论著

以重组蛋白为抗原的ELISA法检测旋毛虫抗体

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摘要

目的 寻求特异性强、敏感性高的旋毛虫病诊断抗原。 方法 以旋毛虫新生幼虫期特异性T668基因在 E.coli高效表达的重组蛋白为抗原,分别以兔、猪和健康者血清及旋毛虫病阳性血清为一抗,以辣根过氧 化物酶(HRP)标记的山羊抗兔IgG、山羊抗猪IgG和山羊抗人IgG为二抗,建立检测旋毛虫抗体的间接 ELISA方法,并以旋毛虫肌幼虫排泄?鄄分泌(ES)抗原作为检测对照。 结果 以T668重组蛋白为抗 原,对兔、猪和人旋毛虫病血清进行检测,阳性检出率为100%,且敏感性高(0.016 μg/孔),与 ES抗原检测结果完全一致。 结论 T668重组抗原有望替代ES抗原检测旋毛虫抗体。

旋毛虫 重组蛋白 新生幼虫期特异性基因 间接ELISA 关键词 分类号

Detection of Anti-Trichinella spiralis Antibody by Indirect ELISA Using Recombinant Protein as Antigen

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Abstract

Objective To study the specificity, sensitivity of diagnostic antigen for Trichinella spiralis (T. s) . Methods T668 recombinant protein from highly efficient expression of newborn larvae stage-specific gene of T. s in E. coli as diagnostic antigen. By using negative and positive sera from rabbit, pig, human as the first antibody, and goat-anti-rabbit IgG, goat-anti-pig IgG, goat-anti-human IgG labeled with HRP as the secondary antibody, indirect ELISA method was established for detecting anti-T.s antibody, while excretorysecretory (ES) antigen of T.s muscle larvae was used as control. Results Sera from rabbit, pig and human were detected by T668 recombinant protein as antigen, which showed a positive rate of 100% with 0.016 µg/well. There was no difference on the results between the T668 recombinant antigen and the ES antigen for diagnosing T. sinfection. Conclusion The T668 recombinant antigen is promising in substituting ES antigen in the detection of anti-T. s antibody.

Key words Trichinella spiralis Recombinant protein Newborn larvae (NBL) stagespecific gene Indirect ELISA

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