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### 粉尘螨第7类变应原 (Der f 7) 基因的克隆表达及免疫学特性鉴定

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### Cloning, Expression and Identification of Der f 7 gene from Dermatophagoides farinae and its Immunological Characteristics

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**摘要** 目的 克隆和表达粉尘螨第7类变应原基因, 并鉴定重组蛋白的免疫原性。方法 提取粉尘螨总RNA, 根据GenBank提供的Der f 7 编码区 (CDS) (登录号为AY 283292) 序列设计特异性引物, 逆转录PCR (RT-PCR) 克隆Der f 7基因。将测序正确的目的片段克隆至pET-32a表达载体, 得到的重组质粒pET-32a-Der f 7在大肠埃希菌 (E. coli) BL21 (DE3) 中用异丙基-β-D-硫代半乳糖苷 (IPTG) 诱导表达, 通过镍离子亲和层析纯化目的蛋白。十二烷基硫酸钠-聚丙烯酰胺凝胶电泳 (SDS-PAGE) 检测目的蛋白表达和纯化结果, 蛋白质印迹 (Western Blotting) 分析重组蛋白的免疫原性。结果 RT-PCR结果显示, Der f 7基因片段大小约为650 bp。测序结果表明, Der f 7基因片段与已发表的粉尘螨Der f 7基因 (登录号为FJ436108) 同源率为99%。SDS-PAGE结果显示, 重组质粒pET32a-Der f 7在BL21 (DE3) 中高效表达, 重组蛋白相对分子质量 (Mr) 约为23 000。Western Blotting分析结果表明, Der f 7重组蛋白可被尘螨过敏患者血清识别。结论 成功构建了粉尘螨第7类变应原的原核表达载体, 并获得具有免疫原性的Der f 7重组蛋白。

**关键词:** 粉尘螨 Der f 7 变应原 表达 纯化

**Abstract:** Objective To clone and express Der f 7 gene of Dermatophagoides farinae, and identify its immunogenicity. Methods Total RNA was extracted from D. farinae mites. A reference sequence (Accession No. AY283292) was used to design specific primers. The Der f 7 gene fragment was amplified by RT-PCR, and cloned into pET-32a vector. The recombinant plasmid was transformed into E. coli BL21 (DE3) and induced with IPTG for protein expression. The recombinant protein was purified by Ni<sup>2+</sup> chelating affinity chromatography and analyzed by SDS-PAGE and Western blotting. Results The Der f 7 gene fragment was about 650 bp, and shared 99% homology with the published one (Accession No. FJ436108). SDS-PAGE result showed its relative molecular weight (Mr) of 23 000. The recombinant protein showed appropriate combination ability with IgE in sera of mite allergic patients. Conclusion Der f 7 gene has been expressed in prokaryotic expression system and shows allergenicity.

**Keywords:** Dermatophagoides farinae Der f 7 Allergen Expression Purification

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