

## **Centre for AIDS Reagents.**



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## **Data Sheet**

NAME:	pRHPA4259 clone 7 (SVPB14)
REPOSITORY REFERENCE:	ARP2061
	<b>NOTE:</b> This clone is also available as a member of a panel set, see <b>ARP2066</b>
PROVIDED:	20 μg plasmid DNA/per vial (0.5 mg/ml)
HOST STRAIN:	MAX Efficiency STBL2 <sup>TM</sup>
CLONING SITE:	The env/rev cassette was TA cloned into the HindIII and BamHI cloning sites of pcDNA3.1(+), in the correct orientation with the CMV promoter. The size of the insert is 2946 bp.
CLONING VECTOR:	pcDNA3.1(+). The size of the cloning vector including the insert is 8356 bp.
DESCRIPTION:	A PCR fragment containing full-length env and rev genes was derived from plasma virion-associated RNA from a subject infected with a clade B virus by reverse transcription and nested PCR amplification procedures. The env/rev cassette was ligated into pcDNA3.1(+) expression vector using the HindIII and BamHI sites. A single transformed ampicillin-resistant <i>E. coli</i> colony was selected and expanded. Sequence information is available upon request.
SPECIAL CHARACTERISTICS:	The clone represents env/rev cassette from a subject with early subtype B infection (male to female transmission). The clone expresses a functional env/rev cassette and can be used to generate pseudotyped infectious virions. pRHPA4259.7 Env containing pseudovirions are included in a standard virus neutralization panel for subtype B strains (SVPB14).

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PLASMID EXPANSION:

This plasmid can be expanded using MAX Efficiency STBL2<sup>TM</sup> Competent Cells or DH5 $\alpha$ <sup>TM</sup> Competent Cells in

LB medium at 34°C.

**GENE BANK:** 

Accession number is AY835447

STORAGE:

-80°C

SOURCE:

Drs. B. H. Hahn and Dr. J. F. Salazar-Gonzalez (Courtesy of NIH AIDS Research and Reference Reagent Programme.)

**REFERENCE:** 

Li, M., Gao F., Mascola J.R., Stamatatos L., Polonis V.R., Koutsoukos M., Voss G., Goepfert P., Gilbert P., Greene K.M., Bilska M., Kothe D.L., Salazar-Gonzalez J.F., Wei X., Decker J.M., Hahn B.H., and Montefiori D.C. Human immunodeficiency virus type 1 env clones from acute and early subtype B infections for standardized assessments of vaccine-elicited neutralizing antibodies. J. Virology 79(16): 10108-10125, 2005.

**ACKNOWLEDGEMENTS:** 

Publications should acknowledge the donor of the reagent and the Programme EVA Centre for AIDS Reagents. Suggested wording can be found on our website at http://www.nibsc.ac.uk/spotlight/aidsreagent/index.html in the "Acknowledgements" section.

Please also ensure that you send us a copy of any papers resulting from work using reagents acquired through CFAR (this can be electronically or as a paper copy)

NOTE:

Scientists at for-profit institutions or who intend commercial use of this reagent must contact William S. White, UAB Research Foundation, The Office of Intellectual Property Management, AB 1120G, 1530 3rd Ave. S, Birmingham AL 35294-0111, Tel: 205-996-2550 Fax: 205-934-5427, email: wswhite@uab.edu, before the reagent can be released.

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