**Product Sheet** 



## pGreenTIR 87572™

## Description

This is a green fluorescent protein (GFP) cloning vector designed specifically for use in the construction of prokaryotic transcriptional fusions. The gfp gene, along with the translation initiation region (TIR) can be excised with one of eight restriction enzymes (HindIII, PstI, Sall, Xbal, BamHI, Smal, SacI or EcoRI). The gfp allele in pGreenTIR contains both the F64L and S65T mutations that increase protein solubility and cause a ?red-shift? in the excitation maximum from 395 to 490 nm. The vector was constructed by cloning a mutant GFP gene into the EcoRI site of pUC1813. The resulting construct was mutagenized via PCR to 1) restore the 5? end of the gene to wild-type, 2) incorporate an upstream translational enhancer and 3) change the Shine-Delgarno region (and surrounding bp) to consensus.

-----Gene 191: 149-153, 1997

**Clone type:** Vector **Shipping information:** *Escherichia coli* containing the plasmid in glycerol stock

Storage Conditions Product format: Frozen Storage conditions: -80°C or colder

## 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.

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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.

#### Certificate of Analysis

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

#### **Vector Information**

Construct size (kb): 3.483999967575073 Vector name: pGreenTIR (plasmid) Construction: pUC1813 Coding sequence: gfp Enhancer: from the phage T7 gene10 Markers: ampR MCS: EcoRI...HindIII; HindIII...EcoRI Operator: lac Promoters: lac Replicon: pMB1 Ribosome-binding site: Shine-Dalgarno sequence Translational enhancer: from the phage T7 gene10

#### **Growth Conditions**

**Medium:** ATCC Medium 1227: LB Medium (ATCC medium 1065) with 50 mcg/ml ampicillin **Temperature:** 37°C



## pGreenTIR

#### Notes

Restriction digests of the clone gave the following sizes (in kb): EcoRI 2.6, 0.75; KpnI 3.5. ATCC Staff

#### **Material Citation**

If use of this material results in a scientific publication, please cite the material in the following manner: pGreenTIR (ATCC 87572)

#### References

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

#### Warranty

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### Revision

This information on this document was last updated on 2024-10-25

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