

DNA Modified Hydrogel for Simultaneous Purification, Concentration and Detection of Targeted cfDNA in Human Serum

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Table S1. DNA sequences used in this work

Oligonucleotide		Sequence (5-3)
20bp-acrydite		Acrydite-gaggagccgaaaaggttcca
20bp+0bp		tggaaccttttcggctcctc
20bp+10bp		tggaaccttttcggctcctc tgccgatcca
20bp+20bp		tggaaccttttcggctcctc tgccgatccatactgcggaa
20bp+30bp		tggaaccttttcggctcctc tgccgatccatactgcggaatgccgatcca
20bp+40bp		tggaaccttttcggctcctc tgccgatccatactgcggaatgccgatccatactg cgga
20bp+50bp		tggaaccttttcggctcctc tgccgatccatactgcggaatgccgatccatactg cggagttcgaactg

Table S2. The primers used to amplify different lengths of dsDNAD in this work

The		Sequence (5-3)
20bp		tggaaccttttcggctcctc
100bp		aagcataaagtgtaaagcct
150bp		taattgcgttgcgctcactg

200bp		cagctgcattaatgaatcgg
250bp		tgggcgctcttccgcttct
300bp		ggctgcggcgagcggatca

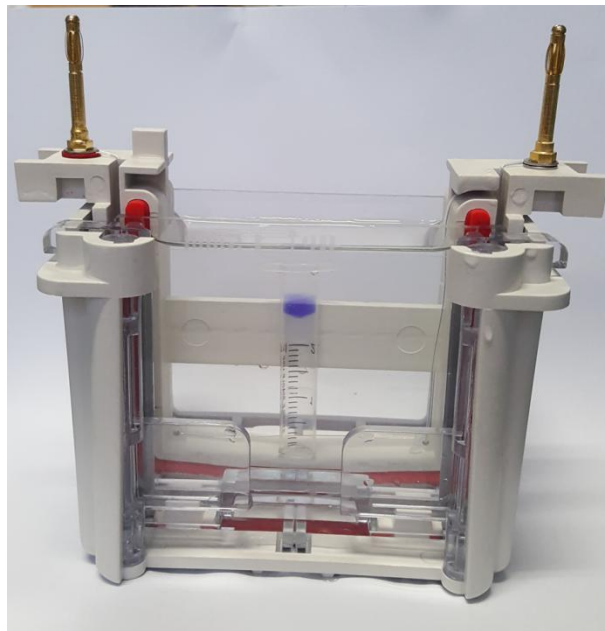


Figure S1. The electrophoresis apparatus of the DNA-Gel

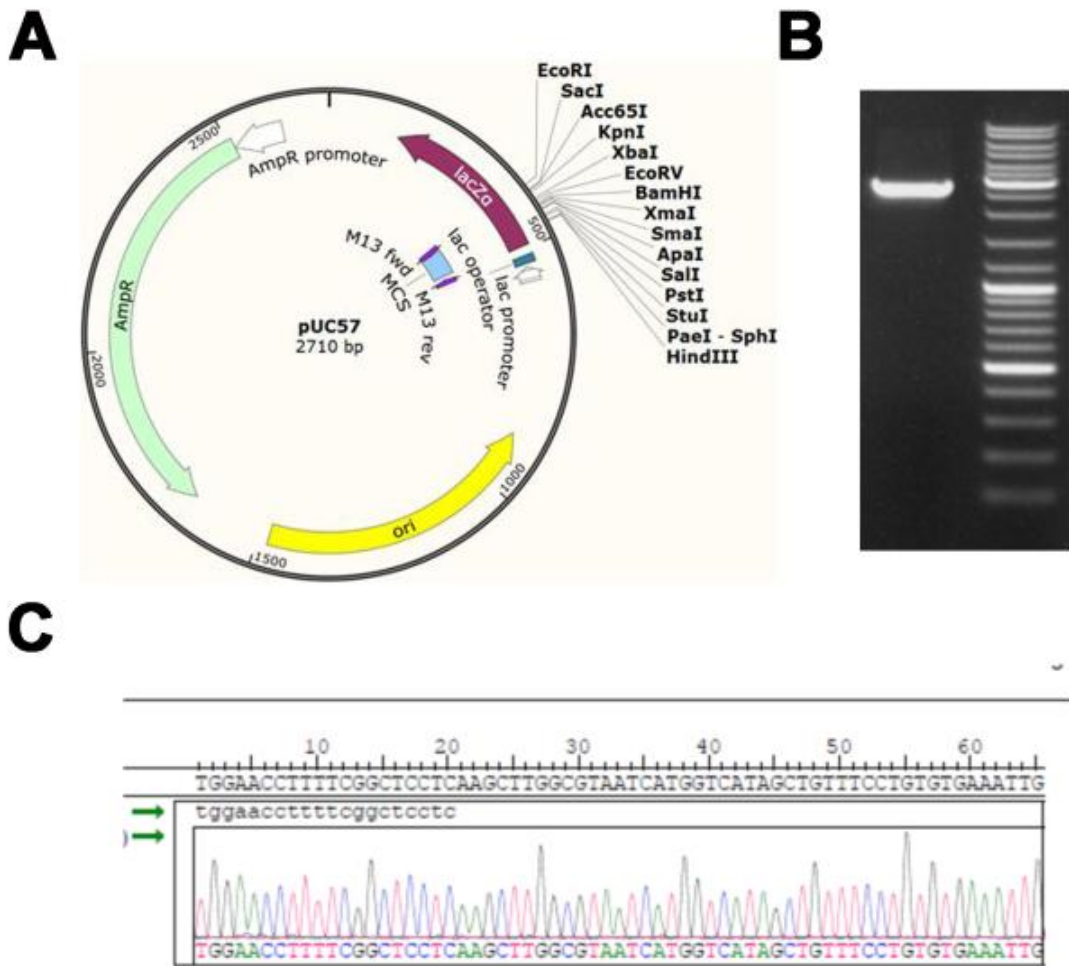


Figure S2. (A) the EcoRI/HindIII of pUC57; (B) the the EcoRI/HindIII of pUC57 ; (C) the sequencing analysis of pUC57

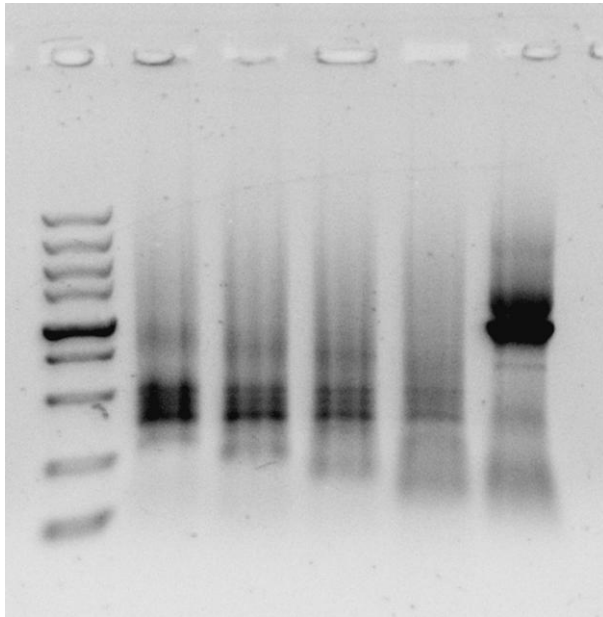


Figure S3. Different lengths of dsDNAD amplified by PCR; the lanes were maker, 100bp, 150bp, 200bp, 250bp and 300bp, respectively. The DNA maker band from the bottom up were 100, 250, 500, 750, 1000, 1500, 2000, 3000 and 5000 bp, respectively.