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## 重组人CK2β亚基的原核表达、纯化与鉴定

Prokaryotic Expression, Purification and Characterization of Recombinant Human Protein Kinase CK2  $\beta$  Subunit

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中文关键词: <u>人蛋白激酶CK2β亚基</u> 表达载体pT7-7 原核表达 蛋白质纯化 酶动力学分析

英文关键词: <u>human protein kinase CK2β subunit</u> <u>expression vector pT7-7</u> <u>prokaryotic expression protein purification</u> <u>enzyme</u> <u>kinetic analysis</u>

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中文摘要:

将构建成功的人蛋白激酶CK2β亚基cDNA的重组质粒,转化大肠杆菌BL21 (DE3),在IPTG诱导下表达.表达蛋白大多数以不溶形式存在.6L(约10.2 g)表达菌抽提得到约20 mg的可溶性表达产物,通过P11磷酸纤维素一步层析分离,得到6.8 mg纯化蛋白.SDS-聚丙烯酰胺凝胶电泳结果显示纯化的蛋白为一分子质量26 ku的单一蛋白带.蛋白质印迹结果证明:纯化的表达产物与抗人CK2β抗体可发生特异性免疫反应.CK2β亚基对CK2α有激活作用,纯化的CK2α和β亚基在等摩尔混合时即可组成有最大生物活性的全酶.实验结果有力地证明了克隆表达与纯化的重组蛋白是人蛋白激酶CK2β亚基.

## 英文摘要:

Protein kinase CK2 is a heterotetramer ser/thr protein kinase composed by two catalytic subunits ( $\alpha$  or  $\alpha'$ ) and two regulatory subunits ( $\beta$ ). The recombinant plasmid containing human CK2 $\beta$  subunit cDNA was transformed into *E. coli* BL21(DE3) and expressed induced by IPTG. Most of the expressed CK2 $\beta$  proteins were insoluble. From 6 L(about 10.4 g) bacteria, 20 mg soluble protein was extracted from the insoluble pellet and purified by a single step P11 phosphocellulose chromatography, the yield of purified CK2 $\beta$  subunit was 6.8 mg. SDS-PAGE analysis of the purified protein showed only one band with molecular mass of 26 ku. Western blot analysis confirmed that the expressed product was human CK2 $\beta$  subunit. Addition of the CK2 $\beta$  subunit to CK2 $\alpha$  subunit led to maximum stimulation at a 1:1 molar ratio of both subunits. These results demonstrate d strongly that the cloned, expressed and purified recombinant protein was human CK2 $\beta$  subunit. The large amount of purified recombinant CK2 $\beta$  protein lays solid basis for further study directly the characteristics of the enzyme and the relationship of structure and function between CK2 $\beta$  subunit and its interacted proteins.

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