基本信息

启动子:	Tac
平台编号	bio-114980
复制子:	ColE1 ori
终止子:	rrnB T1 terminator
质粒分类:	大肠杆菌载体；pMal 系列表达质粒
质粒大小:	5677bp
原核抗性:	氨苄青霉素 Amp
克隆菌株:	DH5α
培养条件:	37℃，有氧，LB
表达宿主:	BL21(DE3)
培养条件:	37℃，有氧，LB
诱导方式:	IPTG 或乳糖及其类似物

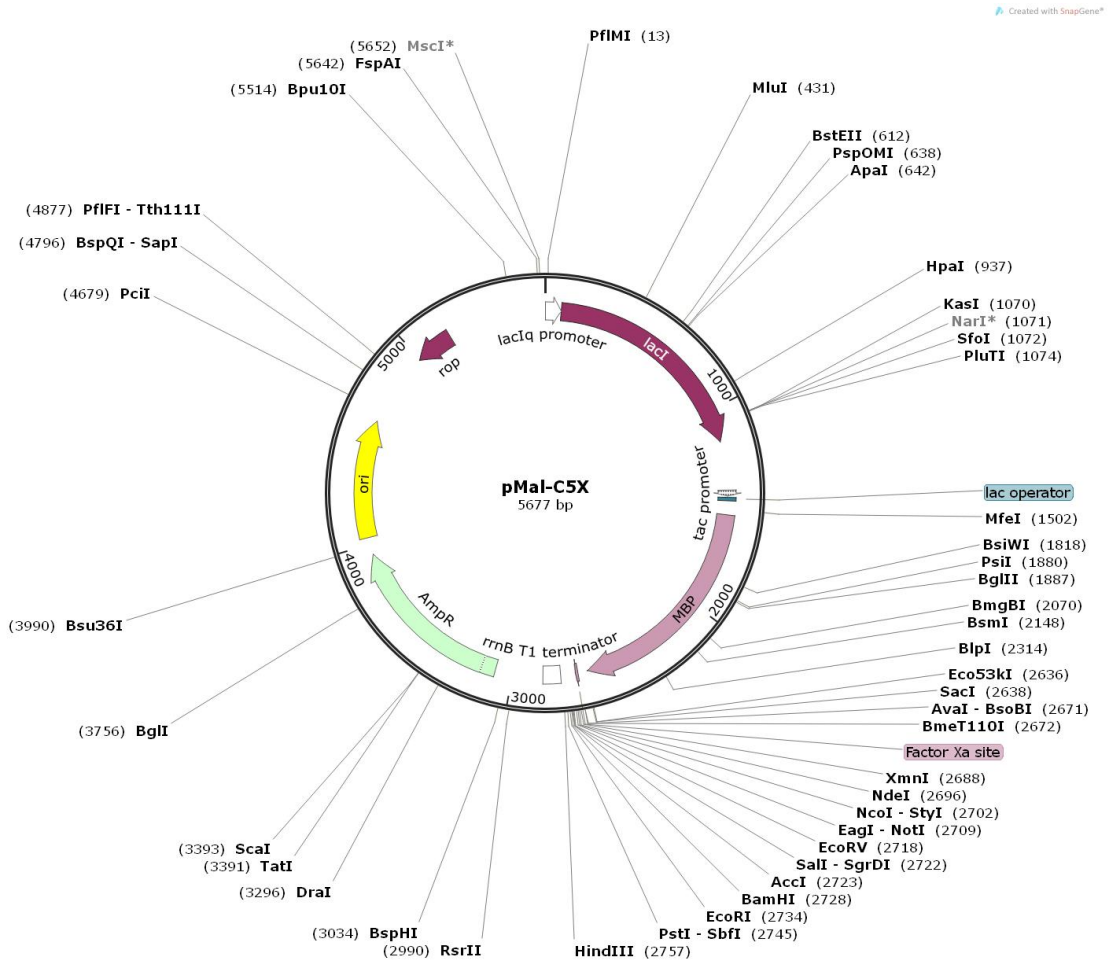
质粒简介

The vector pMAL-c5X is designed to produce maltose-binding protein (MBP) fusions, where the protein of interest can be cleaved from MBP with the specific protease Factor Xa .

MBP fusions made with this vector are expressed cytoplasmically. The MBP has been engineered for tighter binding to amylose resin. A gene or open reading frame is inserted into a restriction site of the vector polylinker, in the same translational reading frame as the malE gene (encoding maltose-binding protein). The fusion protein thus produced can be purified by amylose affinity chromatography. The sequence coding for the four amino acids Ile-Glu-Gly-Arg is present just upstream of the XmnI site. This allows the protein of interest to be cleaved from maltose-binding protein with the specific protease Factor Xa. Fragments inserted in the XmnI site (cleaves GAAGG↓ATTTC) will produce a fusion protein that, after Factor Xa cleavage, contains

no vector-derived residues on the protein of interest.

质粒图谱



质粒序列

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DEFINITION  synthetic circular DNA
ACCESSION   .
VERSION     .
KEYWORDS    pMal-c5X
SOURCE      synthetic DNA construct
  ORGANISM  synthetic DNA construct
REFERENCE   1  (bases 1 to 5677)
  AUTHORS   .
  TITLE     Direct Submission
  JOURNAL   Exported 2015-8-17 from SnapGene Viewer
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