

## pENTR\_L4R1\_pSKN7

SB-1105<sup>™</sup>

#### Description

An entry vector carrying the yeast constitutive promoter SKN7 (750 bp) flanked by two *att* sites (*att* L4 and *att* R1). It is one of promoters in the ATCC® Synthetic Biology Yeast Tool Kit (Detailed information is described and in the ATCC® Synthetic Biology Solutions User Guide).

**Deposited As:** R Weiss, Massachusetts Institute of Technology

Volume: 2 µg to 3 µg

### Storage Conditions

Storage conditions: 2°C to 8°C

#### Intended Use

This product is intended for laboratory research use only. It is not intended for any animal or human therapeutic use, any human or animal consumption, or any diagnostic use.

### BSL<sub>1</sub>

ATCC determines the biosafety level of a material based on our risk assessment as guided by the current edition of *Biosafety in Microbiological and Biomedical Laboratories* (*BMBL*), U.S. Department of Health and Human Services. It is your responsibility to understand the hazards associated with the material per your organization's policies and procedures as well as any other applicable regulations as enforced by your local or national agencies.



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## Certificate of Analysis

For batch-specific test results, refer to the applicable certificate of analysis that can be found at www.atcc.org.

#### Insert Information

Insert size (kb): 0.75

**M13 forward:** 5'- GTTTTCCCAGTCACGAC - 3' **M13 reverse:** 5'- CAGGAAACAGCTATGAC - 3'

#### **Vector Information**

Construct size (kb): 3.197

Type of vector: Entry vector

Markers: kanR

## Handling Procedures

Before opening the vial, centrifuge at 6,000 x g for 30 seconds. Add 30  $\mu$ L of Molecular Grade Water and incubate the vial at 4°C overnight to dissolve the DNA. Each vial contains 2-3  $\mu$ g plasmid DNA (measured by PicoGreen® dsDNA quantitation assay).

#### Material Citation

If use of this material results in a scientific publication, please cite the material in the following manner: pENTR\_L4R1\_pSKN7 (ATCC SB-1105)

#### References



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References and other information relating to this material are available at www.atcc.org.

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#### Revision

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