

Limonene Production from Acetate and Ethanol to Enhance Carbon Flow into the Mevalonate Pathway using Recombinant *Saccharomyces cerevisiae*양정모<sup>1</sup>, 서교연<sup>1</sup>, 조숙형<sup>2</sup>, 나정걸<sup>1</sup>, 이진원<sup>1,2,†</sup><sup>1</sup>서강대학교 화공생명공학과; <sup>2</sup>서강대학교 C1 가스 리파이너리 사업단  
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Biological production of limonene using microorganisms has been actively studied because it can replace hazardous organic solvents in chemical industry. In this study, we developed a bioconversion process from acetate and ethanol to limonene using *Saccharomyces cerevisiae*. Acetate and ethanol can be directly converted to acetyl-CoA not pyruvate providing, which has the advantage of solving the problems of carbon loss. To produce limonene from acetate and ethanol, we introduced limonene synthase(LS) gene from *Mentha spicata* which converts geranyl pyrophosphate(GPP) or neryl pyrophosphate(NPP) to limonene with GPP synthase(GPPS) from *Abies grandis* or NPP synthase(NDPS) from *Solanum lycopersicum* to *S. cerevisiae* using modified pESC-TRP vector. To improve the production of limonene, we optimized the mevalonate pathway by over-expression of acetyl-CoA thiolase, 3-hydroxy-3-methyl glutaryl-CoA(HMG-CoA) reductase, mevalonate kinase and IPP delta-isomerase which are the main bottle-neck of GPP synthesis using modified pESC-LEU and pESC-HIS vector.