Bio Academia
Manufacturer

AS ONE

cDNA Library, *S. pombe*; Meiosis

#02-705 500 ng

This cDNA library (plasmid DNA) is constructed from *Schizosaccharomyces pombe*, strain CD16-1(h+/h-) derived poly(A)$^+$ RNA at the state of meiosis by the Linker-Primer method (Ref.1) by Professor Hiroshi Nojima of Research Institute for Microbial Diseases, Osaka University. cDNAs in this library are unidirectionally cloned by using the oligo $(dT)_{18}$ linker primer which contains the restriction enzyme site of *Not* I, and *Bam*HI (*Bgl* II)-*Sma* I adaptor.
The pAP3*neo* vector used in this library can express the *S. pombe* genes in mammalian cells as it contains SV40 promoter. It also contains f1 ori which is necessary for ssDNA synthesis, and bacteriophage T7 and T3 promoter for RNA synthesis (see Figure). GenBank Accession No. AB003468

Application

PCR screening of known or unknown gene: Prepare the primers for the known or unknown gene (cDNA) and amplify the gene by PCR from this library followed by cloning to an appropriate vector. It is useful for large-scale protein productions, and preparation of probes, etc. Standard amplifying conditions: 35 cycles of PCR reactions using 10-100 ng of cDNA as a template. (Change the quantity of template and the number of cycles depending on the expression levels of mRNA of the genes.)

Specification

Quantity: 500 ng (40 ng/ul, 13ul) in 10 mM Tris-HCl-1mM EDTA (pH 7.5)
Quality: 1) Number of independent clones: 1.3×10^6
2) Average insert size : longer than 1 kb
Storage: -20°C

References

1. Kobori M *et al* " Large scale isolation of osteoclast-specific genes by an improved method involving the preparation of a subtracted cDNA library." *Genes Cells* **3**: 459-475 (1998) PMID: 9753427
2. Sambrook J and Russell DW *Molecular Cloning* Chapter 11 "Preparation of cDNA libraries and gene identification." CSHL Press (2001)

Note

* This library is to be used only by the the purchaser. It is not allowed to amplify and transfer the library to a third person.
* For custom order of cDNA cloning from the libraries, construction of protein expression systems, and production and purification of proteins, contact info@asone-int.com

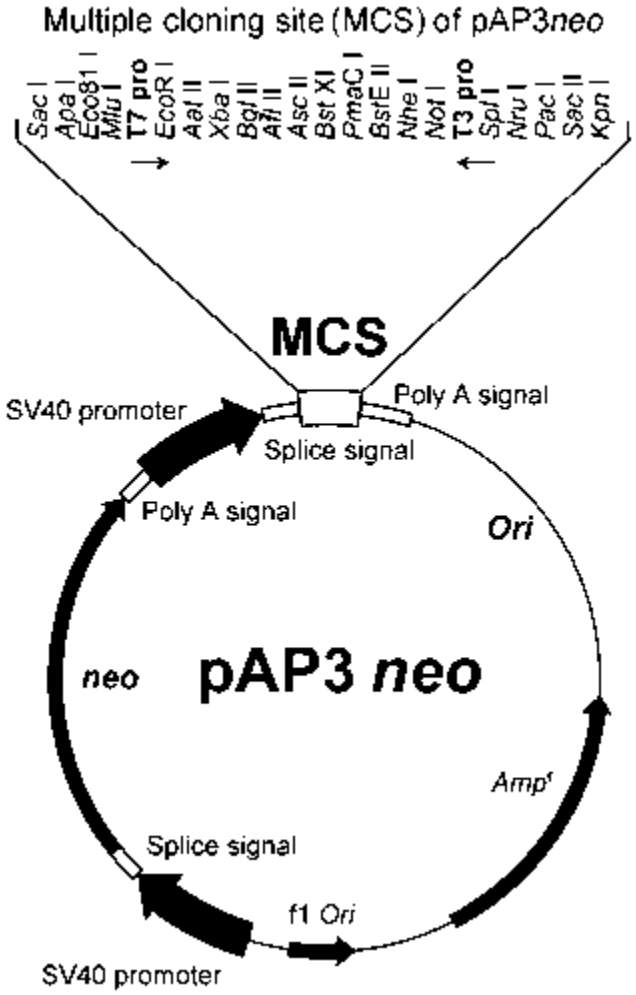

Fig. Structure of pAP3*neo* and the restriction sites