6–6 Energy Saving of the Main Reaction in the IS Process

- Membrane and Electrode Development for the Bunsen Reaction Overvoltage Reduction -



Fig.6-14 Schematic of the Bunsen reaction using the ion-exchange membrane

Hydriodic and sulfuric acids were produced on the cathode and the anode, respectively, in an electric cell equipped with an ion-exchange membrane.



Fig.6-15 Microscopic outlook of the porous Au electrode

The porous Au surface increases the effective reaction area and allows the anode reaction to easily proceed. Consequently, the overvoltage that originated from the electrode reaction can be reduced.

The iodine–sulfur (IS) water-splitting H_2 production process as the heat application of high temperature gascooled reactors is investigated to realize the hydrogen (H_2) economy. The IS process is composed of sulfur and iodine chemical reactions.

The Bunsen reaction, which is a starting reaction of the IS process, produces sulfuric and hydriodic acids by mixing iodine and sulfur dioxide with water. The membrane for the Bunsen reaction method is investigated to allow the Bunsen reaction to proceed using an electric cell equipped with an ion-exchange membrane (Fig.6-14). The conventional Bunsen reactor required the separation process of two acid solutions after a reaction. In contrast, no solution separation process is needed because the two acids can be separately produced in a cell partitioned into two channels by the membrane. However, in the membrane for the Bunsen reaction assembled with the usual electrodes and membrane, the overvoltage was excessively high for practical application to the IS process. The overvoltage originated from the anode reaction at the anode electrode and the membrane resistance caused by the proton permeation. Therefore, the novel electrode and the ion-exchange membrane were developed to reduce the overvoltage.



Fig.6-16 Current density dependence of the Bunsen reaction overvoltage



First, the porous Au electrode was developed in collaboration with Shibaura Institute of Technology (Fig.6-15). The porous electrode that enhanced the effective surface area allowed the anode reaction to easily proceed, indicating that the overvoltage was reduced.

Second, the ion-exchange membrane, which introduced many ion-exchange groups compared with the usual membrane to improve the conductivity, was prepared through the National Institutes for Quantum Science and Technology technique. The prepared membrane enabled the reduction of the membrane resistance.

Finally, the membrane for the Bunsen reaction was operated using the electric cell assembled with the developed electrode and membrane. The stoichiometric Bunsen reaction was confirmed to proceed. Moreover, the overvoltage was 0.21 V at 200 mA/cm², which is 1/3 lower than that of the usual setup (Fig.6-16). In summary, the practical process efficiency for the technical applicability of the membrane for the Bunsen reaction can be achieved.

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Reference

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