Direct Observation of Active Nickel Oxide Cluster in Nickel–Borate Electrocatalyst for Water Oxidation by In Situ O K-Edge X-ray Absorption Spectroscopy

Masaaki Yoshida,†‡ Yosuke Mitsutomi,† Takehiro Mineo,† Masanari Nagasaka,§ Hayato Yuzawa,§ Nobuhiro Kosugi,§ and Hiroshi Kondoh†

†Department of Chemistry, Keio University, 3-14-1 Hiyoshi, Kohoku-ku, Yokohama 223-8522, Japan
‡Cooperative Research Fellow, Catalysis Research Center, Hokkaido University, Kita21, Nishi10, Kita-ku, Sapporo, Hokkaido 001-0021, Japan
§Institute for Molecular Science, 38 Nishigo-Naka, Myodaiji, Okazaki 444-8585, Japan

ABSTRACT: In situ O K-edge X-ray absorption fine structure (XAFS) spectroscopy was applied to investigate the electronic and structural change in the nickel–borate (Ni–B) electrocatalyst during the oxygen evolution reaction (OER). An absorption peak was observed around 528.7 eV at 1.0 V versus Ag/AgCl in a potassium borate aqueous solution, which relates with the formation of nanoscale order domains of edge-sharing NiO₆ octahedra in the Ni–B electrocatalyst. XAFS spectra were measured with variation of the electrode potential from 0.3 up to 1.0 V. The measured absorption peaks suggest that the quantity of NiO₆ octahedra increased in correlation with the OER current; however, when the potential was changed downward, the XAFS absorption peak assigned to NiO₆ octahedra remained constant, even at the electrode potential for no OER current. This difference implies that the water oxidation catalysis proceeds at the domain edge of NiO₆ octahedra. The XAFS technique provides the first successful direct probing of the active species in the Ni–B electrocatalyst during electrochemical reaction.

1. INTRODUCTION

Water splitting using solar energy is a promising approach to produce renewable hydrogen gas toward a sustainable society.1−7 Various types of photoelectrodes and photovoltaics for water splitting have been developed with high solar-to-hydrogen efficiencies of over several percent.8−12 Recently, a perovskite photovoltaic attained a solar-to-hydrogen efficiency of 12.3% using hydrogen and oxygen evolution electrocatalysts.8 For the realization of efficient photovoltaic and photoelectrochemical systems for water splitting, oxygen evolution electrocatalysts have often been used to catalyze water oxidation due to their lower activation energies. Therefore, the development of efficient catalysts for the oxygen evolution reaction (OER) has attracted considerable attention due to the potential toward highly efficient energy conversion.

Nocera and colleagues reported that nickel–borate electrodeposited from a dilute Ni²⁺ solution in a borate-buffered electrolyte (Ni–B) functions as an efficient OER electrocatalyst.13−15 The Ni–B OER catalyst is unique because it is composed of inexpensive and earth-abundant materials, and it operates safely with high activity.13−15 As similar materials, cobalt–phosphate (Co–P) and cobalt–borate (Co–B) complexes have also been reported as efficient electrocatalysts for OER in phosphate and borate electrolytes, respectively.16−19 These OER electrocatalysts can enhance the activities for water splitting, when the electrocatalysts are deposited on a semiconductor photoelectrode or photovoltaic surfaces such as Si, Fe₂O₃, and BiVO₄.9,10,20−23

Mechanistic studies that employ various techniques have been aimed toward determining the origin of efficient oxygen evolution for Ni–B electrocatalysts or similar materials (Co–P, Co–B).24−35 An electron paramagnetic resonance (EPR) spectroscopy study revealed the presence of Co⁴⁺ species in a Co–P electrocatalyst in a frozen electrolysis solution.35 In addition, Ni K-edge X-ray absorption fine structure (XAFS) analysis demonstrated that the average valence state of the nickel species in a Ni–B electrocatalyst changes from +2.0 to +3.6 in response to an applied positive voltage.24 It was concluded that oxidation of the nickel species is due to the generation of bis-oxo/hydroxo-bridged nickel centers organized into sheets of edge-sharing NiO₆ octahedra within ordered domains 2 to 3 nm in diameter by extended X-ray absorption fine structure (EXAFS) analysis.24 The nanoscale order structures of the Co–P and Co–B electrocatalysts were...
investigated by X-ray pair distribution function (PDF) analysis, and a combination study using infrared spectroscopy (IR), atomic force microscopy (AFM), and X-ray grazing incidence diffraction (GID), from which it was determined that nanoscale CoO6 clusters can assemble into stacked superstructures with highly porous morphology. It is suggested that the porous electrocatalyst film enables proton-coupled electron hopping to proceed in the electrocatalyst, which can increase the OER sites with growth of the film thickness. Therefore, the unique characteristics of these electrocatalysts are currently of particular interest as a material for efficient OER.

XAFS spectroscopy is a powerful tool to investigate the local electronic states for OER catalysts. The in situ Ni K-edge XAFS technique using hard X-rays was applied to the Ni−B electrocatalyst; however, it is difficult to quantitatively discuss the active species in the OER electrode potential with this technique because the Ni K-edge XAFS spectra become a superposition of each spectrum for Ni2+ and Ni3.6+ species in the Ni−B electrocatalyst and Ni2+ cations in the electrolyte solution. On the contrary, O K-edge XAFS spectroscopy using soft X-rays has been useful to directly observe the oxygen species in transition-metal oxides at the liquid/solid interface. Therefore, in this study, the Ni−B electrocatalyst was investigated using in situ O K-edge XAFS measurements with control of the potential. An attempt was made to directly detect the active oxygen species in the Ni−B electrocatalyst and discuss the relationship between the amount of active species and the OER activity.

2. EXPERIMENTAL SECTION

2.1. Sample Preparation. Au thin films (20 nm) were deposited on Si substrates (10 × 10 mm2) with an evaporated Cr layer (5 nm thick) and SiC (150 nm thick, 2 × 2 mm2, NTT AT) or SiN4 (500 nm thickness, 2 × 2 mm2, CORNES Technologies) windows (Au/Cr/SiC or Au/Cr/SiN4, respectively) for use as working electrodes for electrochemical reactions. An in-house-built electrochemical cell was used with a Pt mesh counter electrode and a Ag/AgCl (saturated KCl) reference electrode. The potential was controlled using a potentiostat. A Ni−B electrocatalyst was deposited on the Au thin film electrodes in 0.1 M K2B4O7·4H2O (potassium borate, K−B, 99.0%, Showa Chemical) aqueous solution containing 0.4 mM Ni(NO3)2·(99.9%, Wako Pure Chemical Industries) with bubbling He, according to a previous report.24 Milli-Q water (total organic carbon <5 ppb, resistivity >18 MΩ-cm) was used for all experiments. The electrochemical activity of the Ni−B electrocatalyst for the OER was tested in K−B, aqueous solution with bubbling He by measurement of the current–voltage characteristics. The surface morphology, chemical state, and film thickness of each prepared sample were observed using scanning electron microscopy (SEM; FEI, Inspect S50), energy-dispersive X-ray spectroscopy (EDX; EDAX, Genesis), and confocal laser scanning microscopy (CLSM; Keyence, VK-9710), respectively.

2.2. In situ O K-Edge XAFS. Electrochemical soft X-ray XAFS measurements were performed with the transmission mode at the BL3U beamline of the UVSOR Synchrotron, in accordance with previous work.40,41 The radiation was monochromatized with a varied-line-spacing plane-grating monochromator (VLS-PGM). The intensities of the incident X-rays were monitored using a gold mesh under ultrahigh vacuum in front of the sample (I0) and a photodiode (IRD AXUV100) under a helium atmosphere after the sample (I1). The XAFS data were collected as transmission spectra with the Lambert–Beer formula ln(I0/I1). The photon energy was calibrated according to the O 1s−π* peak (530.80 eV)42 for O2 gas mixed in a buffer gas with He gas. The O K-edge XAFS spectrum of Milli-Q water was measured as a reference sample for sample sandwiched between two SiC membranes (100 nm thickness, 2 × 2 mm2, NTT AT). On the contrary, the O K-edge XAFS spectra for the Ni(OH)2, NiO, β−NiOOH, and K−B reference samples were obtained by measuring the sample drain current under ultrahigh vacuum. For the investigation of electrochemical reactions, we sandwiched a liquid layer of a several hundreds of nanometers between the SiC and Au/Cr/SiC membranes by setting Teflon and Au spacers (100 μm thickness) to the window frames of the membranes in an in-house-built electrochemical XAFS cell with a Pt mesh counter and Ag/AgCl (saturated KCl) reference electrodes (Figure 1).
The electrolyte thickness was adjusted by changing the outer helium gas pressure of the SiC and Au/Cr/SiC membranes, according to the previous works. The potential of the Au/Cr/SiC working electrode was controlled using a potentiostat connected to the Au spacer with a conducting wire. The Ni–Bi electrocatalyst was deposited on the Au thin film electrode in 0.1 M K–Bi aqueous solution containing 0.4 mM Ni(NO₃)₂, with bubbling He. The Ni–Bi electrocatalyst was then anodized at 1.0 V for 2 h in 0.5 M K–B, aqueous solution with bubbling He by exchanging the solutions with a tubing pump system. In situ O K-edge XAFS spectra were measured for the Ni–Bi electrocatalyst before and after anodization while maintaining the electrode potential at 1.0 V. The potential dependence of the XAFS spectra was tested in 0.1 M K–Bi aqueous solution with bubbling He by exchanging solutions with the tubing pumping system during current monitoring for the OER with the potentiostat. The XAFS spectra were taken with scanning time of 20 min at cumulated number of 3 times, which results in each electrode potential being applied for 1 h.

3. RESULTS AND DISCUSSION

3.1. Electrochemical Properties and Surface Morphologies. The current during electrodeposition of the Ni–Bi electrocatalyst increased for 2 h, as shown in Figure S1a, which indicates growth of the Ni–Bi thin film and the evolution of oxygen gas not only at the Ni–Bi surface but also from the Ni–Bi substrate, as previously reported. The anodization reaction of the Ni–Bi electrocatalyst was monitored, as shown in Figure S1b, and the current continued to increase for 2 h as in the previous work. Figure 2a shows the electrochemical reaction of Ni–Bi thin film at 1.0 V versus Ag/AgCl in a 0.1 M K–Bi aqueous solution containing 0.4 mM Ni(NO₃)₂, together with those for H₂O, β-NiOOH, NiO, Ni(OH)₂, K–B, and O₂ gas as references that relate with the Ni–Bi catalysis. A peak associated with an oxygen species was observed at ca. 528.7 eV, which was similar to that in the β-NiOOH reference. To evaluate the electrode potential dependence of the Ni–Bi electrocatalyst, we changed the electrolyte solution to 0.5 M K–Bi aqueous solution without nickel ions, and we conducted XAFS measurements while changing the applied electrode potential. The peak at ca. 528.7 eV disappeared at the lower potential (0.5 V) and reappeared at the higher potential (1.0 V) accompanied by the OER activity, as shown in Figure 4a. The same behavior was observed even in 0.05 M K–Bi aqueous solution (Figure 4b). Previous Ni K-edge XAFS measurements suggested that the Ni–Bi electrocatalyst before anodization forms μ-oxo/hydroxido nickel centers organized into higher order domains of edge-sharing NiO₆ octahedra with β-NiOOH structure at higher potential. Thus, in the present study, the formation of NiO₆ octahedra domains can be monitored by the direct observation of oxygen species in the Ni–Bi thin film. The NiO₆ absorption was observed as a single peak at 528.7 eV, which originates from transition to hybridized Ni(3d)–O(2p) states with eg orbital symmetry and the low-spin d⁷ (Ni³⁺) electron configuration. It should be noted that the small peak around 530.8 eV in Ni–Bi is attributed to the strong 1s–π* resonance absorption of residual oxygen gas. No detection of the Ni(OH)₂ peak at lower potential in the O K-edge spectra may be mostly because the electronic state of oxygen species of Ni(OH)₂ structure in the Ni–Bi electrocatalyst is not uniform due to hydration or protonation on the Ni–Bi film.

Figure 3. (a) In situ O K-edge XAFS spectrum for Ni–Bi electrocatalyst under electrochemical control at 1.0 V in 0.1 M K–Bi aqueous solution containing 0.40 mM Ni(NO₃)₂. XAFS spectra for H₂O, β-NiOOH, NiO, Ni(OH)₂, K–B, and O₂ gas are shown for reference. (b) In situ O K-edge XAFS spectra (i) before and (ii) after anodization for 2 h at 1.0 V in 0.5 M K–Bi aqueous solution.
Anodization of the Ni–Bi electrocatalyst caused the NiO$_6$ octahedra peak to increase, as shown in Figure 3b. Bediako et al. reported that the inactive Ni–Bi species with β-NiOOH structure (Ni$^{3+}$) changed to the active Ni–Bi species with γ-NiOOH structure (Ni$^{3.6+}$), which resulted in the relaxation of Jahn–Teller distortion with nonequivalent Ni–O and Ni–Ni distances.24 We also verified the existence of γ-NiOOH structure for our Ni–Bi sample by in situ Ni K-edge XAFS measurements as well as the previous work,24 as shown in Figure S3. Both NiOOH structures have two kinds of oxygen species derived from hydroxide (OH) and oxide (O) at a different OH/O ratio (e.g., β-NiOOH = Ni$^{3+}$(O)$_{1.6}$(OH)$_{0.4}$, γ–NiOOH = Ni$^{3.6+}$(O)$_{1.6}$(OH)$_{0.4}$). It is known that the 1s core level of oxygen species of hydroxide (OH) is shifted to higher binding energy with respect to that of oxygen species of oxide (O) in transition-metal oxyhydroxides, which results in the oxide oxygen peak being observed at a lower energy than the hydroxide oxygen peak in the O K-edge XAFS spectra.43 Thus, in the present work, the first peaks in the O K-edge XAFS spectra are likely attributed to the oxide oxygen species in the β- and γ-NiOOH structures, and the increase of the XAFS peak is most likely due to an increase in the oxide oxygen species by the structural change from β-NiOOH (Ni$^{3+}$) to γ-NiOOH (Ni$^{3.6+}$). On the contrary, the first peak positions in the O K-edge XAFS spectra for transition-metal oxides are empirically related with the cation valences to some extent.44,45 Therefore, the local nickel valence bound by oxide oxygen species in the Ni–Bi electrocatalyst was tentatively estimated as ca. 4+ using the linear fitting of first peak positions of reference compounds against the cation valences, as shown in Figure S4, which suggests that the highly oxidized Ni species such as NiO$_2$ are formed during catalytic water oxidation. It should be noted that the O K-edge XAFS spectra are sensitive to chemical bonding (molecular orbitals), valence, spin state, and so on.38,39,43–46 Further investigation is required to evaluate the cation valence quantitatively.

### 3.3. Electrode Potential Dependence of XAFS Spectra and Reaction Model

The upward potential dependence of the Ni–Bi electrocatalyst on the Au thin film was evaluated. Figure 5a shows O K-edge XAFS spectra for this sample measured under various controlled potentials. As the potential was changed from 0.3 to 1.0 V, the intensity of the peak for the NiO$_6$ octahedra increased from 0.6 V and remained constant over 0.8 V, which indicates that the nickel species was oxidized from Ni$^{2+}$ species with the Ni(OH)$_2$ structure to NiO$_6$ octahedra on the basis of previous works.24,36 The increase in...
the edge of XAFS peak area for NiO$_6$ octahedra was consistent with the onset of OER activity (Figure 5b), which indicates that the high activity of the Ni–Bi thin film for the OER is derived from the formation of NiO$_6$ octahedra domains. On the contrary, the XAFS peak remains constant at higher electrode potentials (≥0.8 V) because the nickel oxide nanoclusters were completely oxidized to NiO$_6$ octahedra. The increase in the OER current even in the suppression region of peak growing in XAFS spectra suggests that the reaction rate for OER catalysis at higher potentials is probably restricted according to the Tafel equation, relating the rate of an electrochemical reaction to the overpotential as $V = a + b \log I$, where $V$ is the potential (overpotential), $I$ is the current density, and $a$ and $b$ are the Tafel equation constants.\(^{47}\)

To compare the difference according to the direction of potential sweep, we measured the downward potential dependence of XAFS spectra under potential control (Figure 6a). The NiO$_6$ XAFS peak at 528.7 eV was observed at higher potential as well as the result of upward potential dependence; however, the peak was observed even at 0.6 V, which indicates that the generated NiO$_6$ octahedra were not easily reduced to the Ni$^{2+}$ species. The relationship between the electrode potential and XAFS peak area suggests that the downward potential dependence is different from that of the upward potential dependence, as shown in Figure 6b. The result indicates that the nickel oxide nanocluster was gradually reduced from the domain edge because the oxidation and reduction of domain edge is likely to proceed by lower activation energy than those of domain inside; however, the currents for OER activity were the same for the upward and downward sweeps (Figure 6c), which implies that the interior of the NiO$_6$ octahedra structure does not function as an OER site. Figure 7 shows a schematic model for water oxidation during upward and downward potential sweeps. The upward
potential change results in oxidation of the Ni$^{3+}$ species with Ni(OH)$_2$ structure to the Ni$^{3+}$ species with NiO$_6$ octahedra structure accompanied by OER activity. In contrast, the downward potential change gradually reduces the Ni$^{3+}$ species to Ni$^{2+}$ from the domain edge and the OER activity is suppressed when the outer NiO$_6$ octahedra domain is reduced. On the basis of these results, water oxidation is likely to proceed at the domain edge of NiO$_6$ octahedra.

Reaction pathways for OER catalysis with Ni$\cdots$B, Co$\cdots$B, and Co$\cdots$P electrocatalysts were previously proposed.

Bediako et al. elucidated that the OER over the Ni$\cdots$B electrocatalyst in borate-buffered electrolyte entails the reversible dissociation of borate anions from the domain edge of the NiO$_6$ octahedra structure and a two-electron and three-proton equilibrium, followed by a chemical turnover-limiting step. Our present study spectroscopically confirmed the report of Bediako et al. in that the active species functions at the domain edge of the NiO$_6$ octahedra structure. Thus, an increase in the number of NiO$_6$ domains is necessary to achieve an efficient OER catalyst.

The octahedra structures of transition-metal oxides (MnO$_6$, CoO$_6$, and NiO$_6$) are generally considered to exhibit high OER activity; however, the direct spectroscopic detection of active octahedral species had not been achieved around the OER electrode potential. The XAFS technique used here has provided the first successful direct probing of the active species of Ni$\cdots$B$_{14-19}$, Co$\cdots$B$_{17-18}$, and MnO$_{10,50}$ to reveal the relationship between the active species and OER activity. The knowledge obtained by this study will be useful to develop more efficient water-splitting systems.

4. CONCLUSIONS

In summary, in situ O K-edge XAFS spectra were measured for a nickel–borate OER electrocatalyst. The XAFS peak was observed at 28.7 eV by changing the electrode potential for each sample, which demonstrates that higher order domains of the edge-sharing NiO$_6$ octahedra structure are generated at higher potentials. The upward potential dependence of the XAFS spectra resulted in an increase in the intensity of the absorption peak for the NiO$_6$ octahedra with the OER current. In contrast, the downward potential dependence of the XAFS spectra revealed that the NiO$_6$ peak remains even if the OER current is suppressed. The different behavior between the upward and downward potential sweeps indicates that water oxidation catalysis is likely to proceed at the domain edge of the NiO$_6$ octahedra. This is the first successful direct observation of active nickel oxide species reported for OER electrocatalysts.

ASSOCIATED CONTENT

2 Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acs.jpcc.5b06102. Experimental procedures of in situ Ni K-edge XAFS, time course of current during electrodeposition and anodic oxidation, estimation of film thickness, in situ Ni K-edge XAFS spectra, and estimation of cation valence. (PDF)
Oxides


