Red重组系统及在微生物基因敲除中的应用 The Red Recombination System and Its Application to Gene Knockout in Microorganism

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在完成了对各种微生物基因组的测序以后,功能基因学的研究变得尤为重要。研究基因功能最直接的方法 便是将待研究的基因失活。最初构建基因突变体是采用大肠杆菌的RecA系统,但是RecA重组系统操作复杂,重组 效率低。最近建立了Red重组系统,该系统由3个蛋白组成:α蛋白(即λ核酸外切酶),β蛋白,Gam蛋白。应用Red系▶文章反馈 统进行基因敲除,可以直接利用线性打靶DNA,两侧同源臂长度在35~60 bp即可发生同源重组,且重组效率高。 Abstract: Since many DNA-sequencing projects of varied microorganisms have been completed, studies on their functional genomics become more important. Inactivation of an interesting gene is a direct method to characterize its function. Though the Esherichia coli RecA recombination system can be used to produce gene mutants, it needs a complex manipulation process. Furthermore, its efficiency is very low. Recently a Red recombination system was developed. This recombination system consists of three proteins: α protein (λ exonuclease), β protein and Gam protein. In this system, the linear targeting DNA which contains a selectable marker flanked with a homologous region as short as only 35~60 bp can be directly targeted for gene knock-out with a higher efficiency.

Red重组系统 基因敲除 _ 抗药性基因 Key words _ Red recombination system _ gene knock-out _ resistant 关键词 gene

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Abstract

Key words

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