫FCC1/HN株新抗原表达序列标记位(ESTs)的获得

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目的: 以筛选恶性疟原虫FCC1/HN 株 λ g t 1 1 c DNA 表达文库所获得的强阳性克隆作基础,对上述强阳性克隆的 c DNA 插入片段进行DNA 序列测定,阐明相对应的新表达序列标签(ESTs),作为发现新抗原基因的线索。方法:以 c DNA 表达文库接头的较长链作PCR引物、扩增 c DNA 插入片段,将扩增产物克隆入M13 m p 18测序载体,进行部分DNA 序列测定、编辑,将之在GenBank中进行DNA 序列同源性搜索比较和分析。结果: 获得1个C03序列为已知恶性疟原虫热休克蛋白70-2基因片段,发现5个新的具有抗原意义的恶性疟原虫表达序列标记位(ESTs)。结论: 这5个新的恶性疟原虫表达序列标记位为发现新的恶性疟原虫抗原基因奠定了基础。

关键词 <u>恶性疟原虫</u> <u>抗原</u> <u>cDNA</u> <u>表达序列标记位</u> 分类号

FINDING OF NEW FCC1/HN ANTIGENIC EXPRESSED SEQUENCE TAG(ESTs) OF PLASMODIUM FALCIPARUM

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Abstract

AIM: To sequence the strong positive clones obtained by immuno-screening of Plasmodium falciparum FCC1/HN λ gt11 cDNA expression library, and to elucidate the antigenic expressed sequence tags through sequencing the cDNA insert of these positive clones, and new antigenic ESTs could serve as a resource to pursue their corresponding antigen genes. METHODS: cDNA inserts of positive λ gt11 phage clones were amplified by PCR. The PCR products, after purification, were cloned into the M13 mp18 sequencing vector. Single-stranded M13 DNA was prepared and sequenced. Then the acquired sequences were compared in homologies with EMBL/GenBank database on the PC/GENE software system and searched in NCBI (National Center for Biotechnology Information) GenBank using BLAST (Basic Local Alignment Search Tool) commond. RESULTS: Sequence CO3 was part of the known P.falciparum antigenic heat shock protein 70 (Pfhsp70) gene, while the other 5 sequences were new P.falciparum antigenic expressed sequence tags (ESTs). CONCLUSION: The 5 new antigenic ESTs generated could serve as the breaking through points in our efforts to find out new P.falciparum antigen genes.

Key words Plasmodium falciparum antigen cDNA expressed sequence tags

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