

CORRECTION

# Correction: Gateway Vectors for Efficient Artificial Gene Assembly *In Vitro* and Expression in Yeast *Saccharomyces cerevisiae*

The PLOS ONE Staff

There are errors in the Results, [Table 2](#) and [Table 3](#), which were introduced during the typesetting process. The publisher apologizes for these errors.

There is an error in the penultimate sentence of the third paragraph of the “Construction of Yeast Gateway Vectors for One-step Gene Assembly” subsection of the Results. The correct sentence is: We compared the fluorescence produced by yEVENus-Cln2<sub>PEST</sub>-NLS and yEVENus-NLS when expressed by the constitutive promoter ADH1 (Fig 4B, C).

There is an error in [Table 2](#), “Entry clones created in this study.” Please see the corrected [Table 2](#) here.

**Table 2. Entry clones created in this study.**

Plasmid	Promoter
pYS1	<i>S. cerevisiae</i> CUP1
pYS2	<i>S. pombe</i> ADH1
pYS3	<i>S. cerevisiae</i> TEF
pYS6	TetO <sub>7</sub> -CYC1TATA
pYS7	TetO <sub>2</sub> -CYC1TATA
Plasmid	ORF
pCG32	yEGFP
pDHM7	yEGFP-Cln2 <sub>PEST</sub>
pCG55	yEVENus
pYS61	yEVENus-NLS* <sup>1</sup>
pCG98	yEVENus-Cln2 <sub>PEST</sub> -NLS
pCG40	mCherry
pYS60	mCherry-NLS
pYS19	tTA
pYS20	rTA
pYS58	AIDtTA
pYS57	AIDrtTA
pCG72	OsTIR1-9Myc

Promoter Entry clones were created with pDONR221P5-P2 and ORF Entry clones with pDONR221P1-P5r. \*<sup>1</sup> Significant cytoplasmic fluorescence was observed when overexpressed, for example, by TEF promoter.

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There is an error in [Table 3](#), “Expression plasmid vectors constructed by Gateway recombination method in this study.” Please see the corrected [Table 3](#) here.

**Table 3. Expression plasmid vectors constructed by Gateway recombination method in this study.**

Plasmid	Promoter	ORF	Marker	Destination vector
pCG52	<i>S. cerevisiae</i> TEF	mCherry	LEU2	pDEST415TEFt7
pCG109	<i>S. pombe</i> ADH1	mCherry-NLS	TRP1	pDEST414TEFt7
pCG57	<i>S. cerevisiae</i> TEF	yEVenus	LEU2	pDEST415TEFt7
pRN1	<i>S. pombe</i> ADH1	yEVenus-NLS	LEU2	pDEST415TEFt7
pDHM57	<i>S. pombe</i> ADH1	yEVenus-Cln2 <sub>PEST</sub> -NLS	LEU2	pDEST415TEFt7
pCM25	<i>S. cerevisiae</i> CUP1	yEVenus-Cln2 <sub>PEST</sub> -NLS	LEU2	pDEST415TEFt7
pCG87	<i>TetO<sub>7</sub></i> -CYC1TATA	mCherry-NLS	TRP1	pDEST414TEFt7
pCG103	<i>TetO<sub>7</sub></i> -CYC1TATA	yEVenus-Cln2 <sub>PEST</sub> -NLS	TRP1	pDEST414TEFt7
pCM20	<i>TetO<sub>7</sub></i> -CYC1TATA	yEVenus-Cln2 <sub>PEST</sub> -NLS	LEU2	pDEST415TEFt7
pCG84	<i>S. pombe</i> ADH1	tTA	HIS3	pDEST413TEFt7
pCG85	<i>S. pombe</i> ADH1	rtTA	HIS3	pDEST413TEFt7
pDHM19	<i>S. pombe</i> ADH1	AIDtTA	HIS3	pDEST413TEFt7
pDHM20	<i>S. pombe</i> ADH1	AIDrtTA	HIS3	pDEST413TEFt7
pCG112* <sup>1</sup>	<i>S. cerevisiae</i> TEF	tTA	HIS3	pDEST413TEFt7
pCG113	<i>S. cerevisiae</i> TEF	rtTA	HIS3	pDEST413TEFt7
pCG106* <sup>1</sup>	<i>S. cerevisiae</i> TEF	AIDtTA	HIS3	pDEST413TEFt7
pCG107	<i>S. cerevisiae</i> TEF	AIDrtTA	HIS3	pDEST413TEFt7
pMM6* <sup>2</sup>	<i>S. cerevisiae</i> TEF	AIDrtTA	URA3, MET15	pDEST375
pCG81	<i>S. pombe</i> ADH1	OsTIR1-9Myc	URA3	pDEST416TEFt7

\*<sup>1</sup> Yeast cells with these expression vectors showed poor growth with an unknown reason.

\*<sup>2</sup> Integration vector.

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## Reference

1. Giuraniuc CV, MacPherson M, Saka Y (2013) Gateway Vectors for Efficient Artificial Gene Assembly *In Vitro* and Expression in Yeast *Saccharomyces cerevisiae*. PLoS ONE 8(5): e64419. doi:[10.1371/journal.pone.0064419](https://doi.org/10.1371/journal.pone.0064419) PMID: [23675537](https://pubmed.ncbi.nlm.nih.gov/23675537/)