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犬恶丝虫酪蛋白激酶2 β 亚基基因片段的克隆和原核表达

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Cloning and Prokaryotic Expression of Casein Kinase II Subunit Beta Gene Fragment of *Dirofilaria immitis*

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摘要 目的 对犬恶丝虫酪蛋白激酶2 β 亚基(Csnk2b)部分基因片段进行克隆、原核表达及免疫反应性分析。方法 根据犬恶丝虫cDNA文库中筛选出的Csnk2b部分基因片段设计引物,以含有插入Csnk2b部分基因片段的噬菌体DNA为模板,进行PCR扩增。产物亚克隆至原核表达载体,构建重组载体pGEX-4T-1-Csnk2b,双酶切鉴定。将其转入大肠埃希菌(E. coli)Rosetta(DE3)中,异丙基- β -D-硫代吡喃半乳糖苷(IPTG)诱导表达,通过十二烷基硫酸钠-聚丙烯酰胺凝胶电泳(SDS-PAGE)分析表达的重组蛋白,蛋白质印迹(Western blotting)以小鼠抗犬恶丝虫阳性血清为一抗,分析重组蛋白免疫反应性。结果 Csnk2b部分基因片段PCR扩增产物约为700 bp,测序结果与cDNA文库中筛选得到的序列一致。重组表达载体pGEX-4T-1-Csnk2b双酶切鉴定正确。SDS-PAGE结果显示, IPTG诱导后重组蛋白GST-Csnk2b表达,相对分子质量(Mr)约为45 000,与预期大小一致。Western blotting分析表明,重组蛋白能被小鼠抗犬恶丝虫阳性血清识别。结论 克隆并表达了犬恶丝虫Csnk2b重组蛋白,且该蛋白具有免疫反应性。

关键词: 犬恶丝虫; 酪蛋白激酶2 β 亚基; 原核表达

Abstract: Objective To clone and express the partial fragment of Csnk2b gene of *Dirofilaria immitis* in prokaryotic cells, and analyze the immunoreactivity. Methods The partial fragment of Csnk2b gene was amplified by PCR with a pair of specific primers. The PCR product was cloned into pMD18-T, and then sub-cloned to pGEX-4T-1 expression vector. The constructed plasmid pGEX-4T-1-Csnk2b was transformed into E. coli Rosetta (DE3) and followed by expression of the protein induced by IPTG. The recombinant protein was analyzed by SDS-PAGE and identified by Western blotting. Results The PCR product was about 700 bp. Enzyme digestion and DNA sequencing confirmed that the recombinant plasmid pGEX-4T-1-Csnk2b was constructed. SDS-PAGE results showed that the relative molecular weight (Mr) of the fusion protein (GST-Csnk2b) was about 45 000. GST-Csnk2b reacted positively with mouse anti-D. immitis serum. Conclusion The partial Csnk2b gene has been expressed in prokaryotic expression system and shows immunoreactivity.

Keywords: *Dirofilaria immitis*; Csnk2b; Prokaryotic expression

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