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Graphene-supported Pt and PtPd Nanorods with Enhanced Electrocatalytic Performance for Oxygen Reduction Reaction

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The combinational modification of the morphology, alloying, and support for Pt catalyst has been optimized towards oxygen reduction reaction. Graphene-supported PtPd nanorods have lower unfilled Pt d-states than carbon-supported Pt nanoparticles (Pt/C) and their specific and mass activities after accelerated durability test is about 6.5 and 2.7 times higher than those of Pt/C, attributed to the synergistic electronic modification effect and graphene-metal interaction.

The development of highly effective and stable electrocatalysts toward oxygen reduction reaction (ORR) has attracted lots of attention recently.1 Carbon-supported Pt nanomaterials with different alloying components and morphologies especially nanorods (NRs) or nanowires (NWs) have shown enhanced ORR performance when compared with Pt nanoparticles (NPs).2 The ORR promotion for Pt or Pt-based 1-dimensional (1-D) nanomaterials may be related to their larger surface contact with the carbon support than the polyhedral NPs, better electron conductivity between the reaction sites and the carbon support, preferential exposure of certain crystal facets and less surface defects bearing.2,3 Besides, recent studies have found that the electronic and chemical states of Pt NPs can be changed through the formation of 1-D structure and alloying effect, thus modifying the oxophilicity of Pt and enhancing the ORR performance.2b,2c

Another approach for the optimization of the cathode catalysts is to use graphene as support. Because of high surface area, high conductivity, unique graphitized basal plane structure and potential low manufacturing cost, graphene has become a promising candidate as catalyst support in low-temperature fuel cells.4 It has been reported that the application of graphene as the support for Pt 1-D nanomaterials can successfully improve the dispersion and performance of the composite electrocatalyst materials and further improve the catalytic activity and stability by tuning the arrangement of surface active Pt atoms.5 However, this research paid little attention on the effect of alloying of graphene-supported Pt NPs on the ORR performance.5 Therefore, in this study, the graphene-supported Pt and PtPd NPs (G-Pt and G-PtPd) have been prepared and applied as electrocatalysts for ORR. Formic acid method was employed to prepare the 1-D nanomaterials without the use of sacrificial templates and surfactants. To the best of our knowledge, this is the first time that graphene has been used as supports for the Pt alloy NRs in order to enhance ORR performance through electronic modification. The effect of morphological changes, Pd alloying and supports modifications on their ORR performance has been elucidated by the combination of the rotating disc electrode (RDE), high resolution transmission electron microscopy (HRTEM), and X-ray absorption spectroscopy (XAS) characterizations.

Figure 1 shows the morphologies changes of carbon-supported Pt NRs (C-Pt) and G-PtPd during accelerated durability test (ADT) characterized by HRTEM. In Figure 1 (a), NRs of C-Pt with a diameter of 3 nm and a length of 15 nm are grown on the surface of carbon supports while NRs with a larger diameter about 4.4 nm and a longer length about 25 nm can be obtained for G-Pt as displayed in Figure S1 in supporting information (SI). For G-PtPd NRs, their length is 35 nm as displayed in Figure 1 (b), suggesting that graphene can promote the anisotropic growth of Pt. Moreover, after ADT, it can be noted clearly that the rod-like structure almost disappears for C-Pt shown in Figure 1 (c) while NRs with different degrees of aggregations can still be observed for G-Pt, carbon-supported PtPd NRs (C-PtPd), or G-PtPd samples displayed in Figure S1 (c), Figure S1 (d), or Figure 1 (d), respectively, suggesting that either graphene support or Pd alloying may stabilize Pt during ADT, thus enhancing the ORR performance.

Figure S2 in SI shows the X-ray diffraction (XRD) patterns of various NRs. In the view point of bulk analysis, the C-PtPd and G-PtPd samples have fcc alloy structures. However, based on the extended X-ray absorption fine structure (EXAFS) results displayed in Figures S3 and S4, these alloys have a Pt core in the inner core and Pd rich on the outer shell structure. Moreover, Table S1 shows the data analyses of XRD patterns. The peak areas integrated from (111), (200), and (220) planes for various NRs are calculated and used to describe the degrees of anisotropic growth, which influence the ORR performance significantly. It has been reported that the ORR performance increases on the order of Pt(100) << Pt(111) < Pt(110), in which the Pt(111) and Pt(110) planes have similar activity.2b Therefore, the peak area ratio of [(111)+(220)/(200)] for Pt/C (46%, TKK) and various NRs are compared in Table S1. It seems that there is strong disagreement on the NPs and alloying with Pd and using graphene support, the growth of (111) and (200) planes is promoted, suggesting that G-Pt and G-PtPd NRs have potentials to show better ORR performance due to the morphological effect.
Furthermore, Figure S5 displays the X-ray absorption near edge spectroscopy (XANES) spectrum at Pt L\_III edge of Pt/C and various NRs. The absorption at 11564 eV is attributed to the 2p\_3/2-5d electronic transitions and the magnitude of white line is related to the Pt oxidation state. As shown in Figure S5, the differences in the white lines indicate a change in Pt 5d band occupancy due to the adsorption of oxygenated species. The affinity of OH chemisorption on Pt depends on alloying\(^8\) in this study, in order to investigate the effect of morphological changes of catalysts on the d-band vacancies of Pt, XANES results extracted from the Pt L\_III and L\_II white lines are applied to calculate the total number of unoccupied d-states, \((H_{3s})^2\), \(H_{5s}\) can be a good indicator to describe the effect of alloying, morphological changes, and modification on d-band of Pt.\(^7\) It has been reported that the d-band density of states (DOS) is structure-sensitive and the position of the d-band center shifts for different surfaces of Pt\_Ni single crystals,\(^9\) thus affecting their ORR activity. However, for powder nanomaterials, the precise characterization of DOS under ultrahigh vacuum seems to be difficult and the calculation of \(H_{3s}\) becomes an alternative way to get an insight into the d-band. In this study, the \(H_{3s}\) value listed in Table S2 of Pt/C is decreased due to the formation of NRs, and further decreased due to the use of graphene support and alloying. In the viewpoint of d-band, the effect of using graphene as support and alloying with Pd for Pt is similar, whose \(H_{3s}\) is 0.307 and 0.306, respectively. Besides, the Pd alloying and graphene support can modify the d-band synergistically so that the \(H_{3s}\) of G-PtPd is as low as 0.295, suggesting that they have lower unfilled Pt d-states, and more d-band electrons transfer from Pd to Pt, which may lead to the enhanced ORR activity.\(^2b,2c\)

Figure 2 (a) shows the linear sweep voltammetry (LSV) results of Pt/C and various catalysts in O\_2-saturated 0.5 M HClO\_4 solution at 1600 rpm. The onset potentials of various catalysts follow the trend: G-PtPd > C-PtPd > G-Pt > C-Pt > Pt/C, which are inversely proportional to their \(H_{3s}\) values as compared in Table S2. Moreover, their kinetic current densities (\(I_{\text{K}}\)) also have the same sequence, suggesting that the modification of d-band can decisively promote their ORR performance. Besides, the specific activity (SA) and mass activity (MA), which is the \(I_{\text{K}}\) normalized to the electrochemical surface area (ECSA) and Pt loading are summarized in Table S2. The Pt loading is decreased due to the formation of NRs, and further decreased due to the use of graphene support and alloying. In addition, the combination of the both. The SA of as-prepared catalysts after ADT in O\_2 atmosphere is decreased compared with before ADT,\(^4\) the MA retention, which is the ratios of MA/MA after ADT (MA\_ADT) and MA before ADT (MA\_0), are compared in Table S2. It is interesting mentioning that although the graphene support and Pd alloying can both promote ORR of Pt NRs through the modification of Pt, the stability of NRs can be specifically enhanced due to the graphene support. Moreover, Figure S7 summaries the SA changes of Pt/C, C-Pt, and G-PtPd of ADT in O\_2 and their SA after 2000 cycles is about 5, 40, and 81 \(\mu\text{A/cm}^2\), respectively, confirming that the graphene-supported Pd nanomaterials have superior ORR performance to carbon-supported Pt NPs and NRs during long-term test.

Figure 3 summarizes the correlation between \(H_{3s}\) and SA before and after ADT of Pt/C and various NRs. It can be seen that d-band of Pt/C can be modified by formation of NRs, and further decreased through using graphene support, addition of Pd, and the combination of the both. The SA of as-prepared catalysts basically follows the same trend in which lower \(H_{3s}\) suggests more d-band electrons transfer to Pt and higher ORR activity. Moreover, during the ADT, another factor, the resistance to the dissolution, corrosion and migration in acidic media, will influence the SA. The stabilization effect of graphene support is more significant than that from Pd alloying when the SA of G-Pt and C-PtPd is compared. This stabilization can be optimised by synergistic addition of graphene and Pd to Pt NRs. Among all, G-PtPd shows the superior ORR performance to Pt/C and other NRs, implying that the electronic modification effect from Pd and graphene-metal interaction can be promoted significantly for Pt.

The modification and stabilization effect from graphene and Pd also affect the surface chemical states of Pt, as shown in Figure S8 of the X-ray photoelectron spectroscopy (XPS) spectra. The fitting results suggest that the surface Pt/PtO ratios of Pt/C are changed due to the formation of NRs, using graphene support, and alloying with Pd. The formation of surface PtO is inhibited for the G-PtPd to 19 \%, suggesting that the synergistic effect of Pd alloying and graphene support not only influence the electronic state of the bulk but also modify the surface chemical state of the NRs.

**Conclusions**

Graphene has been applied as supports to prepare Pt and Pd/NRs with enhanced ORR performance. The effect of graphene support and Pd alloying on the d-band vacancy and ORR performance has been elucidated. It is observed that G-PtPd has lower unfilled Pt d-states, and more d-band electrons transfer from Pd to Pt, and their SA and MA after ADT is about 6.5 and 2.7 times higher than those of Pt/C, confirming that the electronic modification effect from Pd and graphene-metal interaction can be promoted synergistically for Pt NRs.

**Notes and references**
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Figure 1 HRTEM images for the as-prepared (a) CAPt and (b) GAPtPd catalysts. (c) CAPt and (d) GAPtPd catalysts after ADT.

Figure 2 (a) LSV of Pt/C NP, C-Pt, C-PtPd, G-Pt, and G-PtPd NRs before and (b) after ADT.

Figure 3 Comparison of SA and H charge of the Pt/C NP, C-Pt, C-PtPd, G-Pt, and G-PtPd NR before and after ADT.