Title: Isolated Zn-Co Atomic Pair for Highly Active and Durable Oxygen Reduction

Authors: Ziyang Lu, Bo Wang, Yongfeng Hu, Wei Liu, Yufeng Zhao, Ruouo Yang, Zhiping Li, Jun Luo, Bin Chi, Zheng Jiang, Minsi Li, Shichun Mu, Shijun Liao, Jiujun Zhang, and Xueliang Sun

This manuscript has been accepted after peer review and appears as an Accepted Article online prior to editing, proofing, and formal publication of the final Version of Record (VoR). This work is currently citable by using the Digital Object Identifier (DOI) given below. The VoR will be published online in Early View as soon as possible and may be different to this Accepted Article as a result of editing. Readers should obtain the VoR from the journal website shown below when it is published to ensure accuracy of information. The authors are responsible for the content of this Accepted Article.

To be cited as: Angew. Chem. Int. Ed. 10.1002/anie.201810175
Angew. Chem. 10.1002/ange.201810175

Link to VoR: http://dx.doi.org/10.1002/anie.201810175
http://dx.doi.org/10.1002/ange.201810175
Isolated Zn-Co Atomic Pair for Highly Active and Durable Oxygen Reduction


Abstract: N-coordinated nonprecious dual metal catalysts have demonstrated superior catalytic activity by offering more concentrated active sites. However a rational design of such system remains conceptually challenging which requires in-depth research both experimentally and theoretically. Herein, we develop a competitive complexation strategy to construct a novel electrocatalyst with Zn-Co atomic pairs coordinated on N doped carbon support (Zn/CoN-C). Such architecture offers enhanced binding ability of O₂, significantly elongates the O-O length (from 1.23 Å to 1.42 Å), and thus facilitates the cleavage of O=O bond, showing a theoretical overpotential of 0.335V during ORR process. As a result, the Zn/CoN-C catalyst exhibits outstanding ORR performance in both alkaline and acid conditions with a half-wave potential of 0.861 and 0.796 V respectively. The assembled zinc-air battery with Zn/CoN-C as cathode catalyst presents a maximum power density of 230 mW cm⁻² along with excellent operation durability. The excellent catalytic activity in acid is also verified by H₂/O₂ fuel cell tests (peak power density of 705 mW cm⁻²).

Electrochemical oxygen reduction reaction (ORR) has gained widespread attentions due to its significance in electrochemical energy storage and conversion including proton-conducting membrane fuel cells or metal-air batteries. Platinum-based catalysts are generally efficient for ORR, but their wider applicabilities are restricted by its high price, and limited stability. Replacement of noble metal materials with less expensive, highly active and durable electrocatalysts for ORR is thereby increasingly attractive but arduous with great challenges ahead.

Nonprecious metal catalysts with M-Ni-C (M stands for nonprecious metal elements) coordination, represent a new type of effective catalysts as possible alternatives to Pt. Recent study found that, achieving atomic-level regulation of metal atoms is critical to designing high catalytic active catalysts. Isolated single-atom iron or cobalt catalysts supported on carbon materials have demonstrated unexpected ORR reactivity than commercial Pt/C. Follow-up study revealed that, catalysts with bimetallic or multimetallic active sites can further enhance the catalytic activity and selectivity in numerous catalytic conversion reactions. A recently reported Pt free catalyst with N-coordinated Fe-Co dual sites delivers outstanding activity for ORR in acidic electrolyte. Compared to the single metallic site, the structure formed by the coordination of dual-metal-atom with nitrogen atoms is more favorable for cracking O₂ bond. Nevertheless, the current research efforts about M-Ni-C catalysts mainly focused on Ni, Co or Fe/N-doped carbon materials catalysts. Exploring new metal components and new catalytic site may be an effective way to increase catalytic activity under the premise of ensuring high dispersion. In nature, besides Fe and Co porphyrin, Zn porphyrin structure can also be found in carbonic anhydrase with four-coordinated structure. In view of that, Zn has a smaller electronegativity (χ=1.65) compared to Fe (χ=1.83) and Co (χ=1.88) which makes it easier to donate its outer electron. When atomic pairs formed between Zn and Co or Fe, the interactions between the electronic structures might result in a more optimized structure, which makes it possible for the adsorption and desorption of the reaction intermediate to reach a suitable state, so that the diatomic pair has a higher catalytic activity. Thereby, introducing more metal species to construct bimetallic or multimetallic active sites are challenging yet of great significance to achieve best performing catalysts.

Herein, we report a new design of discrete Zn, Co bimetallic sites supported on N doped carbon through a competitive complexation strategy, whereby chitosan was used as C and N sources, zinc chloride, and cobalt acetate are selected as metal precursors (Figure 1a). The 4s and 4p electrons of Co²⁺ or Zn²⁺ can be coordinated with the -NH₂ and –OH group on chitosan chains through a simultaneous competitive complexation process, thus allowing the homogeneous dispersion of Zn/Co. And due to their similar coordination abilities, a bimetallic...
Zn/CoN\textsubscript{2}-C nanostructure with atomic dispersion is finally achieved after pyrolysis at 750°C, which is denoted as Zn/CoN\textsubscript{C}. The weight percentage of Zn, Co is determined as 0.33 wt% and 0.14 wt% respectively, by inductively coupled plasma mass spectrometry (ICP-MS). This synthetic method allows scalable synthesis of catalysts with good reproducibility.

In order to confirm the existential form of Zn and Co dual atoms, we carried out aberration-corrected atomic-resolution high-angle annular dark field scanning transmission electron microscopy (HAADF-STEM) measurements (Figure 1b). Small bright spots are evenly distributed in carbon substrate, which are attributed to the relatively high atomic numbers of Zn and Co. Further enhancing the magnification, small bright dual dots can be observed more clearly (Figure 1c), in which Zn/Co atomic pair can be recognized by bright spots marked with red circles. Some individual Zn or Co atoms can be distinguished and marked with orange circles. Statistical analysis of more than 40 pairs of dual atoms showed a Zn-\text{Co} distance of 0.22 ± 0.04 nm (Figure 1d). The well dispersed Zn-Co dual-sites can be clearly identified in accessional intensity profiles (Figure 1d insets). HAADF-STEM (Figure 1e) and elemental mapping (Figure 1f and S1) reveal that N, Zn and Co are homogeneously distributed in the ZnCoN\textsubscript{C}. The (HR) TEM image (Figure 1g) demonstrates lattice distortion defects characteristic, which might be attributed to the coordination of single or dual zinc/cobalt atoms with nitrogen. For comparison, single-metal atom Co or Zn doped catalysts, denoted as CoN\textsubscript{C} and ZnN\textsubscript{C} were also synthesized through similar approaches.

X-ray diffraction (XRD) patterns of the samples (Figure S2) confirm the non-formation of metallic and oxide crystals on carbon supports. Nitrogen adsorption measurement illustrates the high BET (Brunauer-Emmet-Teller) specific surface area (1343 m\textsuperscript{2}/g) of ZnCoN\textsubscript{C} with an average pore width of 2.46 nm (Figure S3). Raman spectra (Figure S4) and XPS spectrum of C1s (Figure S5) suggest that more defects are produced due to the co-coordination between Co and Zn and the N doped carbon support, which plays an important role in the electronic properties of carbon matrix. As shown in Figure S6, N1s spectrum for ZnCoN\textsubscript{C} can be deconvoluted into four peaks, pyridinic N (398.3 eV), N-Zn/Co (399.7 eV), pyrrolic N (400.6 eV), and graphitic N (401.4 eV), respectively. The existence of N-Zn/Co can also be confirmed at Zn 2p spectrum (Figure S7b), and Co 2p\textsubscript{1/2} signals at 778.7 eV. (Figure S7c).

X-ray absorption near-edge structure (XANES) and extended X-ray absorption fine structure measurements (EXAFS) were conducted to investigate the chemical state and coordination environment of Co and Zn atoms in the catalysts. As shown in the XANES spectra (Figure 2a), the absorption edge of Co K-edge of CoN\textsubscript{C} shifted 3.0 eV towards lower energy relative to CoO, indicating that the valence state of CoN\textsubscript{C} is situated between that of Co\textsuperscript{2+} and Co\textsuperscript{3+}. Furthermore, from the EXAFS spectra of different samples (Figure 2b), a main peak at about 1.5 Å can be observed for both cobalt phthalocyanine (CoPc) and CoN\textsubscript{C}, which is typically assigned to the Co-N coordination, demonstrating the confined Co-N coordination environment in CoN\textsubscript{C}. The Co K-edge EXAFS spectrum of ZnCoN\textsubscript{C} is characterized by three peaks at 1.4 Å, 2.1 Å and 2.4 Å, corresponding to the nearest Co-N, next nearest Co-Metal and long range Co-C coordination, respectively (Figure 2b). From its corresponding Zn K-edge XANES spectrum (Figure 2c), we can see that both the edge position and the peak intensity of ZnCoN\textsubscript{C} located between Zn foil and ZnO, which suggests the electronic state of the Zn species is between Zn\textsuperscript{2+} and Zn\textsuperscript{0}. On the other hand, Zn K-edge EXAFS (Figure 2d) analysis clearly indicates that ZnCoN\textsubscript{C} and zinc phthalocyanine show a similar main peak at 1.4 Å, which could be attributed to Zn-N scattering paths. The two peaks at 2.1 Å and 2.4 Å, should be due to the next nearest Zn-Metal and long range Zn-N coordinations, respectively.

The fitting results of ZnCoN\textsubscript{C} can better reveal the structural information of the material (Figure S8). The Co-N coordination number is 3.5 for ZnCoN\textsubscript{C} suggest that the catalyst was dominated by the Co-N interaction with a mixed Co-N3 and Co-N4 environment. The second shell coordination number of Co-M is given by CN_{Co-M}=0.5±0.1, suggesting a weak Co-M interaction in form of ZnCoN6 structure. Similarly for Zn of the ZnCoN\textsubscript{C}, it is dominated by the Zn-N coordination (N=3.5) in the first shell, together with a weak second shell due to the Zn-M (CN_{Zn-M}=0.5±0.1), further demonstrating the existence of ZnCoN6 structure in ZnCoN\textsubscript{C}. This result is also consistent with the DFT calculations that Zn and Co are bonded, the system has the lowest energy, therefore Zn and Co tend to form Co-Zn diatomic pairs (Figure S9).
commercial Pt/C (0.45V), suggesting the high ORR catalytic activity of ZnCoN-C.

The ORR catalytic activity of ZnCoN-C was investigated in O2-saturated 0.1M KOH with rotating disk electrode (RDE). At 750 °C the ZnCoN-C exhibit the optimal electrocatalytic performance as shown in Figure S10. And Cyclic voltammetry (CV) reveals a significant reduction peak at 0.84V for ZnCoN-C (Figure S11), suggesting a good ORR electrocatalytic activity. The linear sweep voltammetry (LSV) measurements (Figure 3a) show that ZnCoN-C presents the best oxygen reduction activity, with a much more positive E onset (1.004 V) than Pt/C (0.970 V) and a high E 1/2 of 0.861V, which is 35 mV more positive than Pt/C. This E 1/2 value also surpasses those of CoN-C (0.793V) and Zn-N-C (0.760V) (Figure S3b). In addition, ZnCoN-C exhibits a more active J, of 6.1 mA/cm², and this value is far better than those of Pt/C (5.3 mA/cm²), CoN-C (4.7 mA/cm²) and Zn-N-C (4.4 mA/cm²). The catalytic activities of ZnCoN-C with the optimal Zn-Co atomic ratio (Figure S12) surpass those of most non-precious metal ORR electrocatalysts (Table S1). The electron transfer number of ZnCoN-C was also evaluated from the LSVs (Figure 3c) according to Koutecky-Levich (K-L) equation, giving the n value of 3.88. The n and H2O2 yield were further evaluated through rotating ring-disk electrode (RDE, Figure 3d), whereby the direct four electron process is confirmed, with n approaching 4 and H2O2 yield of about 5%. Tafel plots demonstrate much smaller slope of 67 mV dec⁻¹ of ZnCoN-C, indicating the superior reaction kinetics (Figure 3e). The durability of ZnCoN-C was evaluated by CV at a sweep rate of 100 mV/s. After 5000 continuous cycles (Figure 3f), there was 5 mV change in E 1/2, and after 10000 cycles, there was only 9 mV change in E 1/2, demonstrating that the ZnCoN-C had the superb stability. And the Zn-Co atomic pairs of the ZnCoN-C are still well preserved after 10000 cycles of testing, which is verified by HAADF-STEM (Figure S13) and XANES (Figure S14). Furthermore, the in situ XANES spectra of ZnCoN-C (Figure S15) suggests that Co is more actively involved, and function as active center during the ORR, which likely undergoes a slight redox switch, in agreement with the FeN4 system. Upon the closer inspection of the Co K-edge result, the Co K-edge is slowly shifting to higher energy as the potential increases. After the oxygen reduction process, the structure of the ZnCoN-C has not changed substantially whether for Co K-edge or Zn K-edge, during the whole ORR process which guarantees the high oxygen reduction activity. The fuel crossover effect evaluation of ZnCoN-C demonstrates excellent tolerance to methanol (Figure S16) and SCN⁻ ions (Figure S17).

A primary Zn-air battery was assembled to evaluate the practical application of ZnCoN-C. Significantly, a maximum power of 230 mW cm⁻² is achieved from the battery using ZnCoN-C catalyst...
Figure 3. (a) ORR Linear scan voltammogram (LSV) curves for different catalysts in O₂ saturated 0.1 M KOH solution. (b) Limit current density and half-wave potential for different catalysts. (c) LSV curves for ZnCoN-C at different rotating rates from 400 to 2000 rpm and the inset is K-L plots at various potentials. (d) Electron transfer number and H₂O₂ yield for different catalysts. (e) Corresponding Tafel plots obtained from the RDE polarization curves. (f) The durability tests of ZnCoN-C.

Furthermore, no significant potential change was detected at a current density of 5 mA cm⁻² for 100,000 s (Figure 4b), indicating a stable practical application performance. As shown in Figure 4b inset, two series-connected Zn-air batteries based on the ZnCoN-C as cathode can power a series of red light-emitting diode (LED) in parallel (LED, 2.2 V) with an outstanding stability.

Promisingly, the ZnCoN-C also presents outstanding ORR activity in acidic conditions. In 0.1M HClO₄, ZnCoN-C shows a considerably high ORR catalytic activity with E₁/₂ of 0.796 V and Eᵢₚₛₛ of 0.97 V, which is comparable to that of Pt/C (Figure 4c), and superior to most of non-precious metal ORR electrocatalysts (Table S3). And after 10,000 cycles of CV, there was no obvious change in E₁/₂, demonstrating the excellent stability of ZnCoN-C (Figure S19). In the SCN⁻ poisoning test (Figure S20), the E₁/₂ of ZnCoN-C negatively shifted by 30 mV and the Eᵢₚₛₛ remained unchanged after adding SCN. This apparent resistance to SCN⁻ poisoning can be attributed to the binmetallic coordination in the ZnCoN₅ structure that regulates the interfacial electron structure, which is detrimental to the binding to KSCN. These test results clearly show that dual-metal sites ZnCoN₅ are the major active sites. A H₂/O₂ fuel cell was assembled to verify the practical application potential in acidic solution. As a result, a kinetic current of 999 mA cm⁻² is achieved at 0.6 V and a peak power density of 705 mW cm⁻² at 0.5 V, which is ~78% of that based on commercial Pt/C (Figure 4d). Furthermore, after eight hours of stability testing, there was no attenuation of activity (Figure S21).

In summary, this work demonstrates that the Zn/Co dual atoms sites can serve as efficient ORR catalyst with outstanding activity both experimentally and theoretically. The Zn/Co dual atoms sites were confirmed by a combination of aberration-corrected HAADF-STEM, XAFS and DFT calculations. DFT calculations show that ZnCoN₅ site is the main active site and the theoretical overpotential of oxygen reduction can be as low as 0.335 V. In addition, the catalytic activity can be maintained even under acidic environment. Furthermore, the assembled Zn-air batteries and fuel cell with ZnCoN-C as catalyst show excellent power density and stability, which demonstrates the practical application of the as-prepared ZnCoN-C catalyst. This work may represent a new class of dual atoms sites catalysts for both fundamental research and practical applications.

Acknowledgements
We thank the financial supports from NSFC (51774251), HB NSDFY (B2017203313), HEITSP (SLRC2017057), and National Program for Thousand Young Talents of China. The XAS measurement was conducted at the Canadian Light Source (CLS). CLS is supported by the NSERC, NRC, CIHR of Canada, the University of Saskatchewan, and the BL 14W beamline at the Shanghai Synchrotron Radiation Facility (SSRF).

Keywords: isolated atomic pair, M-N₅-C, oxygen reduction reaction, Zn-air battery


The discrete zinc, cobalt bimetallic sites supported on N doped carbon (Zn/CoN-C) through a competitive complexation strategy. The as-prepared Zn/CoN-C catalyst exhibits outstanding ORR performance in both alkaline and acid conditions, and the assembled Zn-air batteries and H₂/O₂ fuel cells show excellent power density and stability which demonstrate the practical applications.