# pBIND

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| **载体编号 Vector ID** | V000416 |
| **抗性 Antibiotic Resistance** | Ampicillin |
| **长度 Length** | 6360 bp |
| **载体类型 Vector Type** | Mammalian Two-Hybrid System |

## 产品信息

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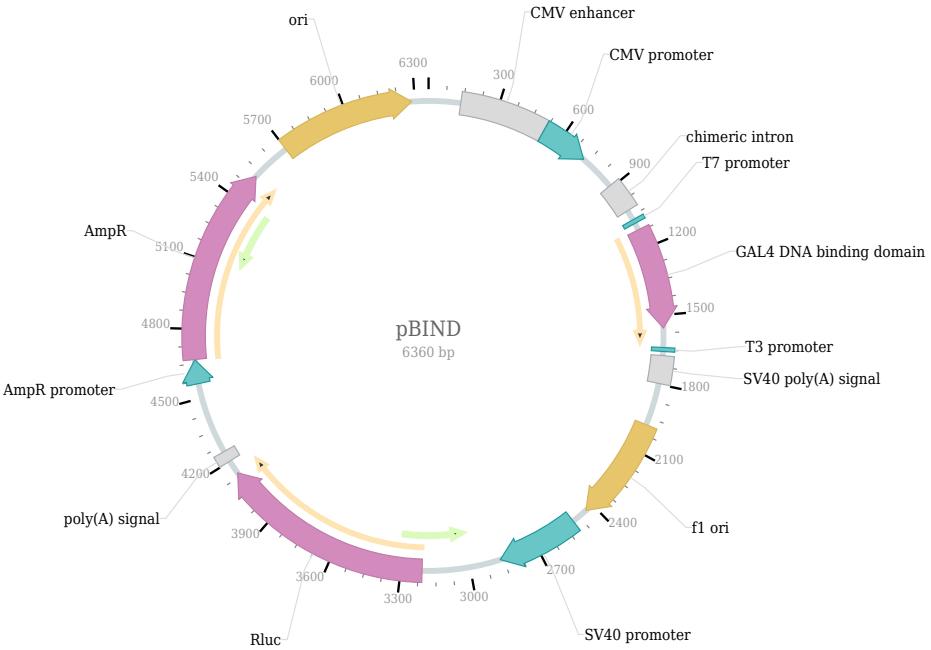
| **复制子 Replication Origin** | ori |
| **启动子 Promoter** | CMV |
| **备注 Comment** | The pBIND Vector is a high-copy plasmid in which the CMV immediate early promoter drives expression of a portion of the yeast GAL4 gene (amino acids 1–147) containing a DNA-binding domain. As with the pACT Vector, the DNAbinding domain sequence is flanked by a chimeric intron, a multiple cloning region, stop codons and an SV40 late polyadenylation region (Figures 4 and 5). The fusion gene region is flanked by T7 and T3 RNA polymerase promoters for the purpose of synthesizing sense and antisense RNA products, respectively. The Renilla luciferase gene on this vector is preceded by the SV40 early promoter and a growth hormone intron. Introns can increase protein expression through mRNA stability and nuclear to cytoplasmic transport effects (9–12). A synthetic polyadenylation sequence resides 3′ of the Renilla luciferase gene. The plasmid backbone contains an f1 origin of replication for the production of ssDNA and the β-lactamase gene (Ampr) for selection in E. coli. |

## 质粒图谱



## 复溶方法

1. Centrifuge at 5,000×g for 5 min.
2. Carefully open the tube and add sterile water to dissolve the DNA: add 20 µl for 5 µg plasmid, and 100 µl for 100 µg plasmid.
3. Close the tube and incubate for 10 minutes at room temperature.
4. Briefly vortex the tube and then do a quick spin to concentrate the liquid at the bottom. Speed is less than 5000×g.
5. Store the plasmid at -20 ℃.