N, B and P co-doped graphene as metal-free catalysts for the oxygen reduction in alkaline media

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Doping graphene with heteroatoms such as N, B or P can tailor the properties of the 2D structure of graphene¹ and increase the catalytic activity towards the oxygen reduction reaction (ORR) with respect to pristine graphene.² Here we present a new systematic study in which we compare the performance of B-N and P-N dual-doped graphene and their single-doped N-free counterparts.

The doping-precursor agents are melamine, boric acid and orto-phosphoric acid for N, B and P, respectively. These are mixed with a certain amount of commercial GO and subjected to a thermal treatment at 900 °C during 2 hours with an incremental step of 5 °C min⁻¹ from room temperature. In the Raman analysis (Fig. 1) the ratio between the I_D peak, linked to defective sites in the structure, and the I_G peak, related to the degree of graphitization,³ is lower when B and P are co-doped with N, with respect to B-Gr and P-Gr, respectively. Regarding the catalytic activity (Fig. 2), doping graphene with one single element does not improve the performance with respect to thermally reduced GO without doping. Nevertheless, when graphene is codoped with two elements (BN-Gr and PN-Gr) there is a significant improvement with respect to the catalysts in which these elements are not combined with N. This effect is considerably more noticeable in the case of PN-Gr.

The better performance of the dual codoped graphene catalysts is attributed to a synergistic effect between the two different elements incorporated into the active 2D graphene structure.⁴

Figures







Figure 2: LSV at 1600 rpm in O_2 saturated 0.1 M KOH solution.

References

- X. Wang, G. Sun, P. Routh, D-H. Kim, W. Huang, P. Chen, Chem. Soc. Rev., 43 (2014), 7067.
- [2] C-H. Choi, S-H. Park, S-I. Woo, ACS Nano, 6, 8 (2012), 7084.
- [3] Z-H. Sheng, L. Shao, J-J. Chen, W-J. Bao, F-B. Wang, X-H. Xia, ACS Nano, 5, 6 (2011), 4350.
- [4] Z. Wei, R. Li, X. Gou, ACS Catal., 5 (2015), 4133.