

*Supporting Information for*

Edge Dislocation Defects Generation in Co<sub>3</sub>O<sub>4</sub> Catalyst: an Efficient  
Tactic to Improve Oxygen Evolution Catalytic Activity

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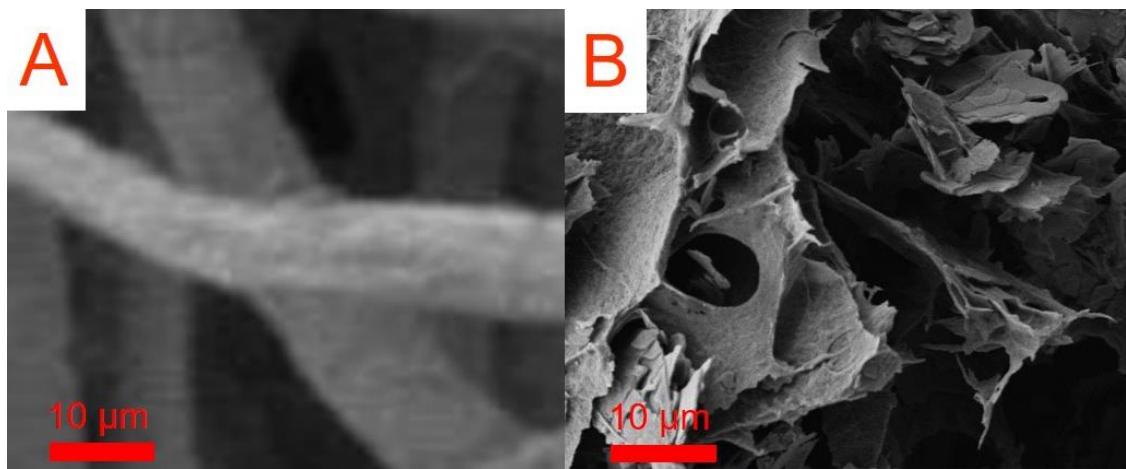
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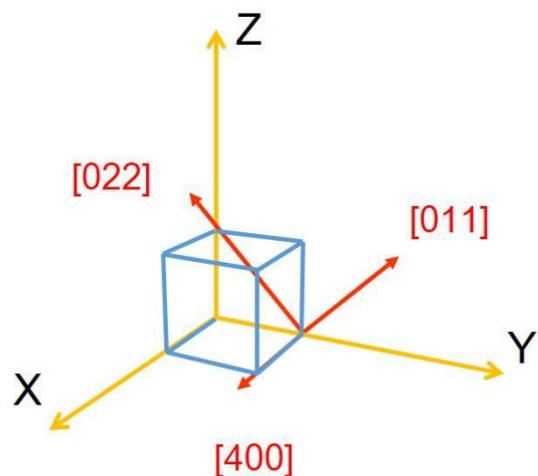
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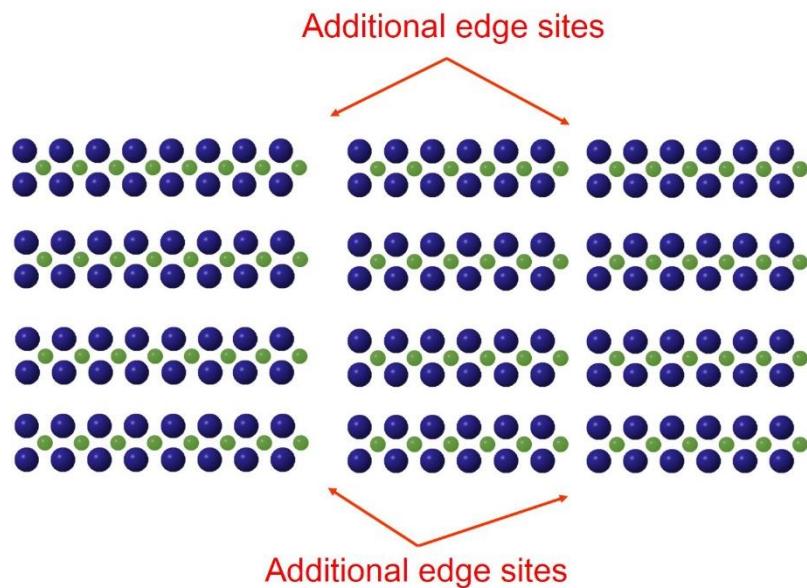
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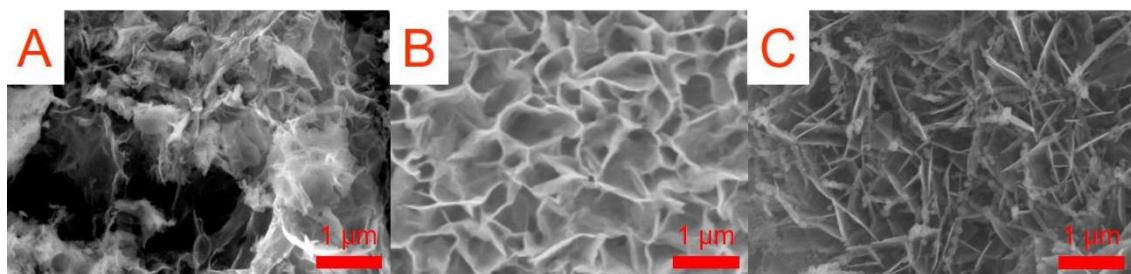
**Fig. S1** SEM images of cellulose I (native cotton linter pulp) (A), and cellulose II (regenerated cellulose) (B).



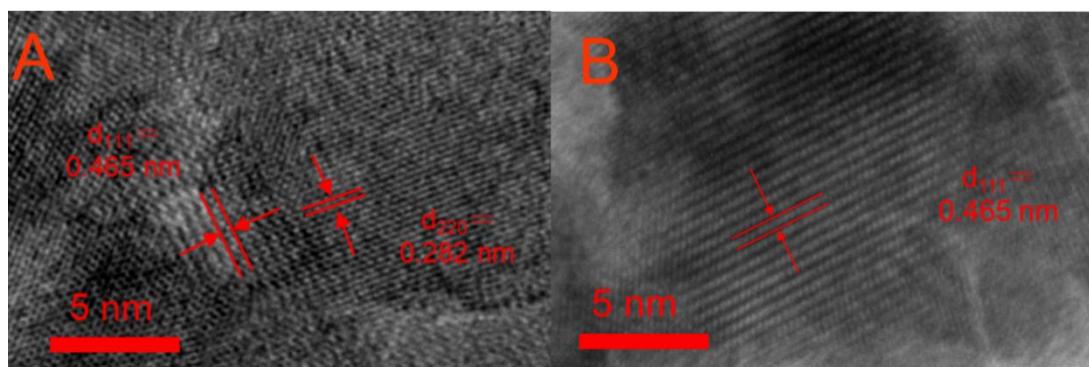
**Fig. S2** Structure model from the obtained fcc of DA-Co<sub>3</sub>O<sub>4</sub>



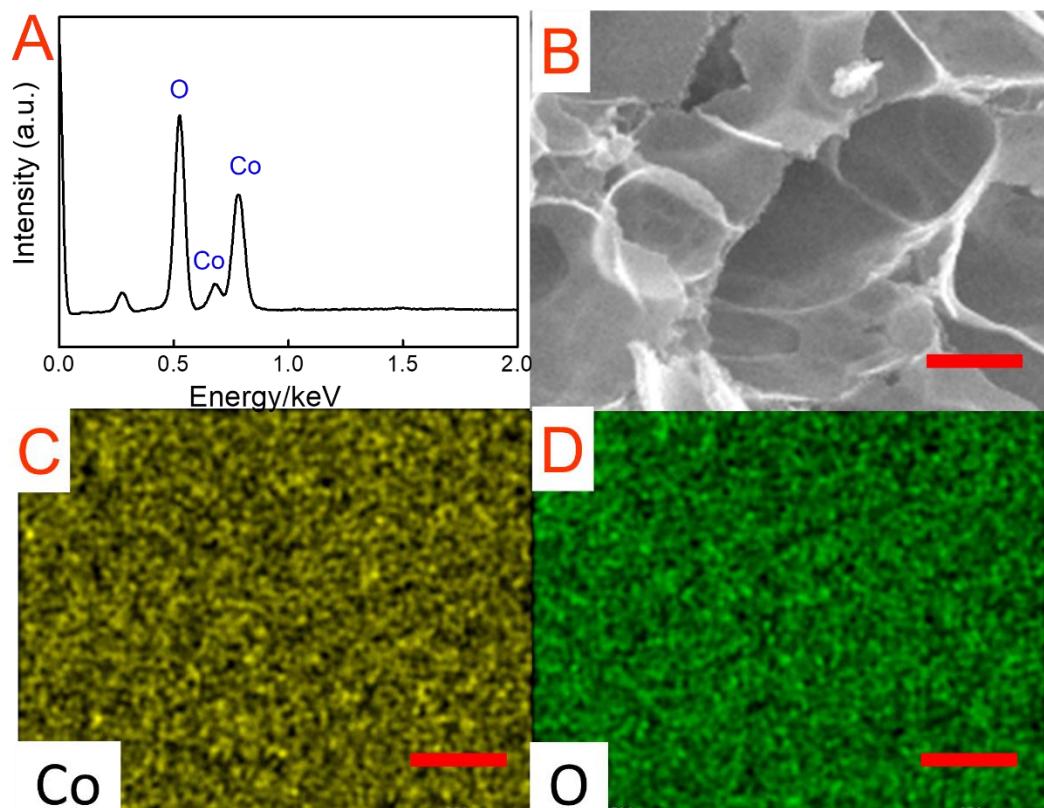
**Fig. S3** Atomic reconstruction of Fig. 3G. Additional active edge sites are marked by red arrow.



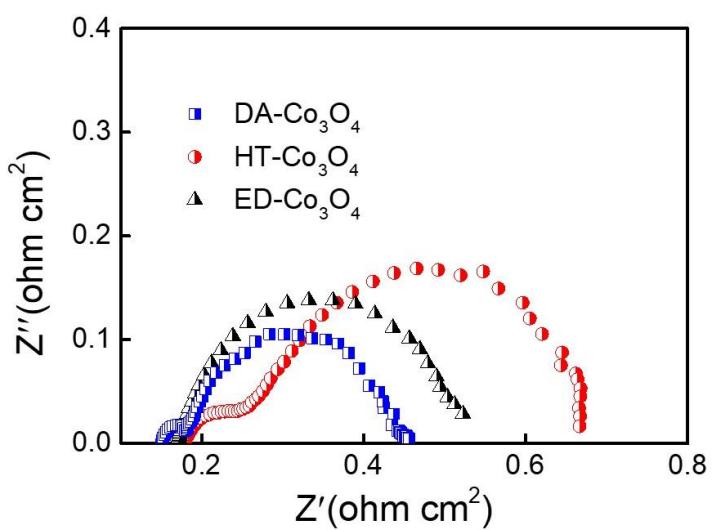
**Fig. S4** SEM images of DA-Co<sub>3</sub>O<sub>4</sub> (A) ED-Co<sub>3</sub>O<sub>4</sub> (B) and HT- Co<sub>3</sub>O<sub>4</sub> (C).



**Fig. S5** TEM images of ED-Co<sub>3</sub>O<sub>4</sub> (A) and HT- Co<sub>3</sub>O<sub>4</sub> (B).



**Fig. S6** EDS spectra, elemental mapping and corresponding SEM image of DA-  $\text{Co}_3\text{O}_4$  nanosheets.



**Fig. S7** Nyquist plots of DA- $\text{Co}_3\text{O}_4$  (A) ED- $\text{Co}_3\text{O}_4$  (B) and HT-  $\text{Co}_3\text{O}_4$  coated electrodes at a 400 mV overpotential.

The resistances of DA-Co<sub>3</sub>O<sub>4</sub>, ED-Co<sub>3</sub>O<sub>4</sub> and HT- Co<sub>3</sub>O<sub>4</sub> coated electrodes for OER process were analyzed by electrochemical impedance spectroscopy technique. As shown in Fig. S7, the resistance in the high frequency region is corresponding to the ohmic resistance R<sub>s</sub>. Here, the charge transfer resistance (signed as R<sub>ct</sub>) was calculated from the arc radius of the Nyquist plot. One can see that the ohmic resistance R<sub>s</sub> of DA-Co<sub>3</sub>O<sub>4</sub> electrocatalysts coated electrodes is slightly smaller than those of ED-Co<sub>3</sub>O<sub>4</sub> and HT- Co<sub>3</sub>O<sub>4</sub> electrodes. In addition, the DA-Co<sub>3</sub>O<sub>4</sub> coated electrode also shows the lowest charge transfer resistance R<sub>ct</sub> among these three Co<sub>3</sub>O<sub>4</sub> catalysts. These results indicate that the interface of DA-Co<sub>3</sub>O<sub>4</sub> electrode had good affinity with the charges and reactants, whose rich defects may promote OH<sup>-</sup> diffusion and adsorption, and enhance charge/mass transfer.

**Table S1.** Comparison of electrocatalytic OER activities of DA-Co<sub>3</sub>O<sub>4</sub> coated electrodes with various state-of-the-art OER catalysts.

Catalysts	Electrolyte	$\eta$ (mV)@ 10 mA cm <sup>-2</sup>	$\eta$ (mV)@ 50 mA cm <sup>-2</sup>	Ref.
DA-Co <sub>3</sub> O <sub>4</sub>	1M KOH	183	261	This work
<b>Noble metal-based</b>				
IrO <sub>2</sub> /C	0.1M KOH	470	N/A	1
IrO <sub>2</sub>	0.1M KOH	450	N/A	2
Ir/C	1M KOH	300	N/A	3
<b>LDH-based</b>				
NiFe LDH /CNT	1M KOH	~235	N/A	3
NiFe LDH	1M KOH	300;	N/A	4
NiCo LDH	1M KOH	330;	N/A	4
CoCo LDH	1M KOH	350	N/A	4
NiFe LDH	1M KOH	224	~300	5
NiFe LDH/RGO	1M KOH	245	~290	6
Ni <sub>2/3</sub> Fe <sub>1/3</sub> -GO	1M KOH	230	N/A	7
NiFe LDH /CQDs	1M KOH	~235	N/A	8
FeNi <sub>8</sub> Co <sub>2</sub> LDH	1M KOH	220	N/A	9
CoMn LDH	1M KOH	324	N/A	10
CoFe LDH	1M KOH	281	341	26
Fe <sub>x</sub> Co <sub>1-x</sub> OOH	1M KOH	266	N/A	27
<b>Transition metal oxides</b>				
Co <sub>3</sub> O <sub>4</sub>	1M KOH	~270	~330	11
CoNiO <sub>x</sub>	1M KOH	336	~360	22
CoFe <sub>2</sub> O <sub>4</sub> /C	1M KOH	240	~290	25
NiCo <sub>2</sub> O <sub>4</sub>	1M KOH	340	N/A	12
Na <sub>1-x</sub> Ni <sub>y</sub> Fe <sub>1-y</sub> O <sub>2</sub>	1M KOH	290		23
Co <sub>3</sub> V <sub>2</sub> O <sub>8</sub>	0.1M KOH	350	N/A	13
CoMoO <sub>4</sub>	1M KOH	312	~390	14
CuO	1 M NaOH	290	~420	20
CuCo <sub>2</sub> O <sub>4</sub> /NrGO	1M KOH	360	~420	15
<b>Other catalysts</b>				
Ni <sub>2</sub> P	1M KOH	310	N/A	16
NiFeP	1 M NaOH	219	~340	21
Co <sub>4</sub> N/CC	1M KOH	257	~270	17
CoP	1M KOH	360	N/A	18
CoS/Ti	1M KOH	361	~400	19
CoS <sub>4.6</sub> O <sub>0.6</sub>	1M KOH	290	N/A	24

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