

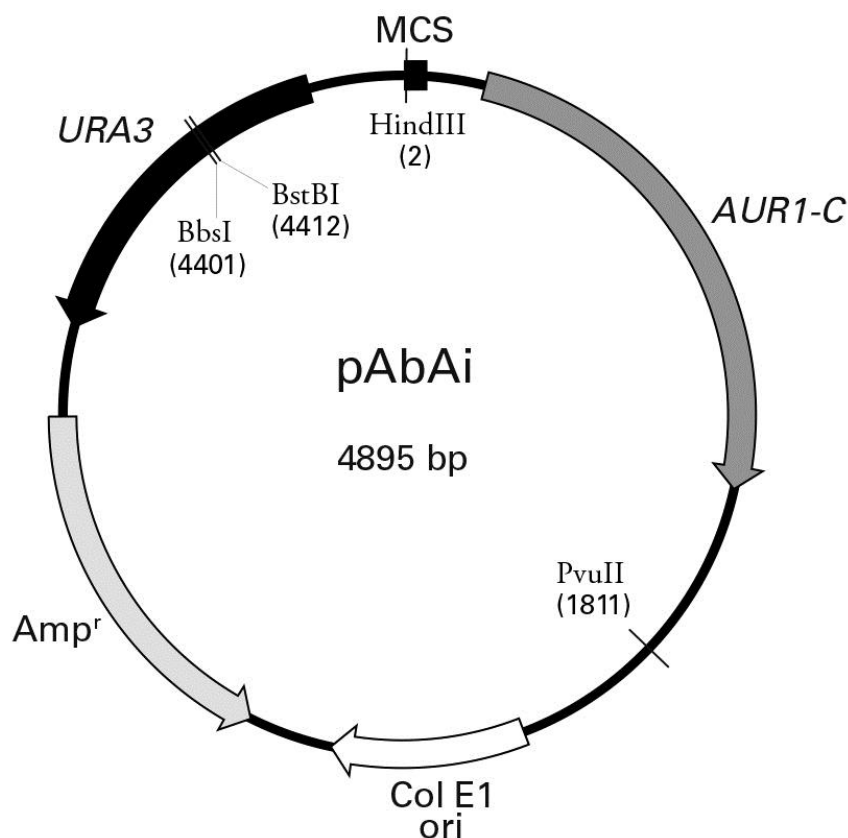
# Vector Map

## pAbAi Vector

### Catalog No.

(Not sold separately)

Sold as a part of 630491



	<u>HindIII</u>	<u>SacI</u>	<u>KpnI</u>	<u>SmaI</u>	<u>SalI</u>	<u>XhoI</u>
1	AAGCTTGAAT	TCGAGCTCGG	TACCCGGGGA	TCTGTCGACC	TCGAGGCATG	
	TTCGAACTTA	AGCTCGAGCC	ATGGGCCCT	AGACAGCTGG	AGCTCCGTAC	

**Figure 1. pAbAi Vector map and multiple cloning site (MCS).** pAbAi cannot be propagated episomally in yeast; it can only be stably maintained through integration into the host genome. Integration is accomplished via homologous recombination between the vector's *URA3* gene and the *ura3-52* locus of the yeast strain provided in the Matchmaker® Gold Yeast One-Hybrid System. To use pAbAi in a one-hybrid assay, clone one or more copies of a cis-acting element into the MCS. To facilitate recombination of the vector with the host's *ura3-52* locus, linearize the vector with either BbsI or BstBI, then introduce the linearized vector into competent yeast cells using the protocols in the Matchmaker Gold Yeast One-Hybrid Library Screening System User Manual (PT4087-1). Insertion of your target sequence may alter the basal expression of *AUR1-C*. Therefore, before starting a one-hybrid analysis, basal levels of *AUR1-C* should be determined as described in the User Manual.

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## Location of Features

- Multiple cloning site: 1–45
- *AUR1-C* (aureobasidin A resistance gene): 185–1390
- Col E1 origin of replication: 2159–2609
- Amp<sup>R</sup> (ampicillin resistance gene): 2806–3666 (complementary)
- *URA3*: 3876–4678 (complementary)
- Sequencing primer: 4655–4679  
5'–GTT CCT TAT ATG TAG CTT TCG ACA T–3'

## Propagation in *E. coli*

- Suitable host strains: DH5 $\alpha$  and other general purpose strains
- Selectable marker: plasmid confers resistance to ampicillin (100  $\mu$ g/ml) to *E. coli* hosts
- *E. coli* replication origin: Col E1
- Copy number: low

## Propagation in *S. Cerevisiae*

- Suitable host strain: Y1HGold
- Selectable marker: *URA3*

**NOTE:** The vector sequence was compiled from information in the sequence databases, published literature, and other sources, together with partial sequences we obtained. This vector has not been completely sequenced.

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This document has been reviewed and approved by the Quality Department.