

Metabolic engineering of *Escherichia coli* for the production of polyhydroxyalkanoates incorporating 2-hydroxybutyrate

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E. coli strain was metabolically engineered to synthesize polyhydroxyalkanoates (PHAs) containing 2-hydroxybutyrate (2HB) monomer from glucose. The recombinant *E. coli* expressing evolved *Clostridium propionicum* propionyl-CoA transferase (PctCp) and *Pseudomonas* sp. MBEL 6-19 PHA synthase 1 (PhaC1Ps6-19) was developed and cultured in a chemically defined medium containing 20 g/L of glucose and varying concentrations of 2HB and 3HB. PHAs consisting of 2HB, 3HB, and a small fraction of lactate were synthesized. Also, heterologous metabolic pathway to supply 2-hydroxybutyrate from glucose was constructed via the citramalate pathway. Recombinant *E. coli* expressing the *phaC1437*, *pct540*, *cimA3.7*, and *leuBCD* genes together with the *L. lactis* II1403 *panE* gene successfully produced PHAs consisting of 2HB, 3HB, and a small fraction of lactate by varying the 3HB concentration in the culture medium. [This work was supported by the Korean Systems Biology Research Project (20090065571) of the Ministry of Education, Science and Technology (MEST), the R&D Program of MKE/KEIT (10032001) and KRICT]