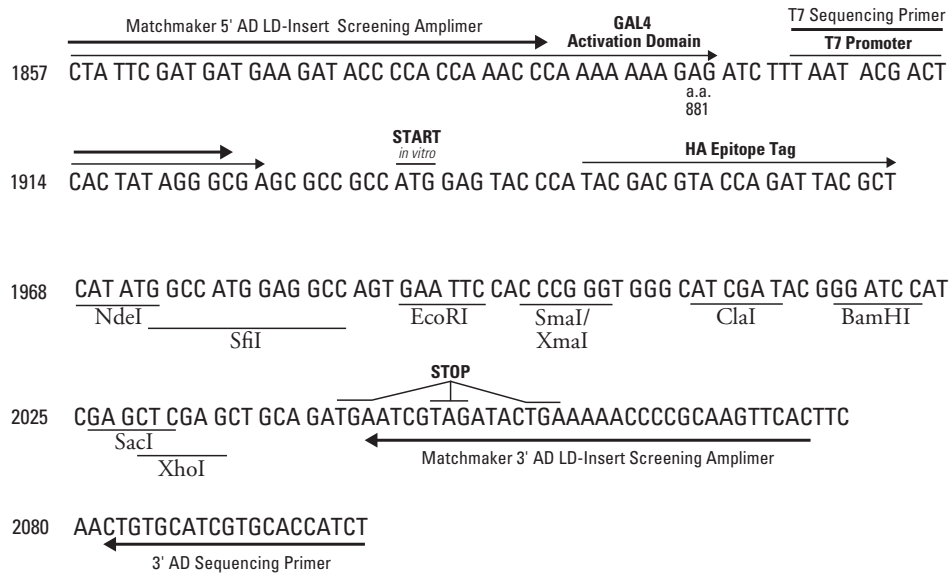
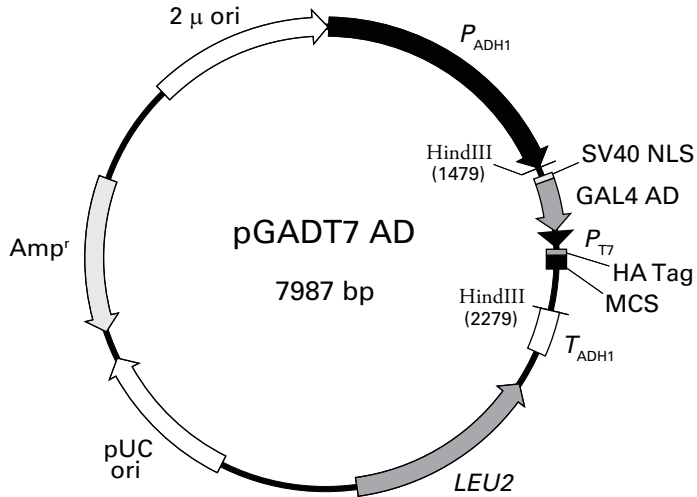


**pGADT7 AD Vector Information**

PT3249-5

Cat. Nos. 630442  
630489  
630491



**pGADT7 AD Vector Map and Multiple Cloning Site (MCS).**

**Description**

pGADT7 AD is a yeast expression vector that is designed to express a protein of interest fused to a GAL4 activation domain (AD; amino acids 768–881). Transcription of the GAL4 AD fusion is driven by the constitutively active *ADH1* promoter ( $P_{ADH1}$ ), and is terminated at the *ADH1* transcription termination signal ( $T_{ADH1}$ ). The GAL4 AD fusion contains an N-terminal SV40 nuclear localization signal (SV40 NLS; 1) that targets the protein to the yeast nucleus, and a hemagglutinin epitope tag (HA Tag), located between the GAL4 AD and the protein of interest, that allows the protein to be easily detected with HA-tag antibodies.

The T7 promoter ( $P_{T7}$ ), located just upstream of the HA tag sequence, allows *in vitro* transcription and translation of the HA-tagged protein of interest (without the GAL4 AD and the SV40 NLS). pGADT7 AD replicates autonomously in both *E. coli* and *S. cerevisiae* from the pUC and 2  $\mu$  ori, respectively. The vector also contains an ampicillin resistance gene (*Amp<sup>r</sup>*) for selection in *E. coli* and a *LEU2* nutritional marker for selection in yeast.

(010312)



**Clontech**

**United States/Canada**  
800.662.2566

**Asia Pacific**  
+1.650.919.7300

**Europe**  
+33.(0)1.3904.6880

**Japan**  
+81.(0)77.543.6116

Clontech Laboratories, Inc.  
A Takara Bio Company  
1290 Terra Bella Ave.  
Mountain View, CA 94043  
Technical Support (US)  
E-mail: tech@clontech.com  
www.clontech.com

## Use

pGADT7 AD is the AD Cloning Vector provided in Clontech's Matchmaker™ Gold Yeast One- and Two-Hybrid Screening Systems (Cat. Nos. 630491 and 630489, respectively). The vector allows the generation of GAL4 AD fusion proteins from either a gene of interest or a cDNA library. **Important:** Genes must be cloned into the MCS so that they are in-frame with the GAL4 AD and HA tag coding sequences. The vector can also be used as a negative "prey" control for the One-Hybrid System.

GAL4 AD/HA-tagged fusion proteins expressed by the vector can be detected with either our GAL4 AD Monoclonal Antibody (Cat. No. 630402) or our HA-Tag Polyclonal Antibody (Cat. No. 631207). Note: *In vitro* transcription/translation from the T7 promoter, located between the GAL4 AD and HA tag sequences, produces an HA-tagged protein that lacks the GAL4 AD. Such proteins can be detected by the HA-Tag Polyclonal Antibody, but not the GAL4 AD Monoclonal Antibody.

## Location of features

- $P_{ADH1}$  (full-length *S. cerevisiae ADH1* promoter): 7–1478
- GAL4 AD (GAL4 activation domain with SV40 Nuclear Localization Signal [NLS])  
SV40 NLS: 1500–1556  
GAL4 (amino acids 768–881): 1560–1898
- $P_{T7}$  (T7 RNA polymerase promoter): 1904–1926
- HA Tag (hemagglutinin epitope tag): 1941–1967
- MCS (multiple cloning site): 1968–2040
- $T_{ADH1}$  (*S. cerevisiae ADH1* Terminator): 22879–2604
- *LEU2* coding sequences: 2722–3813 (complementary)
- pUC ori (pUC replication origin): 4580–5417
- Amp<sup>r</sup> (ampicillin resistance gene): 5573–6433 (complementary)
- 2  $\mu$  ori (Yeast 2  $\mu$  replication origin): 6997–7987

## Location of primers

- T7 Sequencing Primer: 1904–1924
- 3' AD Sequencing Primer: 2101–2082
- Matchmaker 5' AD LD-Insert Screening Amplimer (Cat. No. 630433): 1857–1888
- Matchmaker 3' AD LD-Insert Screening Amplimer (Cat. No. 630433): 2077–2045

## Propagation in *E. coli*

- Suitable host strains: DH5 $\alpha$ , DH10 & other general purpose strains
- Selectable marker: plasmid confers resistance to ampicillin (100  $\mu$ g/ml) to *E. coli* hosts
- *E. coli* replication origin: pUC
- Copy number: ~500
- Plasmid incompatibility group: pMB1/Col E1

## Propagation in *S. cerevisiae*

- Suitable host strains: Y1HGold, Y2HGold, AH109(MAT $\alpha$ ), Y187(MAT $\alpha$ ), Y190(MAT $\alpha$ ), SFY526(MAT $\alpha$ ), CG1945(MAT $\alpha$ ), HF7c(MAT $\alpha$ )
- Selectable marker: *LEU2*
- *S. cerevisiae* origin: 2  $\mu$

## Reference

1. Chien, C.T., Bartel, P.L., Sternglanz, R. & Fields, S. (1991) *Proc. Natl. Acad. Sci. USA* **88**:9578–9582.

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

## Notice to Purchaser

Clontech products are to be used for research purposes only. They may not be used for any other purpose, including, but not limited to, use in drugs, in vitro diagnostic purposes, therapeutics, or in humans. Clontech products may not be transferred to third parties, resold, modified for resale, or used to manufacture commercial products or to provide a service to third parties without written approval of Clontech Laboratories, Inc.

Clontech, Clontech Logo and all other trademarks are the property of Clontech Laboratories, Inc., unless noted otherwise. Clontech is a Takara Bio Company. ©2012 Clontech Laboratories, Inc.