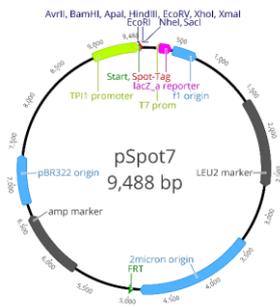


pSpot7 plasmid for expression of N-terminal Spot-Tag® fusion proteins in *S. cerevisiae*



For plasmid sequence, please visit [www.chromotek.com](http://www.chromotek.com).

**Location of features**

- TPI1 promoter: 8914-9
- Start codon (ATG): 10-12
- Spot-Tag®: 13-48
- MCS: 53-112
- T7 primer site: 230-248
- Leucin marker (LEU2): 1564-2670
- 2µ replication origin: 3396-4865
- Ampicillin resistance gene: 5723-6583
- pRB322 replication origin: 6738-7357

Product	Code	Size
pSpot7	ev-7	1.25 µg
Vector type	yeast expression vector	
Tag	Spot-Tag® (PDRVRAVSHWSS) N-terminal	
MCS	BamHI, AvrII, Apal, HindIII, EcoRV, XhoI, XmaI, NheI, SacI	
Promoter	TPI1	
Induction	constitutive	
Host cells	<i>Saccharomyces cerevisiae</i> ( <i>leuΔ</i> )	
Selection	leucin ( <i>S. cerevisiae</i> ) ampicillin ( <i>E. coli</i> )	
Replication	2µ ( <i>S. cerevisiae</i> ) pBR322 ( <i>E. coli</i> )	
Use	Expression of a protein of interest fused to Spot-Tag® (N-terminal) in <i>S. cerevisiae</i> .	

**Vector description**

The plasmid pSpot7 is an expression vector for Spot-Tag® fusion proteins in *S. cerevisiae*. After cloning the protein of interest (POI) into the multiple cloning site (MCS) provided by pSpot7, the Spot-Tag® (sequence: PDRVRAVSHWSS) will be fused to the N-terminus of the POI.

**Expression in *S. cerevisiae***

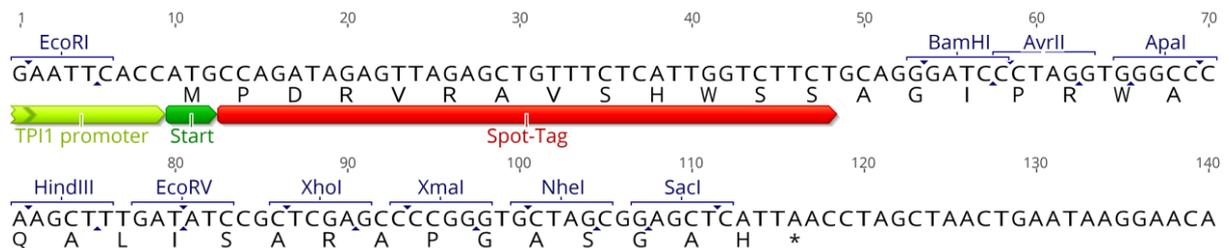
Transform a suitable *leu2* deletion strain of *S. cerevisiae* using pSpot7 and standard methods and grow on media lacking leucine. Protein expression is constitutive under the strong triosephosphate isomerase (TPI1) promoter. The resulting Spot-Tag® fusion protein can be purified using the ChromoTek Spot-Trap®.

**Propagation in *E. coli***

Suitable host strains for propagation in *E. coli* include DH5alpha, HB101, XL1-Blue, and other general purpose strains. The vector confers resistance to ampicillin (100 µg/ml) to *E. coli* hosts.

Note: The plasmid DNA was isolated from dam<sup>+</sup>-methylated *E. coli*. Therefore some restriction sites are blocked by methylation. If you wish to digest the vector using such sites, you will need to transform the vector into a dam<sup>-</sup> host and make fresh DNA

**Multiple cloning site (MCS)**



**Notice to Purchaser:**

This plasmid was designed and generated by Dr. Philipp Kaiser from the Naturwissenschaftliches und Medizinisches Institut (NMI) at the University of Tübingen, Germany and is distributed by ChromoTek GmbH. Please acknowledge Dr. Philipp Kaiser (NMI, Tübingen, Germany) and ChromoTek GmbH (Martinsried, Germany) when using or redistributing this vector. The development of this plasmid was supported by a grant "Zentrales Innovationsprogramm Mittelstand" (ZIM) from the Federal Ministry for Economic Affairs and Energy of Germany.