Improvement of H₂ production by recombinant Escherichia coli strains

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As other microbial production process, the performance of biocatalyst often determines the success of dark fermentative hydrogen ($\rm H_2$) production. However, it is not simple to determine which microorganism is suitable for $\rm H_2$ production process. Here, we compared 4 different Enterobacteriacea (*Enterobacter aerogenes, Citrobacter amalonaticus* Y19, *Escherichia coli* K-12 and *Escherichia coli* DKJ135) for its capability of $\rm H_2$ production. The comparison showed that $\rm H_2$ production activities of microorganisms were not much different except of wild type *E. coli* K-12 from format and modification of gene expression of *fhlA* (encoding activator of formate-hydrogen lyase(FHL)) could affect to improvement of $\rm H_2$ production activity. Therefore, by modifying the expression of relevant genes including *fhlA*, improvement of H2 production activities was attempted. $\rm H_2$ production activity was significantly increased more than 3-fold by deleting both *hycA* (encoding repressor of FHL) and *iscR* (*iron-sulfur duster regulator*) and overexpressing *fhlA*. In addition, we confirmed the high volumetric $\rm H_2$ production activity (2.4 L/L/h) from formate using mutant strain of *E. coli* SH5.