

Certificate of Analysis for NR-13403

Shuttle Vector pMCSU1 for Gene Expression in *Mycobacterium tuberculosis* and *Escherichia coli*

Catalog No. NR-13403

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

Product Description: pMCSU1 is a shuttle vector that can be used for gene expression in either *Escherichia coli* or *Mycobacterium tuberculosis*. The pMCSU1 vector contains origins of replication for both organisms, *Escherichia coli* bacteriophage λ *att*R sites, a *Mycobacterium BCG* hsp60 promoter region, as well as the genes that confer resistance to kanamycin (Km), hygromycin B (Hyg) and chloramphenicol (Cm).

Lot: 59388155 Manufacturing Date: 30JUN2010

QC testing was performed by Colorado State University under the TB Vaccine Testing and Research Materials Contract (NIH). The Colorado State University documentation is attached.

ATCC[®], on behalf of BEI Resources, hereby represents and warrants that the material provided under this certificate has been subjected by the contractor to the tests and procedures specified and that the results described, along with any other data provided in this certificate, are true and accurate to the best of ATCC[®]'s knowledge.

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NR-13403_59388155_13OCT2010

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Recombinant Plasmid Quality Control Record

Plasmid designation	pMCSU1	
BEI Product Item Nu	mber <u>NR-13403</u>	
BEI Lot Number	59388155	
CSU Lot Number	10.pMCSU1.6.30	
Notebook/Pgs	notebook #5, page 23 (NKG)	
Notebook detail		
Media used	LB	
Culture size	250 mL	
Growth conditions:	Temp 37 Time 18 hrs Shaker speed 200	
Plasmid prep type (m (Cat. No. 12643)	ini/maxi, kit name or protocol) Qiagen HiSpeed Plasmid Midi Kit	
Plasmid prep detail:	Midi prep Qia100 tip and elution conditions	
Strain used to produc	e plasmidDB3.1_	
E. coli ori? Y/N	$\underline{\mathbf{Y}}$	
Contains Mycobacter	ial ori? Y/N Y	
Final concentration 5	6.95 ng/μL	
Total Stocks 43		
Buffer TE	_	
Method used for quar	ntifying nanodrop	
QC gel – N/A (no ins	ert)	
Restriction enzymes used in QC analysis N/A		
Expected size of restr	iction fragments	

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Vector	N/A			
Insert	<u>N/A</u>			
Other	N/A			
Gel description	file number, % agarose, buffer	N/A		
Recombination site/region confirmed by DNA sequencing: Y				
Note: Sequencing was performed with two primers – 1. hspF Primer sequence 5' CGGTGAGTGCTAGGTCGGACGG 3' 2. mRev1 Primer sequence 5' GACGTCAGGTGGCTAGCT 3'				
Sequence file: <u>CSU</u>	1-F/CSU1-R	Date: _7/12/10		
Plasmid Map:				
Hyg-r Ndel attR1 Cm-r Scal				

Generated by Nicholas May Date 8/19/10

Supervisor Date 8/2016

pMCSU1 7494 bp

Form 4.2.09.KMD

ccdB

_attR2 _HindIII _6his.stop _mRev1