

Supplementary Material



Human Cytomegalovirus Tegument Protein pUL23 Interacts with Capsid Protein pUL85

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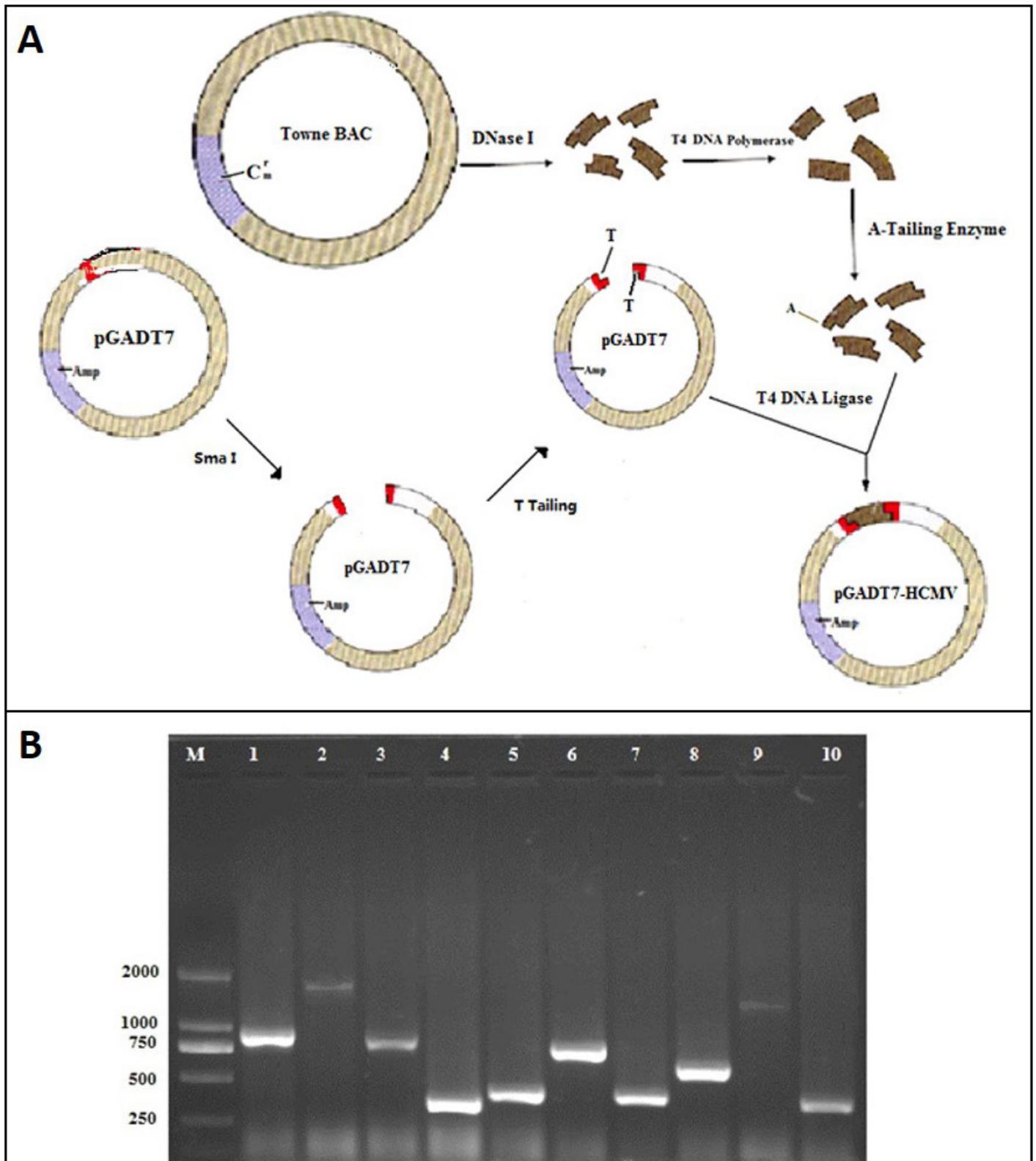
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Supplementary Table S1: Primers for cloning and sequencing

	Sequence (5'-3')
Primers for cloning UL23 gene	
UL23_Forward_EcoRI	CCG <i>GAATTC</i> CGTCGACATGTCGGTAATCAAGGACTGTTTTCTC
UL23_Reverse_XhoI	ATACTCGAGTCACGCGTCGTCAAAAAGTTGGTGGTC
UL23_Reverse_Flag_XhoI	ATACTCGAGTCACTTATCGTCGTCATCCTTGTAATCCGCGTCGTCAAAAA-GTTGGTGGTC
Primers for cloning UL85 gene	
UL85_Forward_EcoRI	CG <i>GAATTC</i> GCCACCATGGCGGCCATGGAG
UL85_Reverse_XhoI	CCGCTCGAGTCAGCCTTTAAATATGCAGGTCG
UL85_Reverse_HA_XhoI	CCGCTCGAGTCAAGCGTAATCTGGAACATCGTATGGGTAGCCTTTAAATATG-CAGG
UL85_N_Reverse_XhoI	CCGCTCGAGCAGCACGCATAGCGCGTC
UL85_M_Forward_EcoRI	CG <i>GAATTC</i> ATGAAGGGCGACGCGCTATGC
UL85_M_Reverse_XhoI	CCGCTCGAGGAGGCAGGCGGTCTTCATGT
UL85_C_Forward_EcoRI	CG <i>GAATTC</i> ATGGACATGAAGACCGCCTGCCTC
Primers for cloning UL46 gene	
UL46_Forward_EcoRI	CGGGTACCTCTACCGTGCCTCGCC
UL46_Reverse_Myc_XhoI	CGCGGATCCAGGTCTTCTTCAGAGATCAGTTTCTGTTTCGACGAAT-TCTCGAAAGTCTCCCA

Restriction enzyme sites used for cloning are marked as *Italic*, Expression tag or "ATG" initiation codon are marked as underline.



Supplementary Figure S1: Construction of HCMV genomic library.

A) Flowchart of the construction of HCMV genomic library; **B)** PCR analysis the inserts sizes of HCMV library. DNA marker (M), PCR amplification of inserted fragments from 10 random recombinant plasmids (1-10)