

Escherichia coli* – *Staphylococcus aureus* Shuttle Vector pCN57, Recombinant in *Staphylococcus aureus

Catalog No. NR-46158

This reagent is the tangible property of the U.S. Government.

For research use only. Not for use in humans.

Contributor:

Richard P. Novick, M.D., Departments of Microbiology, Medicine and Molecular Pathogenesis, New York University School of Medicine, New York, New York, USA

Manufacturer:

BEI Resources

Product Description:

NR-46158 is *Staphylococcus aureus* (*S. aureus*), strain RN 4220 (RN9623, NRS623) containing the *Escherichia coli* (*E. coli*)-staphylococcal shuttle vector pCN57. Vector pCN57 contains the *E. coli* ColE1 replication origin, the *S. aureus* pT181 *cop-wt-repC* replicon, the *P_{blaz}* promoter and a promoterless β -lactamase reporter gene, *gfpmut2*. Transformation with pCN57 leads to ampicillin resistance in *E. coli* and erythromycin resistance in *S. aureus*.¹

The vector sequence was deposited into GenBank as pNR-46158 (GenBank: [KP255997](https://www.ncbi.nlm.nih.gov/nuccore/KP255997)). The complete plasmid sequence and map are provided on the BEI Resources webpage.

Vector pCN57 is a member of a series of novel shuttle vectors that were developed using PCR-designed cassettes to allow for easy exchange of vector components. The base shuttle vectors are comprised of (i) a staphylococcal replicon (pT181-based low-copy number, high-copy-number or thermosensitive replicons or pl258-based low-copy-number theta replicon), (ii) a staphylococcal selectable marker (erythromycin, tetracycline, chloramphenicol, kanamycin or spectinomycin resistance), (iii) an *E. coli* ColE1-based replicon (iv) an *E. coli* selectable marker (ampicillin resistance) and (v) a pUC19-derived expanded multiple cloning site (MCS). Additionally, some of the vectors may contain a staphylococcal ϕ 11 phage fragment, staphylococcal pathogenicity island SaPI1 fragment, an inducible or constitutive promoter, and reporter genes.¹

Material Provided:

Each vial contains approximately 0.5 mL of bacterial culture in Casitone-Yeast broth containing 0.1 M glycerol phosphate and 10 μ g/mL erythromycin supplemented with 10% glycerol.

Note: If homogeneity is required for your intended use, please purify prior to initiating work.

Packaging/Storage:

NR-46158 was packaged aseptically in cryovials. The product

is provided frozen and should be stored at -60°C or colder immediately upon arrival. For long-term storage, the vapor phase of a liquid nitrogen freezer is recommended. Freeze-thaw cycles should be avoided.

Growth Conditions:

Media:

Casitone-Yeast broth containing 0.1M glycerol phosphate and 10 μ g/mL erythromycin

Tryptic Soy agar containing 10 μ g/mL erythromycin

Incubation:

Temperature: 37°C

Atmosphere: Aerobic

Propagation:

1. Keep vial frozen until ready for use, then thaw.
2. Transfer the entire thawed aliquot into a single tube of broth.
3. Use several drops of the suspension to inoculate an agar slant and/or plate.
4. Incubate the tube, slant and/or plate at 37°C for 1day.

Citation:

Acknowledgment for publications should read “The following reagent was obtained through BEI Resources, NIAID, NIH: *Escherichia coli* – *Staphylococcus aureus* Shuttle Vector pCN57, Recombinant in *Staphylococcus aureus*, NR-46158.”

Biosafety Level: 2

Appropriate safety procedures should always be used with this material. Laboratory safety is discussed in the following publication: U.S. Department of Health and Human Services, Public Health Service, Centers for Disease Control and Prevention, and National Institutes of Health. [Biosafety in Microbiological and Biomedical Laboratories \(BMBL\)](#). 6th ed. Washington, DC: U.S. Government Printing Office, 2020.

Disclaimers:

You are authorized to use this product for research use only. It is not intended for human use.

Use of this product is subject to the terms and conditions of the BEI Resources Material Transfer Agreement (MTA). The MTA is available on our Web site at www.beiresources.org.

While BEI Resources uses reasonable efforts to include accurate and up-to-date information on this product sheet, neither ATCC® nor the U.S. Government makes any warranties or representations as to its accuracy. Citations from scientific literature and patents are provided for informational purposes only. Neither ATCC® nor the U.S. Government warrants that such information has been confirmed to be accurate.

This product is sent with the condition that you are responsible for its safe storage, handling, use and disposal. ATCC® and the U.S. Government are not liable for any damages or injuries arising from receipt and/or use of this product. While reasonable effort is made to ensure authenticity and reliability of materials on deposit, the U.S. Government, ATCC®, their suppliers and contributors to BEI Resources are not liable for

damages arising from the misidentification or misrepresentation of products.

Use Restrictions:

This material is distributed for internal research, non-commercial purposes only. This material, its product or its derivatives may not be distributed to third parties. Except as performed under a U.S. Government contract, individuals contemplating commercial use of the material, its products or its derivatives must contact the contributor to determine if a license is required. U.S. Government contractors may need a license before first commercial sale. This material may be subject to third party patent rights.

References:

1. Charpentier E., et al. "Novel Cassette-Based Shuttle Vector System for Gram-Positive Bacteria." Appl. Environ. Microbiol. 70 (2004): 6076-6085. PubMed: 15466553.

ATCC® is a trademark of the American Type Culture Collection.

