Cu$_2$O/AlZn-LDH Preparation on FTO Electrode Substrate for Enhanced Water Oxidation

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Abstract: The electrochemical activity of the Cu$_2$O/AlZn-LDH, deposited on the FTO electrode, was investigated in NaOH (0.1 M) media. The thin film Cu$_2$O/AlZn-LDH was characterized by techniques such as X-ray diffraction (XRD), scanning electron microscopy (SEM), and energy-dispersive X-ray spectroscopy (EDX). Based on the obtained results from XRD, Cu$_2$O/AlZn-LDH was prepared in the nanometer size with good crystalline quality. The SEM images were shown the synthesis of a thin film Cu$_2$O/AlZn-LDH nanocomposite on the FTO electrode. The water oxidation results show that Cu$_2$O-Zn/Al-LDH modified FTO electrode is an improved electrocatalyst and has high activity at water oxidation in alkaline media with the onset potential about 0.6 V vs. SCE and overpotential of 380 mV at10 mA/cm. The improved water oxidation activity at Cu$_2$O/AlZn-LDH can be attributed to the good conductivity of the nanocomposite.

Keywords: Electro catalyst, Layered Double Hydroxide (LDH), FTO Electrode, Oxygen evolution reaction (OER)

1. INTRODUCTION

In recent years, in order to obtain clean energy, the electrochemical water oxidation has been increasingly studied.1-5 Among all the researches, the oxygen evolution reaction (OER) using electrocatalysts is considered as a key issue at water oxidation. In this process, four-electrons are transferred in an anodic reaction.6-10 In the recent years, various metal electrocatalysts such as oxides,11-13 phosphides,14-17 and hydroxides18, 19 have been applied as new materials for water oxidation. The oxygen generation process is kinetically sluggish, because at water oxidation, potential about 1.23 V is needed due to multistep proton-coupled electron transfer (PCET).20, 21 The water oxidation reaction in base or acidic solution is occurred as follows:

$$4\text{OH}^- \rightarrow 2\text{H}_2\text{O} + \text{O}_2 + 4\text{e}^- \quad \text{base solution} \quad (1)$$

$$2\text{H}_2\text{O} \rightarrow 4\text{H}^+ + \text{O}_2 + 4\text{e}^- \quad \text{acidic solution} \quad (2)$$

The first-row transition metal electrocatalysts in forms such as nanocomposites,22-26 perovskites,27 and layered double hydroxides,28-30 with high activity have been comprehensively studied. In the recent years, important studies on improving of LDH-based materials have been reported. In 2015, Dai et al. published a mini-review for water oxidation using NiFe-based electrocatalysts,31 where the applications and related mechanisms have been bringing up briefly. Shao et al. reported the improvement of Layered based materials for energy conversion using electrocatalysts.32 In the mentioned study, the applications of water oxidation have been discussed. Also, a review article was published by Strasser et. al. which was focused on the development of NiFe-based materials.33 Other privileged researches have been reported LDH-based materials for water oxidation, including the design of electrocatalyst oxides and hydroxides of transition metal by Boettcher et al.34 Layered double hydroxides (LDHs), are a family of anionic clays. In these materials the general formula is reported as [M$^{II}_{1-x}$M$_n$(OH)$_2$]$^{x+}$ [A$^{x-}$]$_{y}$H$_2$O]$^{z-}$, where M$^{II}$ and M$^{III}$ represent trivalent and divalent metal ions, respectively, and A$^{x-}$ is defined as an anion with n valent. The LDH layers consist positive charge due to partial substitution of M$^{II}$ by M$^{III}$. This positive charge was neutralized using anions which intercalated at the interlayer of LDHs.35 Transition metal based LDHs, have
received considerable attention due to their potential efficiency in catalysis, adsorbents, drug delivery, anion exchangers, biotechnology, photoactive materials, electrochemistry, nanocomposites, magnetization, and environmental applications. Also, the LDHs have some interesting properties, such as simple and economical synthesis in the laboratory and industrial scales with properties such as nanoscale size, specific structure, improved chemical composition, large surface areas, variable layer charge density, colloidal properties, flexible tunability, swelling in water, good thermal stability, etc.\(^3^6\) Synthesis of LDHs on the FTO was interested at last years. FTO coated glasses have many advantages; example being: thigh electrical conductivity, good thermal and mechanical resistance, providing in any dimensions necessary, reasonably priced, and using in photo-electronic and photovoltaic devices. They are stable in different environmental conditions and also are chemically inert. In the present work, Cu\(_2\)O/AlZn-LDH was prepared using the simple electrodeposition method on the FTO electrode. The obtained nanocomposite was then applied as electrocatalysts at water oxidation in 0.1 M KOH with high activity and small overpotential.

2. EXPERIMENTAL

Apparatus and software

Electrochemical investigations were achieved using a Sama 500 Electrochemical Analysis System (Sama, Iran) interfaced with a personal computer (PC) and a three-electrode configuration. FTO electrode modified with Cu\(_2\)O/AlZn-LDH was applied as the working electrode. A saturated calomel electrode and a Pt wire were utilized as the reference and counter electrodes, respectively (the working, counter, and reference electrodes were obtained from Azar Electrode Co, Urmia, Iran). The pH measurements were carried out using a Metrohm pH-meter 691. All the measurements were carried out at room temperature (25 ± 0.5 °C). The X-ray diffraction (XRD) patterns were collected with a Bruker AXS model D8 Advance diffractometer using Cu-K\(_\alpha\) radiation (\(\lambda = 1.542\) Å), with the Bragg angle of 2-70°, performing steps of 0.04°(2θ), and counting 4 s/step. The scanning electron microscopy (SEM)/energy dispersive X-ray (EDX) analysis was performed using a MIRA3 TESCAN scanning electron microscope equipped with an EDX system. The accelerating voltage was 10 kV, with a beam current of 1 nA and a spectrum collection time of 100 s. The FT-IR spectra were recorded on a Bruker tensor 27 spectrometer.

Reagents and solutions

Zn(NO\(_3\))\(_2\)·6H\(_2\)O, Al(NO\(_3\))\(_3\)·9H\(_2\)O and CuSO\(_4\)·5H\(_2\)O were applied without further purification and were prepared from Merck Company. Citric acid (0.960 g) and CuSO\(_4\)·5H\(_2\)O (1.248 g) were solved in 100 mL of deionized water for preparation of a 0.05 M solution contain citric acid (C\(_4\)H\(_4\)O\(_7\)) and copper sulfate (CuSO\(_4\)). Also, KCl (0.187 g), [Fe(CN)\(_6\)]\(^{3-}\) (0.008 g) and [Fe(CN)\(_6\)]\(^{4+}\) (0.010 g) were solved in deionized water (25 mL) to prepare solutions contain KCl (0.1 M), and (0.5 mM) [Fe(CN)\(_6\)]\(^{3-}\) and Fe(CN)\(_6\)]\(^{4+}\).

Fabrication of modified electrode

For the preparation of AlZn-LDH, the FTO electrode (0/5 cm × 0/5 cm) was first immersed in an electrochemical cell containing 15 mM Zn(NO\(_3\))\(_2\)·6H\(_2\)O (as the Zn\(^{2+}\) source) and 5 mM Al(NO\(_3\))\(_3\)·9H\(_2\)O (as the Al\(^{3+}\) source) solution. Then AlZn-LDH was deposited on the surface of the FTO electrode as thin film by applying a potential of -1.65 V/SCE for 120 s using the electro-deposition method. For the synthesis of Cu\(_2\)O/AlZn-LDH, the prepared AlZn-LDH electrode was immersed in an electrochemical cell containing 0.05 M CuSO\(_4\) and 0.05 M citric acid solution and nanocomposite was synthesized by chronoamperometry method at potential of -0.5 V for 180 s at the 65 °C. Then the prepared electrode was kept at temperature 80 °C for 2 h.

3. RESULTS AND DISCUSSION

X-ray diffraction

The XRD patterns of AlZn-LDH and AlZn-LDH/Cu\(_2\)O nanoparticles are shown in Figure 1(a) and (b). The X-ray diffraction pattern of the AlZn-LDH synthesized on the FTO electrode is shown in Figure 1(a). The sharp and strong diffractions at 2θ = 9.91° and 19.78°, are related to planes of (003) and (006) at layered double hydroxide, respectively, and also the diffraction in the 2θ = 33.68 ° is related to ZnO (002) plate. Comparison with the reference diffraction pattern indicates that the pattern is completely consistent with the pattern obtained by other researchers and does not contain any impurities.\(^7\) As can be seen, sharp diffractions in 2θ are equal to 33.46° and 43.12°.

![Figure 1. XRD patterns of (a) AlZn-LDH, and (b) Cu\(_2\)O/AlZn-LDH.](image-url)
correspond to the plants of (111) and (200) at Cu$_2$O. Also, the sharp diffractions in the 2θ about 26.29°, 37.55°, 51.31°, 61.40°, and 65.33° are related to the FTO electrode. The obtained results are consistent with the reported results by other researchers. Also, the sharp and strong diffractions in region 2θ are equal to 9.91° are related to (003) LDH plate. The reported patterns indicate that the synthesized sample has a layered double hydroxide crystal structure and completely is composited with Cu$_2$O.

**SEM image**

SEM images of Cu$_2$O/AlZn-LDH coating on the FTO electrode with two different magnifications are shown in Figure 2(a) and (b). The synthesized compound consists of spherical nanoparticles that are stacked together.

The obtained SEM images show the synthesis of Cu$_2$O/AlZn-LDH with the uniformity and homogeneity nanoparticles on the surface of the FTO electrode. According to SEM images, particle sizes are estimated between 20 and 30 nanometers.

**EDX (Energy-dispersive X-ray spectroscopy) and elemental map analysis**

EDX analysis was reported in Figure 3 to investigate the chemical elemental of the synthesized Cu$_2$O/AlZn-LDH. The resulting EDX spectrum shows the existing peaks of the aluminum and zinc cations in the LDH. Also, the peaks of copper and oxygen atoms related to Cu$_2$O in Cu$_2$O/AlZn-LDH compound are also observed. The intensity of oxygen peak is low because the oxygen atom is lighter than Cu, Al and Zn atoms, so other relaxation processes may compete with the EDX. In addition, the presence of tin peaks, are related to the FTO electrode. The Au peak is also appeared due to covering the surface of the material with gold, for SEM imaging.

The elemental map was used to investigate the distribution of Cu$_2$O/AlZn-LDH synthesized on the FTO electrode. Figure 4 confirmed the presence of Zn, O, Al,
and Cu elements in the prepared composite. Based on this distribution map, we conclude that the synthesized CuO/AlZn-LDH composite elements uniformly, homogeneously, and completely cover the surface of the FTO electrode.

**Electrochemical synthesis and behavior**

Electrochemical deposition of AlZn-LDH could be achieved as follows. By applying negative potentials to the electrode, nitrate anions and zinc cations could be reduced on the FTO electrode. Since the reduction of zinc ions occurs at more negative potentials than the nitrate ions, in the presence of nitrate ions the reduction of zinc ions is not possible. The hydrogen evolution reaction was also may be occurred. But at the used conditions for pH, mole ratio of Zn\(^{2+}\)/NO\(^{-}\) and the applied potential, only reduction of nitrate was occurred.\(^{37}\) It should also be noted that at high temperatures and more negative potentials, ZnO is electrodeposited on the electrode.\(^{37}\) Thus, at room temperature and at the optimum conditions, AlZn-LDH is electrochemically deposited on the electrode by following possible mechanism.

\[
\begin{align*}
NO_2^- + H_2O + 2e^- & \rightarrow NO_3^- + 2OH^- \\
(1-x)Zn^{2+} + xAl^{3+} + xNO_2^- & \rightarrow (Zn)_{1-x}(Al)^x((OH)_x(NO_2)_x)
\end{align*}
\]

Figure 5a shows the cyclic voltammogram of AlZn-LDH in the 0.1M KOH solution. This cyclic voltammogram shows the AlZn-LDH was formed on the electrode. By applying -0.5V on AlZn-LDH electrode, a thin layer of CuO was formed on the modified electrode in the presence of 0.05M citric acid as chelating agent \(^{39}\). Figure 5b also shows the cyclic voltammogram of CuO/AlZn-LDH in the same solution.

**Water oxidation**

To evaluate the water oxidation activity of prepared materials, LSV curves were recorded with a scanning speed of 50 mV s\(^{-1}\) in 0.1 M KOH solution using the reference calomel electrode and platinum electrode as the counter electrode. The LSV voltammograms were reported in Figure 6. According to these data, it is clear that in comparison to CuO, AlZn-LDH and CuO/AlZn-LDH, the prepared composite (CuO/AlZn-LDH) has a significant activity for the oxygen evaluation reaction. The onset potential for CuO/AlZn-LDH was reported about 0.6 V vs. SCE, while onset potential at CuO and AlZn-LDH were reported 0.75 and 1.25 respectively. The current density of 10 mA/cm\(^2\) was represented about 10% conversion efficiency at fuel systems using solar energy.\(^{40, 41}\) So, it is important criteria at water oxidation that CuO/AlZn-LDH needs an overpotential of about 380 mV to obtain the current density about 10 mA cm\(^{-2}\) although, at CuO and AlZn-LDH, the current density and catalytic activity were lower than that of CuO/AlZn-LDH. Despite the low overpotentials, the CuO/AlZn-LDH showed the highest current density at a certain applied potential. This result underscores the importance of assembling catalytically active materials at the molecular level when designing high performance electrochemical catalysts.\(^{42}\)

![Figure 5](image5.png)

**Figure 5.** The cyclic voltammograms of (a) AlZn-LDH and (b) CuO/AlZn-LDH modified FTO electrode in the 0.1 M KOH solution at the scan rate 50 mV s\(^{-1}\).

![Figure 6](image6.png)

**Figure 6.** LSV curves of CuO, AlZn-LDH and CuO/AlZn-LDH.

Tafel slope, which describes the influence of potential or overpotential on the current density, is an important factor for the evaluation of OER kinetics. Tafel slope is often influenced by the electron and mass transport.\(^{43}\) Tafel analysis of the OER on CuO/AlZn-LDH and AlZn-LDH based electrocatalysts were conducted and the results are presented in Figure 7. within the electrocatalyst series, the Tafel slope increased in the following order: CuO/AlZn-LDH (593 mV/dec) < AlZn-LDH (677 mV/dec). The small Tafel slope of CuO/AlZn-LDH indicated that the electron and mass transfer easily occurred in CuO/AlZn-LDH compared to AlZn-LDH. Therefore, the Tafel result supports the conclusion that the advanced electrocatalytic performance of CuO/AlZn-LDH for OER was due to the presence of active sites (Al\(^{13+}\), Cu\(^{2+}\), and Cu\(^{1+}\)) in CuO/AlZn-LDH.
The electrochemical stability of Cu$_2$O, AlZn-LDH, Cu$_2$O/AlZn-LDH electrodes were investigated by chronoamperometry in 0.1 M KOH solution. The chronoamperometry method expresses the time-dependent behavior and can report information on the durability of electrocatalyst through water oxidation reaction. As shown in Figure 8, the current densities remained constant for all three electrodes at constant overpotential within 50 s water oxidation. Therefore, the structures of AlZn-LDH, Cu$_2$O, Cu$_2$O/AlZn-LDH are stable in the water oxidation reaction. Also after 5 cycle chronoamperometry test, the stable current density with negligible degradation was observed, revealing their durability under water oxidation conditions.

For the electrocatalytic OER over semiconductors, the possible mechanisms at alkaline solution have been reported. The reaction mechanism reported by Hunter et al. was as follows:\(^5\)

\[
4\text{OH}^- \rightarrow \text{O}_2 \ (g) + 2\text{H}_2\text{O} \ (l) + 4\text{e}^- \quad (5)
\]

\[
4\text{H}_2\text{O} \ (l) + 4\text{e}^- \rightarrow 2\text{H}_2 \ (g) + 4\text{OH}^- \quad (6)
\]

\[
2\text{H}_2\text{O} \ (l) \rightarrow \text{O}_2 \ (g) + 2\text{H}_2 \ (g) \quad (7)
\]

Also, the electrocatalytic activity of Cu$_2$O/AlZn-LDH in comparison with the electrocatalysts reported in the literature, was summarized in Table 1.

<table>
<thead>
<tr>
<th>Electrolyte</th>
<th>pH</th>
<th>Onset potential (V) SCE</th>
<th>Over potential at 10 mA cm$^{-2}$</th>
<th>Ref.</th>
</tr>
</thead>
<tbody>
<tr>
<td>NiCu-LDH/SAV</td>
<td>1 M KOH</td>
<td>0.45</td>
<td>290</td>
<td>45</td>
</tr>
<tr>
<td>Cu@NiFe-LDH</td>
<td>1 M KOH</td>
<td>0.39</td>
<td>199</td>
<td>46</td>
</tr>
<tr>
<td>CC/NiCu-LDH</td>
<td>1 M KOH</td>
<td>0.44</td>
<td>370</td>
<td>47</td>
</tr>
<tr>
<td>NiCu-LDH</td>
<td>1 M KOH</td>
<td>0.54</td>
<td>770</td>
<td>48</td>
</tr>
<tr>
<td>Cu$_2$O/AlZn-LDH</td>
<td>0.1 M KOH</td>
<td>0.6</td>
<td>380</td>
<td>This study</td>
</tr>
</tbody>
</table>

\footnotesize{This study}
4. CONCLUSIONS
The thin film Cu$_2$O/AlZn-LDH was successfully prepared on a FTO electrode using an easy, one-step electrodeposition method. This method can be applied for the synthesis of LDH based nanocomposite at a large scale. The obtained nanocomposite was characterized by different techniques. In this study, the electrochemical water oxidation activity of Cu$_2$O/AlZn-LDH was studied. The obtained results confirm that this nanocomposite can be used to the electrocatalytic water oxidation. The proposed Cu$_2$O/AlZn-LDH exhibited good water oxidation activity with the onset potential about 0.6 V vs. SCE and an overpotential of 380 mV at 10 mA cm$^{-2}$. The increased water oxidation activity of the Cu$_2$O/AlZn-LDH is attributed to a decrease in the charge transfer resistance which influences the electronic structure and consequently increases the electric conductivity and improves the water oxidation.

CONFLICTS OF INTEREST
There are no conflicts to declare.

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