



#### **PURPOSE**

(Empty Backbone) Localization: Protein Expression Vector, Excitation: 488, Emission: 507

#### **DEPOSITING LAB**

Michael Davidson

#### **PUBLICATION**

Michael Davidson Empty Backbones (unpublished) (How to cite ↓)

### **SEQUENCE INFORMATION**

Sequences (3)

ORDERING								
Item	Catalog #	Description	Quantity	Price (USD)				
Plasmid	54762	Standard format: Plasmid sent in bacteria as agar stab	1	\$85 *	Add to Cart			
Cloning Grade DNA	54762- DNA.cg	2 μg of cloning grade DNA in Tris buffer <u>More Information.</u> ↓	1	\$105	Add to Cart			

<sup>\*</sup> Login to view industry pricing.

# **BACKBONE**

Vector backbone: EGFP-pBAD

Backbone size (bp): 4800

Vector type: Bacterial Expression

### Tags / Fusion Proteins:

EGFP (C terminal on backbone)6xHis (N terminal on backbone)

## **GROWTH IN BACTERIA**

Bacterial Resistance(s): Ampicillin, 100 µg/mL

Growth Temperature: 37°C Growth Strain(s): DH5alpha Copy number: High Copy

#### **GENE/INSERT**

Gene/Insert name: None
Tags / Fusion Proteins:

EGFP (C terminal on backbone)6xHis (N terminal on backbone)

# RESOURCE INFORMATION

#### **Supplemental Documents:**

• EGFP-pBAD Data Supplement

## Articles Citing this Plasmid:

• 12 References

#### **TERMS AND LICENSES**

# Academic/Nonprofit Terms:

• <u>UBMTA</u>

# Industry Terms:

• Industry MTA

#### Trademarks:

• Zeocin® is an InvivoGen trademark.

## **DEPOSITOR COMMENTS**

Contains 6x-Histidine tag for protein purification. Excitation = 488; Emission = 507

# Information for Cloning Grade DNA (Catalog # 54762-DNA.cg)

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### **PURPOSE**

Cloning grade DNA is suitable for use in PCR, cloning reactions, or transformation into E. coli. The purity and amount is not suitable for direct transfections.

# DELIVERY

Amount: 2 μg

**Guaranteed Concentration:** 100 ng/μl +/- 5 ng/μl

Pricing: \$105 USD

Storage: DNA can be stored at 4°C (short term) or -20°C (long term).

Academic/Nonprofit Terms:		
• <u>UBMTA</u>		
Industry Terms:		
<ul> <li>Industry MTA</li> </ul>		

### **QUALITY CONTROL**

Addgene has verified this plasmid using Next Generation Sequencing. Results are available <u>here</u>

**TERMS AND LICENSES** 

How to cite this plasmid (Back to top ↑)

These plasmids were created by your colleagues. Please acknowledge the Principal Investigator, cite the article in which the plasmids were described, and include Addgene in the Materials and Methods of your future publications.

For your Materials & Methods section:

 $EGFP-pBAD\ was\ a\ gift\ from\ Michael\ Davidson\ (Addgene\ plasmid\ \#\ 54762\ ;\ http://n2t.net/addgene:54762\ ;\ RRID:Addgene\_54762)$ 

