Supporting Information

## One-pot synthesis of mesoporous NiCo<sub>2</sub>O<sub>4</sub> nanoplatelet and graphene hybrid and its oxygen reduction and evolution activities as an efficient bifunctional electrocatalyst

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\*Corresponding Author Email: zhwchen@uwaterloo.ca, Telephone: 1-519-888-4567 Ext. 38664, Fax: 1-519-746-4979 XPS spectra of NiCo<sub>2</sub>O<sub>4</sub> and NiCo<sub>2</sub>O<sub>4</sub>/G



Figure S1. XPS spectra of  $NiCo_2O_4$  (black) and  $NiCo_2O_4/G$  (red) of the elements (a) Ni 2p (b) Co 2p (c) C 1s and (d) O 1s

For the XPS spectra of the elements Ni, Co, and C, no significant shift in the binding energy is observed between NiCo<sub>2</sub>O<sub>4</sub> and NiCo<sub>2</sub>O<sub>4</sub>/G (Figure S1a, b, c), which is consistent with the previous work reported by Dai's group on Co<sub>3</sub>O<sub>4</sub> and graphene hybrid.<sup>1</sup> However, we have observed a shift in the binding energy for the XPS spectrum of the element O in NiCo<sub>2</sub>O<sub>4</sub>/G to a higher value compared to that of NiCo<sub>2</sub>O<sub>4</sub> (Figure S1d). Hence, this is an indication that the hybridization of the metal oxides and graphene sheets is probably occurring through the oxygen species of GO which are likely reacted with the metal precursors at the time of the one-pot synthesis. In a similar work reported on the hybridization of manganese oxide nanostructures with graphene sheets, the authors have put forward the mechanism as the formation of a covalent coordination bond between the metal cation and the oxygen functionalities of GO which act as anchor sites for the metal oxide growth.<sup>2</sup>

## References

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