

Biological hydrogen production by *Escherichia coli*

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Biological hydrogen production research has been carried out as a means of solving energy and environmental problems concurrently for over quarter century. Among the various approaches, biological hydrogen production by dark-fermentation has been rapidly researched in recent times¹⁾. Dark-fermentation is a process of hydrogen and organic acids production through fermentation a substrate like glucose in anaerobic and dark condition by some anaerobic bacteria. Above all, the most important thing in the dark-fermentation is to maintain anaerobic condition. This problem can be solved by sparging N₂ gas to remove O₂ gas in the reactor. We performed study about hydrogen production mechanism, separation and analysis method.

There are many fermentative bacteria, but widely used and well-known coli-form, *Escherichia coli* is suitable for this experiment. The fermentation products of *Escherichia coli* comprise a mixture of ethanol, acetate, formate, lactate and succinate¹⁾. Especially, we concentrated on formate. Formate is anaerobically decomposed to molecular hydrogen and carbon dioxide by *Escherichia coli* according to the equation : $\text{HCOOH} \rightarrow \text{CO}_2 + \text{H}_2$. This decomposition is done by a multi-enzyme system(formate hydrogenlyase, FHL) containing formate dehydrogenase(FDH) and hydrogenase. Studies identified an absolute requirement of selenium and molybdenium for the synthesis of active formate dehydrogenase(FDH) and hydrogen gas production by *Escherichia coli*^{2,3)}. And fermentative hydrogen production can be maximizing through an active hydrogenase.⁴⁾ This study was aimed at investigating the effect of Mo, Se and Fe on the activity of FDH and Hydrogen production rate for dark fermentation.

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