SUPPORTING INFECTIOUS DISEASE RESEARCH

Shuttle Vector pMCSU7 for Gene Expression in *Mycobacterium tuberculosis* and *Escherichia coli*

Catalog No. NR-13408

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

Product Description: pMCSU7 is a shuttle vector that can be used for gene expression in either *Escherichia coli* or *Mycobacterium tuberculosis*. The pMCSU7 vector contains origins of replication for both organisms, *Escherichia coli* bacteriophage λ *att*R sites, a *Streptomyces coelicolor* tetracycline operator sequence, as well as the genes that confer resistance to kanamycin (Km) and chloramphenicol (Cm).

Lot: 59310226

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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Recombinant Plasmid Quality Control Record

Plasmid designation	pMCSU7	
BEI Product Item Nu	mber <u>NR-13408</u>	
BEI Lot Number	59310226	
CSU Lot Number	10.pMCSU7.6.30	
Notebook/Pgs	Notebook #5; Page 23 (NKG)	
Notebook detail		
Media used	LB	
Culture size	250 mL	
Growth conditions:	Temp <u>37</u> Time <u>18 hrs</u> Shaker speed <u>200</u>	
Plasmid prep type (m (Cat. No. 12643)	ini/maxi, kit name or protocol) <u>Qiagen HiSpeed Plasmid Midi Kit</u>	
Plasmid prep detail:	Midi prep Qia100 tip and elution conditions	
Strain used to produc	e plasmid <u>ccdB</u>	
<i>E. coli</i> ori? Y/N	<u>Y</u>	
Contains Mycobacter	ial ori? Y/N <u>Y</u>	
Final concentration	<u>107.83 ng/µL</u>	
Total Stocks <u>72</u>		
Buffer TE		
Method used for quar	ntifying nanodrop	
QC gel - N/A (no ins)	ert)	
Restriction enzymes	used in QC analysis <u>N/A</u>	
Expected size of restr	iction fragments	

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Vector	<u>N/A</u>			
Insert	<u>N/A</u>			
Other	<u>N/A</u>			
Gel description	file number, % agarose, buffer	<u>N/A</u>		
Recombination site/region confirmed by sequencing: Y				
Note: Sequencing was performed with two primers – 1 tetF Primer sequence 5' TGGCATCCGTGGCGCGGC 3' 2. mRev1 Primer sequence 5' GACGTCAGGTGGCTAGCT 3'				
Sequence files:	CSU7R CSU7F2	Date: <u>7/10/10</u> Date: <u>7/14/10</u>		
Plasmid Map:				
oriM	pMCSU7 BE314 BE315			

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Supervisor	he ke to	Date_ 8/20/10

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