Fabrication of Photocatalytic Solar Cell using Manganese Oxide Solid Electrolyte.

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INTRODUCTION: Metal oxide precursor solution has been made by advanced sol-gel method, in which hydrophilic and hydrophobic mixed solvents were used for hydrolysis and polymerization of alkoxide. Photocatalytic activity of TiO₂ thin film prepared by this method is greater than that of conventional sol-gel method. More efficient photocatalytic activity would be expected for TiO₂-SnO₂ thin film. In this work, we fabricated a photocatalytic solar cell, a kind of capacitor, using manganese oxide solid electrolyte and high photocatalytic TiO₂-SnO₂ thin film. Manganese oxide electrolyte was prepared by advanced sol-gel method. The cell voltage generated by irradiation of black light was measured.

EXPERIMENTAL: SnO₂ precursor solution was prepared as described below. By refluxing tin chloride anhydrate in butanol and toluene solvent, tin alkoxide solution was formed and then concentrated. Toluene and butanol containing distilled water was added in condensed solution to produce tin oxide precursor solution. Manganese oxide solution was prepared by the same preparation process of SnO₂ precursor solution except that the starting material was manganese chloride tetrahydrate and the hydrophilic solvent was ethanol. To prepare the TiO₂-SnO₂ precursor solution, SnO₂ precursor solution was added to TiO₂ precursor solution. TiO₂ precursor solution was made by our method.

 TiO_2 -SnO₂ thin film was made by spincoating on FTO substrate and heat-treatment at 500 °C. Manganese oxide precursor solution was spincoated on FTO substrate, and heat-treated at various temperatures. Schematic view of photocatalytic solar cell is shown in Fig.1. The TiO₂-SnO₂ thin film was suited to the manganese oxide film. The cell voltage was measured under the UV-light irradiation of wavelength 352 nm. The crystal structure of these oxides was measured by XRD. The surface morphologies of these were observed by FE-SEM and AFM.

RESULTS AND CONCLUSIONS: Fig.2 shows XRD patterns of manganese oxide after heat-treatment at 200~500°C. At 350°C the Mn₃O₄ phase appeared. At 500° C the peaks showed the two phases of Mn₃O₄ and Mn₂O₃ structure. Fig.3 shows the cell voltage under UV irradiation when manganese oxide was used as solid electrolyte, heat-treated at 200~500°C. After irradiation of UV-light for about 20 second, a steady state was obtained. The cell voltage using solid electrolyte heat-treated at 350°C was about 0.75 V. This voltage is considered relatively high for a cell made under another condition. When solid electrolyte was heat-treated at a lower or higher temperature than 350°C, cell voltages did not exceed about 0.7 V. The highest voltage is obtained at Mn₃O₄ phase. It is considered that the manganese electrolyte takes part in oxidation-reduction reaction with hole and electron, these species produced by photocatalytic excitation on TiO2-SnO2 film. These mechanisms occurred at the interface between manganese oxide and TiO₂-SnO₂, as shown in Fig.1.



Fig.1. The schematic view of photocatalytic solar cell.



Fig.2. X-ray diffraction patterns of manganese oxide at various temperatures: 200, 250, 350 and 500°C.



Fig.3. Measurement of voltage under the UV-light.