CO₂-Induced Fibrous Zn Catalyst Promotes Electrochemical Reduction of CO₂ to CO

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Abstract: The electrochemical reduction of CO₂ is a promising strategy to achieve efficient conversion and utilization. In this paper, a series of Zn catalysts were prepared by electrodeposition in different atmospheric conditions (N₂, CO₂, H₂, CO). A fibrous Zn catalyst (Zn-CO₂) exhibits high electrochemical activity and stability. The Zn-CO₂ catalyst shows 73.0% faradaic efficiency of CO at -1.2 V vs. RHE and the selectivity of CO almost did not change over 6 h in -1.2 V vs. RHE. The excellent selectivity and stability is attributed to the novel fibrous morphology, which increases the electrochemical active surface area. X-ray diffraction (XRD) results show that Zn-CO₂ catalyst has a higher proportion of Zn (101) crystal planes, which is considered to be conducive to the production of CO. The search further demonstrates the importance of morphology control for the preparation of highly active and stable catalysts.

Keywords: electrocatalysis; electrodeposition; CO₂ reduction; fibrous structure; porous

1. Introduction

Over the past few years, heavy consumption of fossil fuels has increased the concentration of carbon dioxide (CO₂) in the atmosphere, leading to many environmental problems such as greenhouse effect and energy dilemma [1]. It is widely accepted that reducing the concentration of CO₂ is a key to solving environmental problems. The transformation and utilization of CO₂ has become the focus of research for researchers. [2]. CO₂ can be converted to high-value products such as carbon monoxide, methane, formic acid, methyl alcohol, ethyl alcohol, ethylene and so on [3–8], by electrochemical CO₂ reduction reaction (CO₂RR) under mild conditions. Therefore, electrochemical CO₂RR is the most promising technology to achieve the conversion and utilization of CO₂. The challenge of electrochemical CO₂RR is the preparation of catalysts with high activity, selectivity and stability, which is the core for catalytic research.

Up to now, researchers have developed a number of catalysts for CO₂RR. Noble metals such as Au [9–13] and Ag [14–19] are known to be the best metal catalysts for converting CO₂ to CO with high activity and selectivity. However, it is difficult to use in large quantities in industrialization, due to the high price and scarce reserve of precious metals. Therefore, research into efficient non-noble metal catalysts has become an important part of the industrial application of electroreduction of CO₂. Zn, an abundant metal, is proved to be an outstanding electrocatalyst to produce CO [20–23]. However, there are many challenges for the practical application of CO₂ reduction. The reaction still suffers from poor stability and faradaic efficiency (FE) of the primary product. Therefore, the preparation of catalysts with high activity and selectivity is the core of future development.
There are many researchers who have studied Zn catalyst in recent years. In order to improve the performance of Zn catalysts for CO₂RR, many strategies have been used in the preparation of catalysts, such as electrodeposition, anodization, oxide reduction [24]. Preparation of Zn catalyst with large surface area is considered as an effective method to enhance the activity of the catalyst. For example, electrodeposited Zn dendrite electrode has a surface area of around 90 cm²/cm²Geo, corresponding to over 80 times higher surface area than Zn bulk foil electrodes, and the FE of CO (FE₃) is around 3-fold higher [25]. Recently, some researchers have studied the synthesis of catalysts in aqueous electrolyte with CO₂-rich condition to improve the activity of catalysts. For example, Wang et al. synthesized Pb catalyst in an aqueous electrolyte with CO₂-rich condition, and formic acid FE reached 90.5% [26]. Wang et al. reported Cu catalyst, which was synthesized under CO₂ reduction conditions, with a 90% FE for C₂H₄ [27]. This indicates that CO₂ plays a key role in the preparation of highly active catalysts by electrodeposition.

In this work, a series of Zn catalysts were prepared by electrodeposition in different gas-saturated (N₂, CO₂, H₂, CO) Zn(NO₃)₂ aqueous solutions. Linear sweep voltammetry (LSV), cyclic voltammetry (CV) and Tafel slope were employed to study the performances of different Zn catalysts for CO₂RR. Zn-CO₂ catalysts have the best activity and selectivity for electrochemical CO₂RR. The Zn-CO₂ electrode shows 73.0% faradaic efficiency of CO at −1.2 V vs. RHE. The partial current density of CO is 6.1 mA/cm² and the selectivity of CO has almost not changed over 6 h in −1.2 V vs. RHE. X-ray diffraction (XRD), scanning electron microscopy (SEM), transmission electron microscopy (TEM) and X-ray photoelectron spectroscopy (XPS) characterization methods were used to investigate the relation of the structure and excellent activity of as-prepared catalysts in detail. XRD results show that Zn-CO₂ catalyst has a higher proportion of Zn (101) crystal planes. The SEM images show that the surface of Zn-CO₂ catalyst is covered by porous fibers with uniform tiny pores, which increases the electrochemical active surface area (ECSA). This is the main reason for the excellent catalytic performance of Zn-CO₂ catalysts. The research prepared a Zn catalyst with new morphology and excellent activity and stability of electrochemical CO₂RR, which develops a new means and idea for the preparation of more efficient and stable Zn catalysts. This work is of great value and significance in solving the challenge of low activity and poor stability in electrochemical CO₂RR.

2. Results and Discussion
2.1. Electrochemical Performance

The electrochemical performance of the prepared electrodes for CO₂RR was evaluated by linear sweep voltammetry (LSV) at 5 mV/s in pure KHCO₃ electrolyte solution saturated with CO₂. As shown in Figure 1a, the current density of the Zn-CO₂ electrode is the highest and far higher than that of other electrodes, which indicates that the Zn-CO₂ electrode has the best catalytic activity. To evaluate the electrochemical activity of Zn catalysts, the controlled potential electrolysis experiment was conducted at different cathodic potential to the CO₂-saturated KHCO₃ electrolyte. Before the controlled potential electrolysis experiment, cyclic voltammetry (CV) was performed over the potential range from −0.55 V vs. RHE to −1.35 V vs. RHE (50 mV/s) for 20 cycles to reduce the oxide layer of the catalyst. The main gas products which were detected every 30 min by gas chromatography (GC) were CO and H₂. Figure 1b shows that the FE of Zn foil electrode gradually increases with the decrease of applied voltage, and reaches a maximum of 47.0% at −1.3 V. Zn-N₂ electrode has a FE of 66.2% at −1.0 V vs. RHE, and decreases gradually with the decrease of applied voltage which indicates that the H₂ evolution reaction (HER) is more serious at more negative voltage for Zn-N₂ electrode. This can also be demonstrated by the diagram of the FE of H₂ as a function of voltage (Figure 1c), which FE of H₂ increases gradually when the voltage exceeds −1.2 V vs. RHE. The maximal FE of CO₂ is found on Zn-CO₂ electrode with 73.0% at −1.2 V vs. RHE (Figure 1b), and with the lowest FE of 32.6% (Figure 1c), and the partial current density of CO is 6.1 mA/cm² at −1.2 V vs. RHE (see Table S1). This indicates the production of CO is promoted and the production
of H₂ is inhibited at −1.2 V. The Zn-H₂ electrode has a FE₈ of 47.9% at −1.2 V vs. RHE, and the Zn-CO electrode has a FE₈ at 56.7% at −1.1 V vs. RHE. These results show that the catalyst prepared by electrodeposition in CO₂-saturated Zn(NO₃)₂ aqueous solution can effectively promote the production of CO. To evaluate the long-term performance of the Zn-CO₂ electrode, a chronoamperometry experiment at −1.2 V vs. RHE was carried out. As shown in Figure 1d, the current density is stable at 4.6 mA/cm², and the FE₈ remains unchanged (~73.0%) over 6 h. The stability evaluations of other electrodes under the same conditions were also carried out, and the results are shown in Figure S1. The FE₈ for Zn foil and Zn-H₂ electrode decreased gradually with the increase of time during 6 h, while their current density is still stable. The current density for the Zn-N₂ electrode maintains stable before 5 h and then decreases dramatically after reaction of 5 h, at the same time, the FE₈ fluctuates between 40.0% and 60.0%. A similar phenomenon appears in the Zn-H₂ electrode. This shows the Zn foil, Zn-N₂, Zn-H₂ and Zn-CO electrodes have very poor stability. According to these comparative experimental results, the Zn-CO₂ electrode exhibits excellent stability.

Figure 1. (a) Linear sweep voltammetry (b) faradaic efficiency (FE) of CO (c) FE of H₂ for Zn foil, Zn-N₂, Zn-CO₂, Zn-H₂, Zn-CO electrode at 5 mV/s. (d) Current density and FE₈ for Zn-CO₂ electrode during 6 h of long-time operation at −1.2 V vs. RHE.

2.2. Characterization of Catalysts

A lot of characterizations were performed to analyze the physical and chemical properties of the catalysts. First of all, XRD was measured to analyze the crystal structure of electrodeposited Zn. Figure 2a exhibits the XRD patterns of all prepared Zn electrodes and Zn foil electrode. It shows that there appear peaks at 36.3°, 39.0°, 43.2°, 54.3°, 70.1°, 70.6°, and 77.0° corresponding to (002), (100), (101), (102), (103), (110) and (004) lattice planes of metallic Zn [JCPDS file No. 04-0831], respectively [28] indicating that Zn catalysts with high crystallinity can be prepared by electrodeposition in different gas-saturated Zn(NO₃)₂ aqueous solutions. In order to research the influence of different atmospheres for electrodeposited Zn catalysts, the texture coefficient of each crystal face for the Zn catalysts is calculated and dotted in Figure 2d. The texture coefficients of the Zn (101) plane are 16.79%, 19.16%, 20.79%, 20.6%, 17.59% corresponding to Zn
foil, Zn-N\textsubscript{2}, Zn-CO\textsubscript{2}, Zn-H\textsubscript{2}, Zn-CO electrode, respectively, which are consistent with the trend of corresponding current densities. The Zn-CO\textsubscript{2} electrode has the highest Zn (101) plane ratio while Zn foil has the minimum Zn (101) plane ratio. This is due to the strong binding ability of intermediates formed during Zn(NO\textsubscript{3})\textsubscript{2} electroreduction in CO\textsubscript{2}-saturated aqueous solutions, which promotes the specific growth of Zn crystals on electrode [26]. As reported in literature [29,30], the Zn (101) plane is conducive to the CO\textsubscript{2}RR to CO. On the other hand, the Zn (002) plane is responsible for H\textsubscript{2} evolution in electrochemical CO\textsubscript{2}RR [29,30]. The texture coefficients of the Zn (002) plane are 16.00%, 13.16%, 10.51%, 11.27%, 11.85% corresponding to Zn foil, Zn-N\textsubscript{2}, Zn-CO\textsubscript{2}, Zn-H\textsubscript{2}, Zn-CO electrode, respectively. The Zn-CO\textsubscript{2} electrode has a small Zn (002) plane ratio while the Zn foil electrode has the biggest Zn (002) plane ratio. The different proportion of crystal face of Zn catalysts may promote the formation of different morphologies. The greater proportion of Zn (101) promotes the generation of CO on Zn-CO\textsubscript{2} electrode.

Figure 2. X-ray diffraction (XRD) patterns of prepared Zn electrodes: (a) before reaction, (b) after reaction and (c) the reduced Zn-CO\textsubscript{2} electrode; (d) texture coefficients of Zn electrodes before reaction.

The XRD patterns of used catalysts and reduced Zn-CO\textsubscript{2} electrode were also given. As shown in Figure 2b, the XRD patterns of used catalysts are the same as the corresponding fresh catalysts, and there are no any new phases appearing in these catalysts. In the reduced Zn-CO\textsubscript{2} electrode, the XRD pattern also exhibits the characteristic reflections of a single crystalline Zn phase. This indicates that catalyst has been reduced by electrochemical reduction methods.

The surface morphology of representative Zn electrodes was characterized by a scanning electron microscope (SEM) and shown in Figure 3. Obviously, each electrode surface is coarse and porous. On the surface of the Zn-N\textsubscript{2} and Zn-CO electrode, the pore channels with various sizes are found to be partially covered by non-porous sheets. For the Zn-H\textsubscript{2} electrode, these non-porous sheets covered on the surface of the hierarchical pores fully disappear and are replaced by abundant tiny pores. In particular, the surface of the Zn-CO\textsubscript{2} electrode is covered by porous fibers with uniform tiny pores. The different surface morphology should be due to different gas dissolving capacity in Zn(NO\textsubscript{3})\textsubscript{2} solution and interaction among gas,
Zn(NO$_3$)$_2$ and Zn foil. The special morphology of the Zn-CO$_2$ electrode is just due to the higher solubility of CO$_2$ in aqueous solution (see Table S2) and a stronger interaction between CO$_2$ and Zn interface than those of other gases resulting in the stable and uniform growth of Zn crystal during the whole electrodeposition process. Correspondingly, the heterogeneous surface morphology for other electrodes should be related to the low solubility and quick escape of gases from the solution. The surface morphology plays an important role in the CO$_2$RR on a three-phase interface. According to the relative high current density of CO$_2$ conversion on the Zn-CO$_2$ electrode shown in Figure 1a, it can be confirmed that the uniform and abundant tiny pores on the Zn-CO$_2$ electrode may be more beneficial for CO$_2$ enrichment by enhancing surface area than other electrodes with low surface area resulting from the bigger pore channels and even non-porous surface. As for the CO and H$_2$ selectivity, it should be the result of the combined effects, including the pore diffusion and surface enrichment of CO$_2$ and H$_2$O molecules, under efficient three-phase contact of CO$_2$ gas, liquid H$_2$O and solid catalyst [31]. The high FE$_{CO}$ for the Zn-CO$_2$ electrode should be attributed to the special surface morphology, which can increase surface area to accommodate high concentrations of CO$_2$ and the uniform tiny pores on porous fibers are beneficial to CO$_2$ permeation and detrimental to H$_2$O molecular diffusion. Instead, the appearance of big pores and non-porous sheets means the shrinkage of surface area, which results in the decrease of surface concentrations of CO$_2$ and H$_2$O, but is conducive to the H$_2$O molecular diffusion, resulting in the possible increase of FE$_{H_2}$ and decrease of FE$_{CO}$ in a certain degree. Certainly, the catalytic performance should be determined by the competition between two reactions of CO$_2$RR to CO and HER (also known as electrochemical water reduction to H$_2$), which depends on the concentration and diffusion ability of CO$_2$ and H$_2$O molecules.

Figure 3. Scanning electron microscope (SEM) images of representative Zn electrodes: (a) Zn-N$_2$, (b) Zn-CO$_2$, (c) Zn-H$_2$, (d) Zn-CO.

In addition, we compared the morphology of the catalyst after the reaction (Figure 4) and found that the Zn foil, Zn-N$_2$ and Zn-CO$_2$ catalysts accumulated into a regular nanosheet structure. The nanosheet of the Zn-N$_2$ electrode is much larger in size than other electrodes after reaction, and more irregular than other electrodes. It is the reason that current density is higher in the initial stage and more unstable. The nanosheet greatly reduced the ECSA of the catalysts, which was one of the reasons for the catalyst’s death, after 6 h of reaction.
Figure 4. SEM images of representative Zn electrodes after reaction: (a) Zn foil electrode, (b) Zn-N$_2$ electrode, (c) Zn-CO$_2$ electrode.

Furthermore, the representative Zn-N$_2$ and Zn-CO$_2$ electrodes were also analyzed by transmission electron microscope (TEM) and the TEM images for these representative samples are shown in Figure 5. The nanosheets can be found easily on each sample, which constructs the skeleton of pore channels. Obviously, the size of nanosheets on the Zn-CO$_2$ electrode is uniform and far less than that on the Zn-N$_2$ electrode. The results are consistent with those of SEM. In addition, the existence of the Zn (002) plane can further be demonstrated by the lattice spacing of 0.24 nm of two electrodes. Besides the metal Zn phase, the characteristic spacing of 0.26 nm for ZnO (002) for the Zn-N$_2$ electrode, and 0.28 nm for the ZnO (100) lattice plane for the Zn-CO$_2$ electrode can be seen. The results also confirm that ZnO can be formed during the preparation process, indicating the importance of electrochemical reduction of the electrode prior to the reaction.

X-ray photoelectron spectroscopy (XPS) was performed to demonstrate the chemical states of Zn and O elements on the surface of the catalysts. The XPS spectra are given in Figure 6 and the characteristics data are collected in Table 1. According to the XPS survey scan as shown in Figure 6a, it can be further confirmed that Zn and O elements exist on the Zn-N$_2$ and Zn-CO$_2$ electrodes. As for the XPS spectra of Zn 2P region, there are two deconvoluted peaks (Figure 6b). The peak at 1021.8 eV is associated with Zn 2P$_{3/2}$. Figure 6c shows the Zn 2P$_{3/2}$ spectrum of Zn-N$_2$ and Zn-CO$_2$ electrode. It shows Zn in two states, 1021.5 eV and 1022.1 eV, respectively. The low binding energy (BE) peak can be assigned to the metallic Zn (Zn$^0$), while the high BE peak should be ascribed to the oxidized Zn (Zn$^{2+}$). The ratio of Zn$^{2+}$/total of Zn-CO$_2$ electrode is higher than that of the Zn-N$_2$ electrode, it is due to the fibrous structure and pore diameter resulting in the formation of more under-coordinated Zn$^{2+}$ species on the surface of small particles, especially at the tip of pore mouths [25]. Before the reaction, the reduction treatment can transfer the under-coordinated Zn$^{2+}$ species into active Zn species. These oxidized Zn sites may facilitate the adsorption of reactants and *COOH intermediates on the surface of the Zn electrode, thus improving the generation of CO in the reaction [32]. Moreover, three fitted peaks can be seen in the O 1s regions. The peak at 530.6 eV is associated with O$^2$ from ZnO. The peak at 531.8 eV is related to hydroxyl group on the surface [33] and the peak at 532.8 eV is attributed to adsorbed oxygen. The adsorbed oxygen comes from oxygen in the absorbed water molecules [32,34]. The proportion of O 1s peak at
530.6 eV, associated with ZnO in the Zn-CO$_2$ electrode, is higher than the Zn-N$_2$ electrode. It indicated that the presence of CO$_2$ will promote the formation of under-coordinated Zn$^{2+}$ species on the surface of the Zn electrode in the process of electrodeposition. The under-coordinated Zn$^{2+}$ may facilitates the adsorption of *COOH intermediates on the surface of the Zn electrode and enhances the ability of intermediate to bind with the Zn electrode [35]. But this is not the significant distraction factor for catalytic excellent performance of Zn-CO$_2$. It is because Zn-N$_2$ and Zn-CO$_2$ electrodes have the under-coordinated Zn$^{2+}$ and the difference between them is not particularly great.

In order to explore the reasons for the superior CO$_2$RR performance of Zn-CO$_2$ electrode, it is necessary to compare the ECSA of each electrode by measuring their specific double-layer capacitance calculated by a CV scan in Figure S2. The result is shown in Figure S2, and the data of specific double-layer capacitance were calculated from this result. The line regression of the double layer capacitance and Tafel plot are displayed in Figure 7 and the corresponding ECSA and Tafel slope data of all samples are summarized in Table 2. The results show that the Zn-CO$_2$ electrode has the largest ECSA value (19.295 mF/cm$^2$) which is 60 times that of the Zn foil electrode (0.309 mF/cm$^2$), and the other electrodes, Zn-N$_2$, Zn-H$_2$ and Zn-CO, give a lower ECSA value (from 2.210 to 4.835 mF/cm$^2$). The quite high ECSA for Zn-CO$_2$ electrode is related to the special porous fibers with uniform tiny pores, which can enhance the surface concentration and diffusion ability of CO$_2$. In general, during electrodeposition, the presence of CO$_2$ affects the growth of crystal surfaces and promotes the formation of porous fibers morphology, which leads to a larger ECSA. This is the main reason for excellent performance of electrochemical CO$_2$RR for the Zn-CO$_2$ electrode.
Figure 6. XPS spectra of (a) overall survey scan, (b) Zn 2P, (c) Zn 2P\textsubscript{3/2}, (d) O 1s for the Zn-N\textsubscript{2} and Zn-CO\textsubscript{2} electrodes.

Table 1. X-ray photoelectron spectroscopy (XPS) characteristics data of Zn 2P\textsubscript{3/2} and O 1s regions for the Zn-N\textsubscript{2} and Zn-CO\textsubscript{2} electrodes.

| Catalysts | Binding Energy (eV) | Peak Area Ratio \textsuperscript{a} | Binding Energy (eV) | Peak Area Ratio |
|-----------|---------------------|-----------------------------------|---------------------|-----------------|
| Zn-N\textsubscript{2} | 1021.5 | 1022.1 | 0.15 | 532.8 | 531.8 | 530.6 | 0.21:0.61:0.18 |
| Zn-CO\textsubscript{2} | 1021.5 | 1022.1 | 0.22 | 532.8 | 531.7 | 530.7 | 0.20:0.59:0.21 |

\textsuperscript{a} Calculated from the fitting peak area results of Zn 2P\textsubscript{3/2} and O 1s XPS spectra.

Figure 7. (a) The line regression of the double layer capacitance and (b) Tafel Plot for CO\textsubscript{2}RR over Zn foil, Zn-N\textsubscript{2}, Zn-CO\textsubscript{2}, Zn-H\textsubscript{2}, Zn-CO electrode in CO\textsubscript{2}-saturated 0.1M KHCO\textsubscript{3} electrolyte.
Table 2. The electrochemical active surface area (ECSA) and Tafel slope data of all samples.

| Sample  | ECSA (mF/cm²) | Tafel Slope (mV/dec) |
|---------|--------------|----------------------|
| Zn foil | 0.309        | 211.70               |
| Zn-N₂   | 2.210        | 214.80               |
| Zn-CO₂  | 19.295       | 229.34               |
| Zn-H₂   | 4.735        | 239.54               |
| Zn-CO   | 4.835        | 253.28               |

To elucidate the origin of the remarkable CO₂RR performance over our catalysts, we performed detailed electrokinetic studies to compare the CO₂ reduction pathway on all electrodes by comparing the experimentally observed Tafel slope (logarithm of the partial current density towards a specific product vs. potential). There are four elementary reaction steps for CO₂RR to CO:

1. \(^*\text{CO}_2 + e^- \rightarrow ^*\text{CO}_2^-\)
2. \(^*\text{CO}_2^- + H^+ \rightarrow ^\text{COOH}\)
3. \(^\text{COOH} + H^+ + e^- \rightarrow ^\text{CO} + H_2O (l)\)
4. \(^\text{CO} \rightarrow ^\text{CO} + ^*\)

where * represents an active site. Firstly, the absorbed CO₂ gains an electron to form to \(^*\text{CO}_2^-\) anion radical. Secondly, the \(^*\text{CO}_2^-\) anion radical binds with a proton to form the \(^\text{COOH}\) intermediate. Thirdly, the \(^\text{COOH}\) intermediate gains a proton and an electron and is reduced \(^\text{CO}\) intermediated and H₂O. Finally, the \(^\text{CO}\) is desorbed from the surface of catalysts to form CO. It is well known that the value of a Tafel slope can reflect the rate-control steps of the reaction under certain conditions. Figure 7b shows that Zn-CO₂ electrode and Zn foil have Tafel slope of 229.34 mV/dec and 211.7 mV/dec, respectively. These are close to the theoretical value of 250 mV/dec for a rate limiting step at higher overpotential (−0.6 to −0.7 V) [24,36]. The higher value implies that the initial electron transfer step is the rate-determining step (RDS) [28]. The Zn-H₂ and Zn-CO electrodes have higher Tafel slope of 239.54 mV/dec and 253.28 mV/dec. From these data, it can be inferred that the RDS of electrochemical reduction reaction is mainly the first step of electron transfer step on these Zn electrodes. This indicated the \(^*\text{CO}_2^-\) intermediate plays a crucial role in the CO₂RR reaction. The same RDS indicated that the CO₂RR reaction may have similar intrinsic activity on these Zn catalysts [24].

3. Experimental Section

3.1. Fabrication of Zn Catalysts

Zn foil electrodes (1 cm × 2 cm) were prepared by mechanical polishing using sandpaper (800 grit), followed by sonication in acetone, ethanol and deionized water for 30 min. Zn(NO₃)₂ (Tianjin Fengchen Chemical Regents Factory, Tianjin, China)aqueous solution (30 mL, 0.05 M) was purged with different gases (N₂, CO₂, H₂, CO) as a precursor solution. A chronoamperometry experiment setting at −1.85 V vs. SCE (300 s) was used to reduce Zn precursor to metallic Zn on the surface of the prepared Zn foil electrode. After deposition, the electrode was rinsed by deionized water and then immediately taken for electrochemical testing to prevent electrode oxidation again.

3.2. Electrocatalytic Test for CO₂RR

An electrochemical workstation (CHI instrument 660E, Chenhua, Shanghai, China) was used for all electrochemical tests. The electrochemical measurements were performed in a three-electrode system. The reference electrode was a saturated calomel electrode (SCE)
and the counter electrode was a Pt sheet, respectively. All potentials in this work are quoted versus the reversible hydrogen electrode (RHE), according to the Nernst equation of:

$$E_{\text{RHE}} = E_{\text{SCE}} + 0.0591 \times pH + 0.241$$  \hspace{1cm} (5)

CO$_2$RR was conducted in a gas-tight two component H-type electrochemical cell separated by Nafion 117 cation-exchange membrane (Dupont, Wilmington, DE, USA). 0.1 M KHCO$_3$ (99.5%) aqueous solution was used as electrolyte and was purged with CO$_2$ (99.999%) at a rate of 20 mL/min for at least 30 min before the testing of CO$_2$RR. During the testing, the electrolyte was stirred by a magnetic stirring apparatus. The gas phase products were detected by a gas chromatograph (GC, SP2100) which was equipped with a TDX-01 column. A flame ionization detector (FID) was equipped with methanizer to analyze CO and a thermal conductivity detector (TCD) to analyze H$_2$. The FE of product was calculated by Equation (6).

$$\text{FE}\% = \frac{z n F}{Q} \times 100\%$$  \hspace{1cm} (6)

where $z$ is 2 for CO$_2$RR to CO or H$_2$, $n$ is the mole number of the products, Faraday’s constant $F$ is 96,485 C/mol, and $Q$ is the total charge passed.

3.3. Catalyst Characterization

XRD tests were performed on a D8-Foucas (BRUKER AXS GMBH, Karlsruhe, Germany) equipped with a Cu Kα radiation ($\lambda = 0.15418$ nm). The spectra were collected between 2θ ranges of 30–80°, at a scanning speed of 8°/min.

To compare the crystal facet ratio of different Zn catalysts, texture coefficients [29] were calculated by Equation (7).

$$\text{TC}_{hkl} = \frac{I_{(hkl)} / I_{0(hkl)}}{\sum_{i=1}^{n} I_{(hkl)} / I_{0(hkl)}} \times 100\%$$  \hspace{1cm} (7)

where $I_{(hkl)}$ is the diffraction intensity of the $(hkl)$ facet and is automatically calculated by the Jade software, $I_{0(hkl)}$ is the standard diffraction intensity of the $(hkl)$ and taken from JCPDS#04-0831, and $n$ is the number of diffraction peaks of Zn ($n = 7$).

The TEM images of Zn catalysts were tested by a JEM2100-F field-emission transmission electron microscope (JEOL, Tokyo, Japan). The field SEM images of Zn catalysts were tested by a Regulus 8100 instrument (JEOL, Tokyo, Japan) at a 3.0 KV electron beam. The surface chemical states of the Zn electrode were investigated by XPS (ULVACPHI, Japan) with monochromatized Al Kα X-ray radiation (1486.6 eV).

4. Conclusions

In summary, a series of Zn catalysts were prepared by electrodeposition in different gas conditions (N$_2$, CO$_2$, H$_2$, CO). The Zn-CO$_2$ catalyst obtained with porous fibers and uniform tiny pores exhibited high activity and selectivity for CO$_2$RR to CO, with FE$_{CO}$ for 73.0% and FE$_{H2}$ for 32.6% at −1.2 V vs. RHE. The partial current density of CO was 6.14 mA/cm$^2$ and the FE$_{CO}$ was almost unchanged over 6 h in −1.2 V vs. RHE. The electrokinetic study results show that the initial electron transfer step was the RDS for the Zn-CO$_2$ electrode. The presence of CO$_2$ affected the crystal growth of electrodeposited Zn, increased the proportion of the Zn (101) crystal on the Zn electrode surface, and induced the formation of a fibrous and porous structure. The structure increased the ECSA and under-coordinated Zn$^{2+}$ species. The greater ECSA provided more active sites and promoted the production of CO. This was the main reason for the excellent performance of electrochemical CO$_2$RR for the Zn-CO$_2$ electrode. This study demonstrates that the morphology and crystal structure of Zn catalysts have a great influence on the activity for
CO₂RR. This work is of great value and significance to solve the challenge of low activity and poor stability in electrochemical CO₂RR.

Supplementary Materials: The following are available online at https://www.mdpi.com/article/10.3390/catal111040477/s1, Figure S1: Current density and FE CO for (a) Zn foil, (b) Zn-N₂, (c) Zn-H₂ and (d) Zn-CO electrodes during 6 h of long-time operation at −1.2 V vs. RHE; Figure S2: CV curves on (a) Zn foil, (b) Zn-N₂, (c) Zn-CO₂, (d) Zn-H₂ and (e) Zn-CO electrodes with a potential range from −0.415 to −0.515 V vs. RHE in a N₂ bubbled 1 M Na₂SO₄ electrolyte; Table S1: The data of CO partial current density for Zn-CO₂ electrode; Table S2: The solubility data of different gases in water.

Author Contributions: M.G. wrote the manuscript, performed the experiments and analyzed data; M.G., X.L. and Y.H. collected references and characterized the physico-chemical properties of materials; L.L. and J.L. collected and checked data; Y.X. and Y.L. analyzed data and checked the manuscript; L.Z. provided research ideas, analyzed data and revised the manuscript. All authors have read and agreed to the published version of the manuscript.

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