Supporting Information

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In vivo Cloning of PCR Product via Site-Specific Recombination in Escherichia coli

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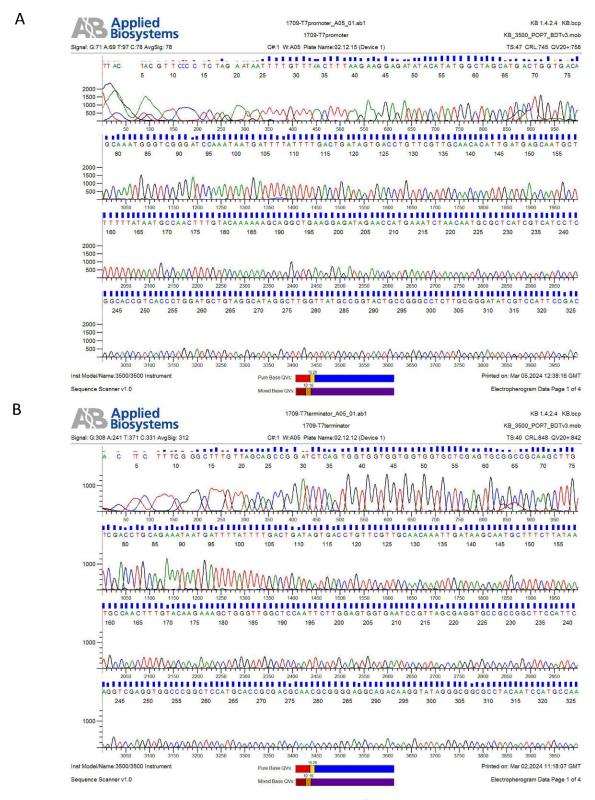


Fig S1. Sequencing chromatogram of the inserted fragment. Related to Fig 5 (A) The performed sequencing with universal primers of the T7 promoter and (B) T7 terminator.

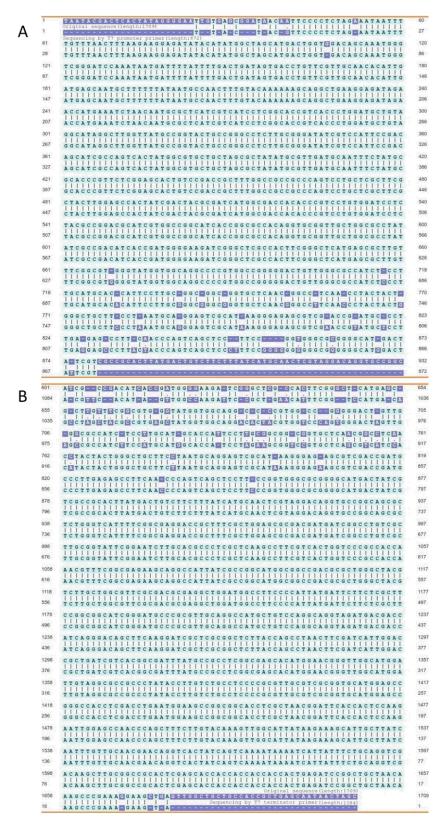


Fig S2. The alignment of sequencing data with the sequence of the *tetA* gene (flanked by *attL* sites). (A) The correct direction of the insert fragment was confirmed by alignment with sequencing data related to the T7 promoter and (B) T7 terminator primers.