Markerless gene deletion of Mannheimia succiniciproducens using a New Temperature Sensitive Plasmid

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Temperature-sensitive plasmid of Mannheimia succiniciproducens MBEL55E was generated from the chemical mutagenesis of plasmid pMVSCS1, which is a native plasmid obtained from Mannheimia varigena. The Ts plasmid in M. succiniciproducens was fully functional at 30°C, but failed to replicate above 42°C. The temperature sensitive plasmid was further modified to generate markerless mutants combined with Cre/loxP system. This result confers that multiple gene deletion in M. succiniciproducens can be applied to generate an enhanced succinic acid producing strain. [This work was supported by the Korea Science and Engineering Foundation (KOSEF) grant funded by the Korea government (MOST) (No. 2005–01294). Further supports by the LG Chem Chair Professorship, IBM SUR program, and by the KOSEF through the Center for Ultramicrochemical Process Systems are appreciated].