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Hydrophilic non-precious metal nitrogen-doped carbon electrocatalysts for enhanced efficiency in oxygen reduction reaction†

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Exploring the role of surface hydrophilicity of non-precious metal N-doped carbon electrocatalysts in electrocatalysis is challenging. Herein we discover an ultra-hydrophilic non-precious carbon electrocatalyst, showing enhanced catalysis efficiency on both gravimetric and areal basis for oxygen reduction reaction due to a high dispersion of active centres.

The oxygen reduction reaction (ORR) is a fundamental electrochemical reaction for fuel cells and metal-air batteries. ORR research has long been focused on the development and understanding of new non-precious nitrogen-doped carbon catalysts that result in significant cost reduction by platinum metal substitution. To catalyze broader commercialization of such devices and technologies, efficient and affordable non-precious electrocatalysts will ultimately be required. One of the most prominent examples is pyrolyzed solid materials consisting of non-precious metal/nitrogen/carbon (M/N/C-) composites,¹ or metal-free, heteroatom-doped nanocarbons.² The nature of the active sites in terms of the modulation of electron donating/withdrawing capability of the carbon basal plane by incorporated heteroatoms of the M/N/C electrocatalysts has been under intensive investigation and has become more and more clear.³ However, a detailed understanding of the effect of surface

hydrophilicity and wettability on the dispersion of metal-related active sites as well as their effects on the catalysis efficiency of the M/N/C materials has remained elusive.

On the one hand, a hydrophilic pore surface benefits a facile loading and a high dispersion of active metal-related species, because hydrophilic affinity can be created between hydrophilic pore walls and precursors, and further inhibits precursor random migration and agglomeration.⁴ On the other hand, a hydrophilic pore surface may also influence the transport of hydrated O₂ to the electrochemically active centres under hydrated conditions, finally affecting the activity.⁵ To our knowledge, such effects stemming from surface hydrophilicity have been rarely investigated to date.

Taking these into consideration, in this contribution, we surface engineered a number of different carbon-based materials with surface characteristics ranging from an ultra-hydrophilic carbon network to an ultra-hydrophobic carbon black. We observe that hydrophilicity, quantified by water adsorption isotherms at 298 K, is correlated with much enhanced ORR catalysis efficiency.

Firstly, a group of ultra-hydrophilic electrocatalysts (Fe/N_{1/3.2}, Fe/Cu/N_{1.3/1/8}, and Cu/N_{1/4}, according to the nominal atomic ratio of Fe/N, Fe/Cu/N and Cu/N in the synthesis, Fig. 1a–c) were fabricated by a facile and scalable impregnation and subsequent pyrolysis and leaching method (Experimental section, ESI†) based on an unprecedentedly hydrophilic carbon network (DUT-110, DUT = Dresden University of Technology, derived from a functional complex⁶). The highly hydrophilic surface property of DUT-110 was confirmed by the sharp uptake in the water vapor adsorption isotherm from the very beginning, showing a record value until $P/P_0 < 0.3$ (Fig. S1a, ESI†).⁷ The narrow and rich micropores were proven by the type I isotherm and pore size distribution based on N₂ physisorption data (Fig. S1b, ESI†) as well as high resolution TEM images (Fig. S1c, ESI†); while the STEM mapping and XPS spectrum revealed the highly heteroatom-doped feature, with a surface composition of C, N, and O with the atomic content of 74.7, 14.3, and 10.33%, respectively (Fig. S1d and e, ESI†). In parallel, hydrophobic non-precious electrocatalysts

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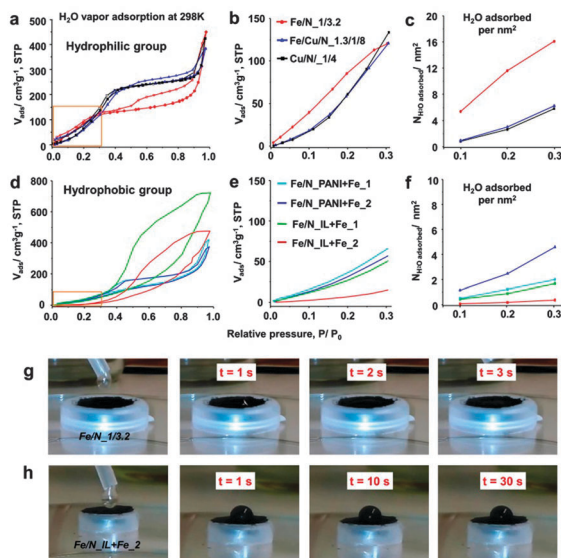


Fig. 1 H_2O vapor adsorption isotherm at different relative pressure range at 298 K of hydrophilic groups (a–c) and hydrophobic groups (d–f). (g and h) Comparison of dynamic water contact angle changes of typical samples of the two groups.

were prepared under the same principle through the modification of N-containing polymers (polyaniline, PANI or N-containing ionic liquid, N,N -ethyl-methyl-imidazolium-dicyanamide) and FeCl_3 , but based on highly hydrophobic carbon black (Ketjen EC 600J). Note that the pyrolysis and leaching treatment as well as the following performance evaluation were kept identical with those of the hydrophilic groups.

Surprisingly, the final hydrophilic electrocatalysts, $\text{Fe/N}_{1/3.2}$, $\text{Fe/Cu/N}_{1.3/1/8}$, and $\text{Cu/N}_{1/4}$, maintained largely the high surface hydrophilicity. They all exhibit relatively good hydrophilic properties, with the water uptake of 120.4, 120.5, and $133.6 \text{ cm}^3 \text{ g}^{-1}$ (equivalent to 5.37, 5.38, and 5.96 mmol g^{-1}) at $P/P_0 = 0.3$ for $\text{Fe/N}_{1/3.2}$, $\text{Fe/Cu/N}_{1.3/1/8}$, and $\text{Cu/N}_{1/4}$, respectively (Fig. 1a). The water adsorption behavior in the range of $0 < P/P_0 < 0.3$ is mainly determined by the surface hydrophilicity;⁷ thus, we further compared the water sorption uptake in this pressure range (Fig. 1b). All the hydrophilic samples, *i.e.*, $\text{Fe/N}_{1/3.2}$, $\text{Fe/Cu/N}_{1.3/1/8}$, and $\text{Cu/N}_{1/4}$, exhibit a higher water adsorption uptake, even though their less developed porosity and low surface area ($200\text{--}574 \text{ m}^2 \text{ g}^{-1}$, Fig. S2, ESI†) were comparable to hydrophobic materials ($340\text{--}1079 \text{ m}^2 \text{ g}^{-1}$, Table S1, ESI†). After normalization to specific surface area (Fig. 1c), the hydrophilicity order is $\text{Fe/N}_{1/3.2} > \text{Fe/Cu/N}_{1.3/1/8} > \text{Cu/N}_{1/4}$. For instance, the water adsorption uptake is calculated to be $16.2 \text{ H}_2\text{O}$ molecules per nm^2 , $6.7 \text{ H}_2\text{O}$ molecules per nm^2 and $6.2 \text{ H}_2\text{O}$ molecules per nm^2 at $P/P_0 = 0.3$ for $\text{Fe/N}_{1/3.2}$, $\text{Fe/Cu/N}_{1.3/1/8}$, and $\text{Cu/N}_{1/4}$, respectively. However, the water adsorption isotherm of the hydrophobic electrocatalysts displays much lower water adsorption uptakes, indicating a much lower surface hydrophilicity (Fig. 1d–f). To illustrate the difference in hydrophilicity vividly, the water contact angles were recorded dynamically after water droplets contacted the carbon pellet and were compared (herein we show the samples of $\text{Fe/N}_{1/3.2}$ and $\text{Fe/N}_{\text{IL}} + \text{Fe}_2$

in Fig. 1g and h). For hydrophilic $\text{Fe/N}_{1/3.2}$, the water droplet can be adsorbed in 3 s with a final contact angle of *ca.* 0, while the sample of $\text{Fe/N}_{\text{IL}} + \text{Fe}_2$ was not wetted until 30 s, again confirming the distinct surface hydrophilicity. Morphologically, all the hydrophilic electrocatalysts exhibit a highly interconnected network structure (Fig. S3, ESI†), but a much denser structure compared with their host carbon networks DUT-110 (Fig. S1d, ESI†). This is due to the shrinkage of the carbon skeletons during high temperature pyrolysis.

Based on the unique hydrophilicity and narrow micropores, a high dispersion of metal-related nanoparticles was expected for hydrophilic electrocatalysts. Thus, the elemental distribution was analysed by STEM images and elemental maps as well as dark-field TEM images (Fig. 2). For $\text{Fe/N}_{1/3.2}$ (Fig. 2a), large and bright particles can be observed, while for $\text{Fe/Cu/N}_{1.3/1/8}$, only very few isolated particles can be detected (Fig. 2b). This observation is consistent with the TEM images (Fig. S4, ESI†). Importantly, except some large particles, the distribution of small metal species is very homogeneous, indicating highly dispersed nanoclusters or even mononuclear metal species embedded in the carbon matrix. The dark-field TEM images (Fig. 2e and f), corresponding to their relevant TEM images (Fig. 2c and d), again confirmed a high and uniform distribution of metal-related nanoclusters over the whole framework. The metal content determined by the ICP technique is 3.8 wt% Fe for $\text{Fe/N}_{1/3.2}$ and 1.22 wt% Fe for $\text{Fe/N}_{1/3.2}$. Note that these metal nanoclusters should be tightly embedded in the carbon matrix, since all these electrocatalysts have been extensively leached in 2.0 M H_2SO_4 at 110°C for 24 h before harvesting for characterisation and application. Furthermore, the hybrid structure composed of graphitic domains and amorphous carbons was revealed by Raman spectra (Fig. S5, fitting details in Table S2, ESI†) with the I_D/I_G ratio of 1.55, 1.82 and 2.11 for $\text{Fe/N}_{1/3.2}$, $\text{Fe/Cu/N}_{1.3/1/8}$ and $\text{Cu/N}_{1/4}$, respectively, confirming the observation by TEM images (Fig. S4, ESI†).

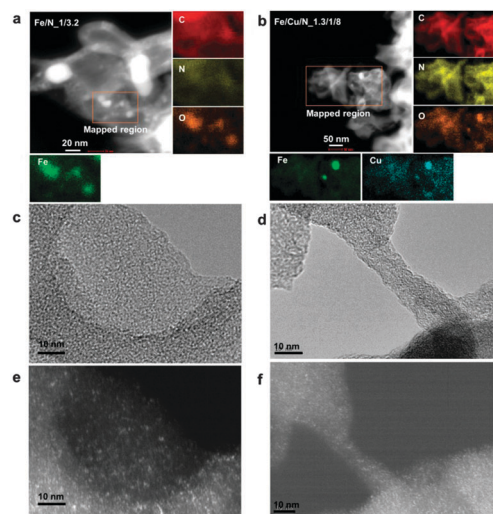


Fig. 2 STEM maps of $\text{Fe/N}_{1/3.2}$ (a) and $\text{Fe/Cu/N}_{1.3/1/8}$ (b), TEM image and the corresponding dark-field TEM images of $\text{Fe/N}_{1/3.2}$ (c and e) and $\text{Fe/Cu/N}_{1.3/1/8}$ (d and f).



For non-metal elements such as C, O, and N, a homogeneous dispersion was also observed by elemental maps (Fig. 2a and b), indicating a uniform doped structure. Interestingly, comparing the Fe or Cu maps with O maps, a strong correlation is found between the metal and O, particularly for large particles, indicating their oxide phase in nature. This observation is further confirmed by their XPS analysis (Fig. S6, ESI†). Furthermore, Table S1 (ESI†) also lists other structural parameters such as the specific surface area analyzed by N₂ adsorption, the surface non-metal compositions determined by XPS and the metal species detected by ICP for all the hydrophilic and hydrophobic catalyst groups in order to obtain a reliable correlation between the structural parameters and the subsequent catalysis performance.

We first evaluated the ORR activity of the hydrophilic group, *i.e.*, Fe/N_1/3.2, Fe/Cu/N_1.3/1/8, and Cu/N_1/4. Linear sweep voltammetry (LSV, Fig. 3a) in 0.10 M KOH was employed to investigate the catalytic activity of the catalysts compared to Pt/C benchmark catalysts. The onset potentials (E_{onset} , a noteworthy onset potential is defined as the potential at which the current density reaches 1.0 mA cm⁻²) are 0.90, 0.92, and 0.89 V for Fe/N_1/3.2, Fe/Cu/N_1.3/1/8, and Cu/N_1/4, respectively (Fig. 3a, Table S1, ESI†). The E_{onset} of Fe/Cu/N_1.3/1/8 is positive and comparable with those of the reported state-of-the-art non-precious catalysts such as Fe/N-doped nanocarbons (*e.g.* N-CNT/Fe₃C, N-doped carbon nanoplate/Fe₃C, Fe@Fe₃C/N-doped carbon),⁸ Fe and/or N-doped porous carbons with higher surface area or larger pores such as mesopores⁹ or hierarchical pores,¹⁰ and the hybrid N-Fe-CNT/carbon nanoparticle with higher Fe content.¹¹ The half-wave potential ($E_{1/2}$) shows a similar trend that is also comparable with that of the state-of-the-art non-precious

electrocatalysts (Table S3, ESI†).^{9–12} Particularly, the high activity reflected by the positive E_{onset} and $E_{1/2}$ of Fe/Cu/N_1.3/1/8 originates from the highly dispersed active sites and highly accessible porosity.

The mass activity indicates the utilization efficiency of catalysts on a gravimetric basis. For the hydrophilic series, relatively higher mass activities up to 413.3, 232.5, and 137.7 mA mg⁻¹ were calculated for Fe/N_1/3.2, Fe/Cu/N_1.3/1/8, and Cu/N_1/4, respectively (Fig. 3b). For the hydrophobic samples, the mass activity is one order of magnitude lower (Fig. 3b). The much higher mass activity of the hydrophilic samples originated from the positively shifted onset and half-wave potentials, indicating the large density of accessible active sites due to the high dispersion of electrochemically active sites benefiting from the highly hydrophilic carbon surface. In order to understand quantitatively, we further normalized the mass activity by the surface area. The obtained specific activities are 2067, 482.4, and 239.9 mA m⁻² for Fe/N_1/3.2, Fe/Cu/N_1.3/1/8, and Cu/N_1/4, respectively. This trend is also consistent with that of mass activity, reflecting the remarkably high surface efficiency (Fig. S7 and Table S1, ESI†).

Moreover, we correlated the Fe and N content, and the surface area as well as surface hydrophilicity with mass activity. However, it is difficult to find a clear trend between either mass or specific activity and doping properties (Fig. 3c, black and grey lines, Fig. S8, ESI†) or specific surface area (Table S1 and Fig. S9, ESI†). In contrast, a clear correlation between mass activity and surface hydrophilicity was observed. This is probably because (1) the higher surface hydrophilicity induces a higher dispersion of active sites, and (2) hydrophilic pores benefit an easy accessibility to the active sites of reactants (such as hydrated O₂ as ORR proceeds). Besides the comparison between specific samples, a comparison between the hydrophobic group and the ultra-hydrophilic group confirmed the same rule (Fig. 3c, red trend line).

The high dispersion of active sites benefiting from surface hydrophilicity has been proven above by STEM mapping and dark field TEM images. However, to observe the effect of hydrophilicity on the diffusion of hydrated O₂ and resultant H₂O is challenging. For the diffusion of hydrated O₂ near reaction interfaces, high surface hydrophilicity may be beneficial.^{5b–d} In order to explain this point, we hypothesized a “physical structure” (Fig. S10, ESI†), where water molecules around hydrated O₂ molecules can be readily stripped by the hydrophilic micropore walls when approaching the carbon slit pores, then the liberated O₂ molecules can freely diffuse to the active sites and thus can accelerate the reaction. The O₂ adsorption was investigated for the active hydrophilic group, which can give the first clue that high surface hydrophilicity enhanced the O₂ adsorption (Fig. S11, ESI†). All hydrophilic samples exhibit a combined type I isotherm (Fig. S9a and b ESI†), indicating a strong interaction between O₂ molecules and doped carbon pore walls. After normalization by the specific surface area, the samples showed a higher areal uptake, indicating the preferential O₂ adsorption and high surface utilization efficiency for trapping O₂ molecules (Fig. S11c, ESI†). Interestingly, the O₂ capture behaviour is consistent with that of water sorption in the same pressure range.

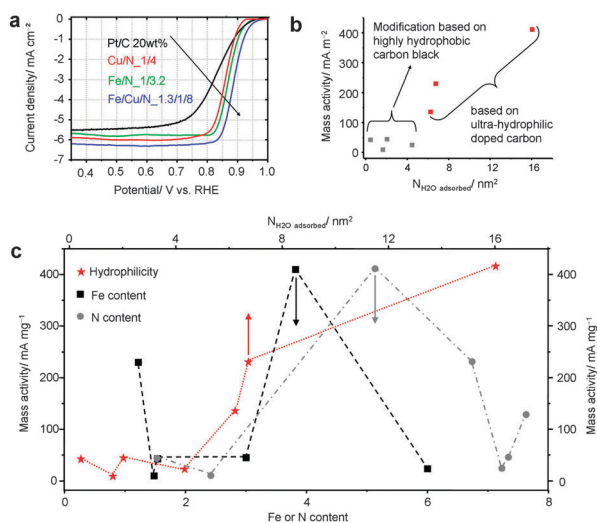


Fig. 3 ORR catalysis evaluation in O₂-saturated 0.10 M KOH. (a) Linear sweep voltammetry (LSV) ORR plots under conditions of room temperature, at a rotation speed of 1500 rpm, scan rate of 10 mV s⁻¹, the non-precious catalysts loading of 0.80 mg cm⁻², and the benchmark Pt loading of 10 μg cm⁻². (b) The relationship between hydrophilicity in terms of water molecules adsorbed per nm² based on water adsorption data at $P/P_0 = 0.3$ and mass activity. (c) Structure–performance comparison including surface hydrophilicity, nitrogen content (atomic%, by XPS) and Fe content (wt%, by ICP) with mass activity in ORR electrocatalysis under identical conditions.



In addition, one would realize that the high surface hydrophilicity of the catalysts may also cause a delay of water desorption when used in applications that generate water such as alkaline fuel cells (AFCs). In this case, further H₂ reduction can be applied to effectively reduce the surface hydrophilicity (Fig. S12, ESI†). Through this way, the possible flooding issues can be avoided in potential applications such as AFCs.

In summary, exemplified using the non-precious carbon based ORR electrocatalyst concept, we surface engineered a number of different carbon based materials with surface characteristics ranging from an ultra-hydrophilic carbon network to ultra-hydrophobic carbon black. A high surface hydrophilicity has been found to give an easily wetted surface which first ensures a high dispersion of metal-related active sites and may also increase the accessibility of reactants to active centres, and thus may increase the surface and mass utilization efficiency of catalysts. This work provides fresh insight into the controlling material parameters of non-precious ORR catalysts, and as such offers new clues and strategies on how to increase the ORR catalysis efficiency by tuning the surface chemistry of non-precious electrocatalysts. The insight into hydrophilicity may also be important to other heterogeneous catalytic reactions such as CO₂ electro-reduction, glucose oxidation, and metal–air batteries, catalyzed on hydrated carbon surfaces.

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