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Electrochemical Property of Fe₂O₃-Cr₂O₃ Thin Films Formed by LP-MOCVD

LP-MOCVD법으로 합성한 Fe₂O₃-Cr₂O₃계 박막의 전기화학적 특성

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The purpose of this study was to examine the relationship between the corrosion resistance and the crystallinity of the Fe₂O₃-Cr₂O₃ artificial passive films and to give some insight on the relationship on the real passive films.

The Fe₂O₃-Cr₂O₃ films were formed on Pt substrates by low pressure MOCVD technique using iron (III) acetylacetonate and chromium (III) acetylacetonate as precursors at substrate temperatures of 150-350°C. The thickness and the optical constants of the films were determined by a rotating analyzer type automatic ellipsometer. The chemical composition of the films was determined by inductively coupled plasma-emission spectroscopy (ICPS) analysis. The oxidation state of constituent elements of the films was examined by X-ray photoelectron spectroscopy (XPS). Micro- and crystal structures were examined by transmission electron microscope (TEM) and electron beam diffraction (ED), respectively. The films coated on Pt plates were immersed in 1.0 M HCl for a given time and the decrease in film thickness was measured by ellipsometry. Potentiodynamic polarization curves were measured in 1.0 M H₂SO₄ and 0.5 M HCl by a potentiostat.

The films deposited above 300°C had crystalline structures and those, deposited below 250°C had amorphous structures. The films deposited above 250°C had a high amount of M-O type chemical bonds, and those deposited below 200°C had high amount of M-OH type chemical bonds. The films deposited above 300°C hardly dissolved in 1.0 M HCl and those deposited below 250°C, however, easily dissolved in the solution. The dissolution rate of the films in the solution increased with decreasing substrate temperature. Passive and transpassive current densities of the films in 1.0 M $\rm H_2SO_4$ were dependent on the substrate temperature and increased with decreasing the temperature. When polarized cathodically in 1.0 M $\rm H_2SO_4$ and 0.5 M HCl, the films deposited below 250°C dissolved due to the reduction of Fe₂O₃ component in the films. The reduction of Fe₂O₃ component was, however, suppressed on the films deposited above 300°C. Therefore, with increasing crystallinity and amount of M-O type chemical bonds, the corrosion resistance of the films increases in HCl and $\rm H_2SO_4$ solutions.