

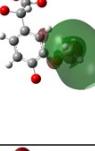
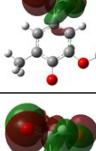
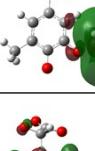
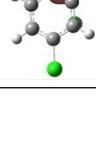
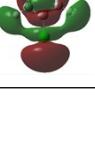
# Supporting Information

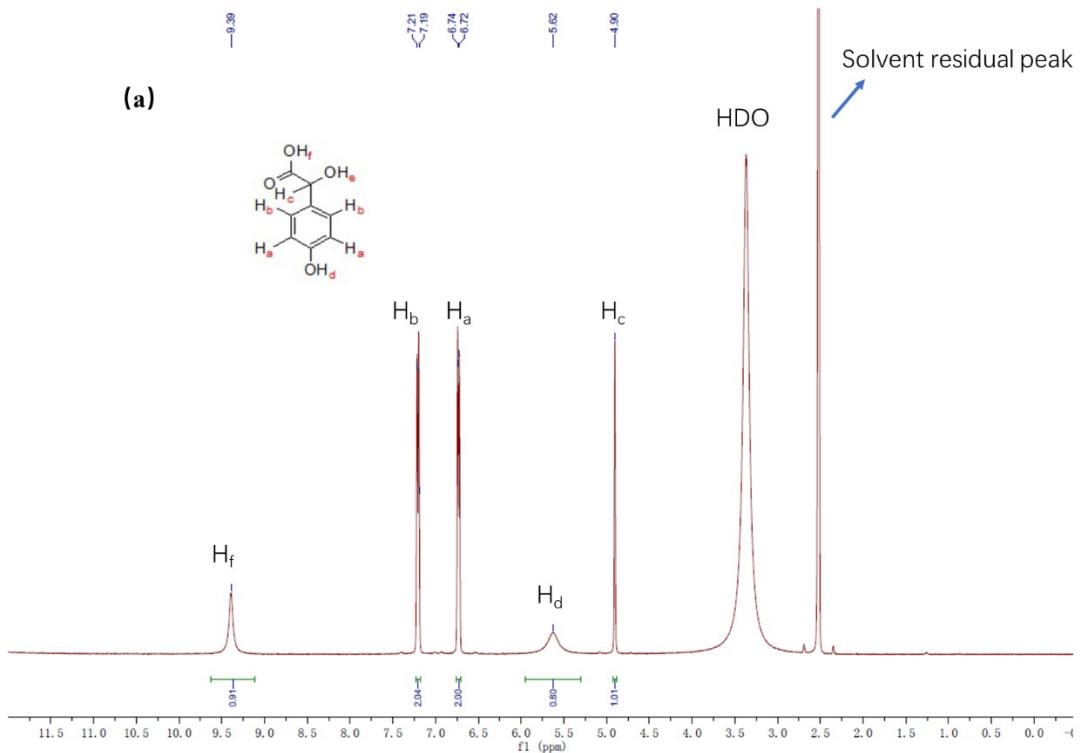
## **Solar-driven Aromatic Aldehydes Green Production from Mandelic Acid Derivatives by Co(II)/C<sub>3</sub>N<sub>4</sub> Combined Catalyst in Aqueous Media**

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Zhenbiao Dong<sup>1</sup>, Ting Tang<sup>\*2</sup>, Wei Zheng<sup>2</sup>, Lehong Jin<sup>2</sup>, Jibo Liu<sup>1\*</sup>

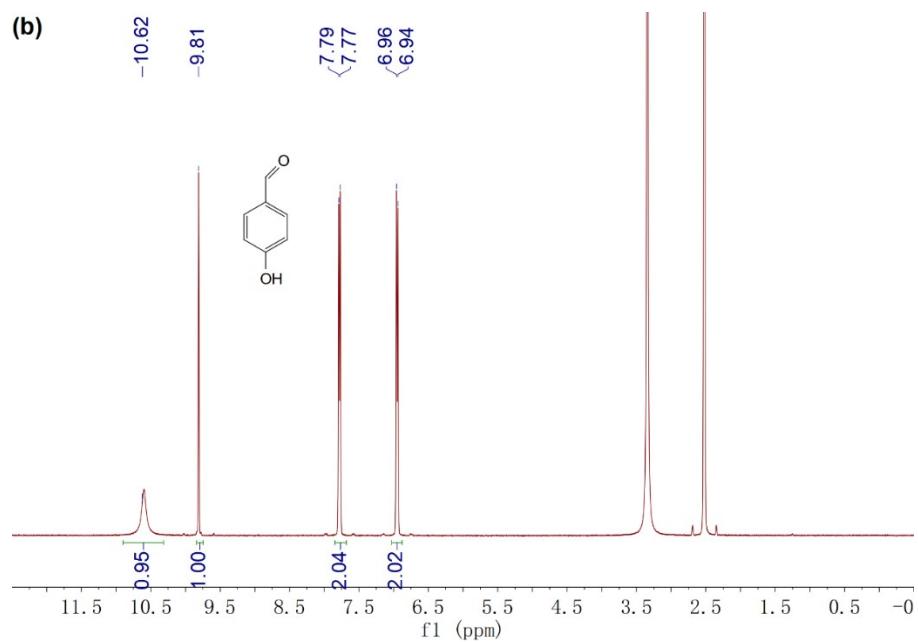
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**Table. S1 The Summering of the DFT Result**

Compound	Charge for C <sub>Bn</sub>	Energy (J/Mol)	HOMO	LUMO	HOMO	LUMO
1	0.132659	290924.7	8.642	11.506		
2	0.129555	373513.0	8.529	9.332		
3	0.127283	445978.0	8.067	8.511		
4	0.127803	443912.1	8.460	9.239		
5	0.113843	286630.9	4.769	6.516		

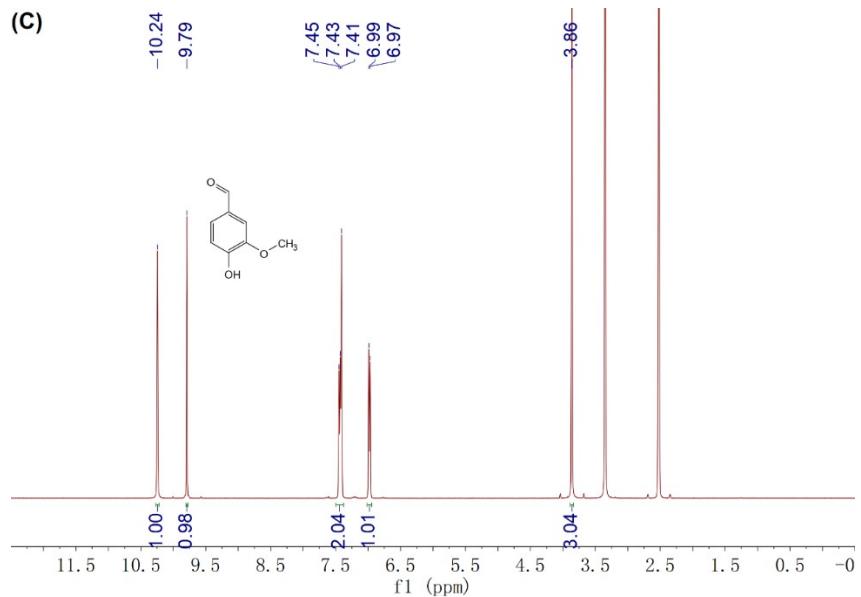


**$^1\text{H-NMR (DMSO-}d_6\text{, TMS, 600MHz)}$**   $\delta$ : 9.39 (s, 1H), 7.19 (d,  $J=12.00$  Hz, 2H), 6.72 (d,  $J=12.00$  Hz, 2H), 5.62 (s, 1H), 4.90 (s, 1H).



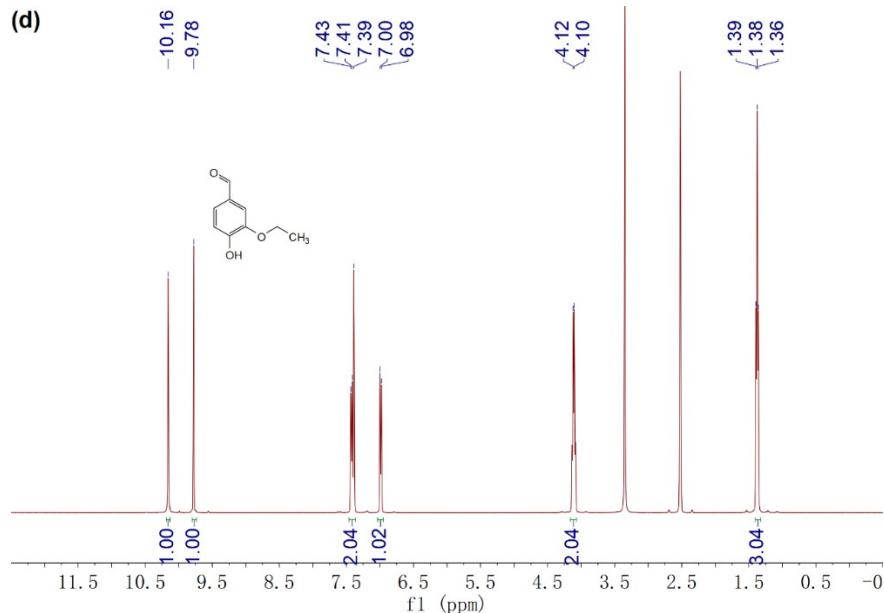
**$^1\text{H-NMR (DMSO-}d_6\text{, TMS, 600MHz)}$**   $\delta$ : 10.60 (s, 1H), 9.81 (s, 1H), 7.77 (d,  $J=12.00$  Hz, 2H), 6.96 (d,  $J=12.00$  Hz, 2H)

(C)

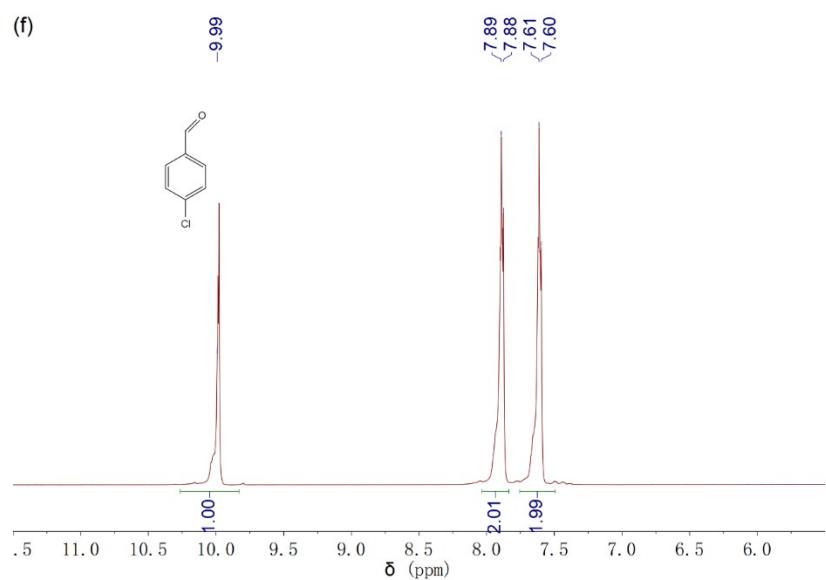
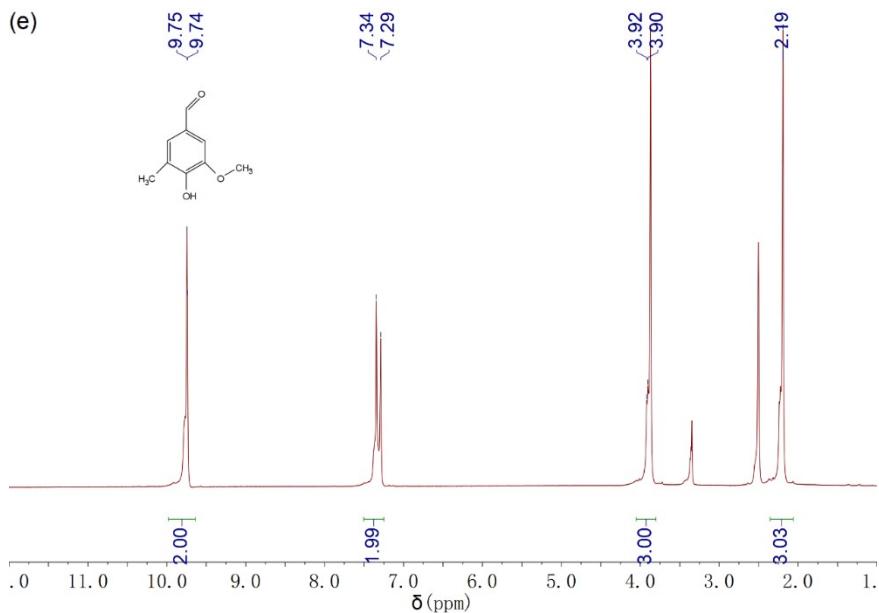


**<sup>1</sup>H-NMR (DMSO-d6, TMS, 600Hz) δ:** 10.28 (s, 1H), 9.78 (s, 1H), 7.43 (t,  $J_1$ =12.00 Hz,  $J_2$ =12.00 Hz, 2H), 6.97 (d, 1H), 3.85 (s, 3H)

(d)



**<sup>1</sup>H-NMR (DMSO-d6, TMS, 600MHz) δ:** 10.16 (s, 1H), 9.78 (s, 1H), 7.37 (t,  $J$ =12.00 Hz, 2H), 6.98 (d,  $J$ = 12.00 Hz, 1H), 4.07 (d,  $J$ = 12.00 Hz, 2H), 1.38 (t,  $J_1$ =12.00 Hz, 3H)



**Fig. S1** The <sup>1</sup>H-NMR spectra of 4-Hydroxyphenylglycolic acid (a), 4-hydroxybenzaldehyde (b), (c) 4-hydroxy-3-methoxymandehyde, (d) 4-hydroxy-3-ethoxymandehyde, (e) 4-hydroxy-3-methoxy-5-methylmandehyde and (f) 4-chlorosylmandehyde.

Fig. S2 The catalytic efficiency of different Co source.

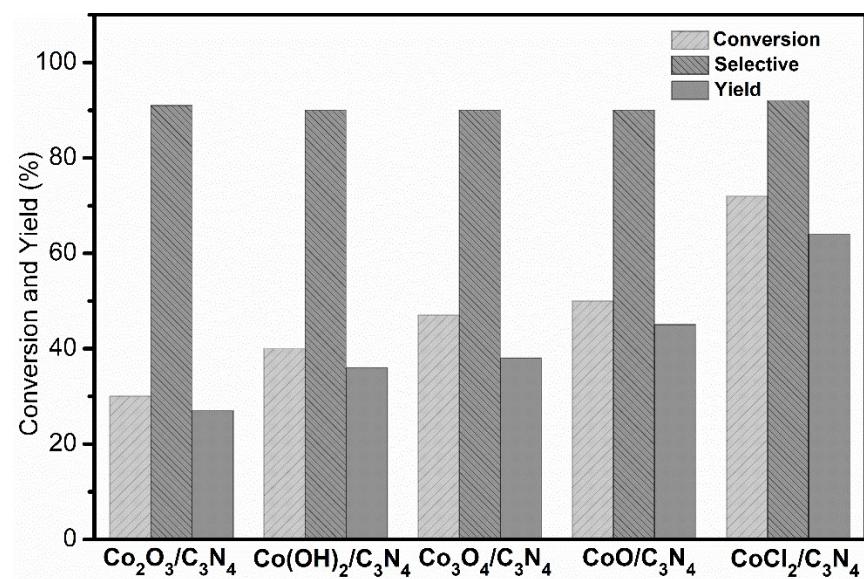
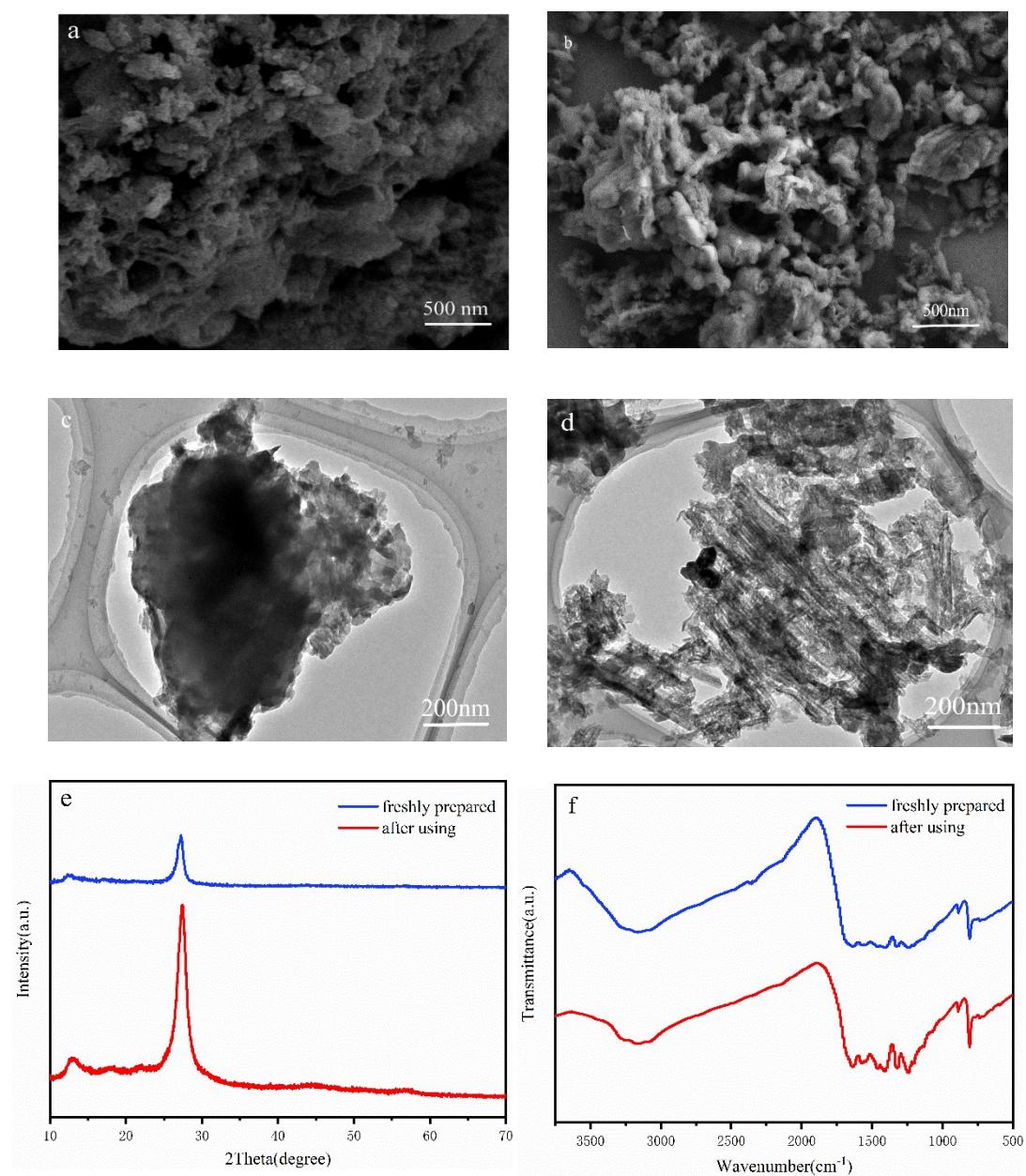
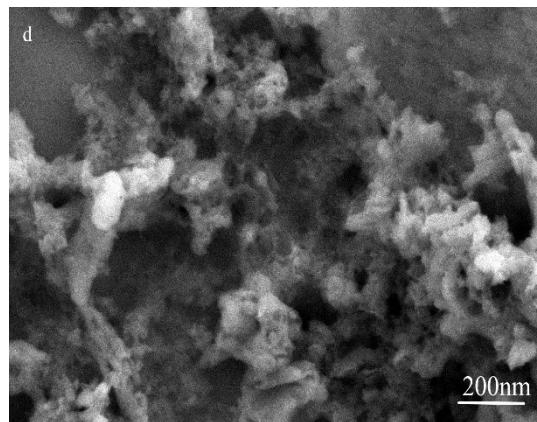
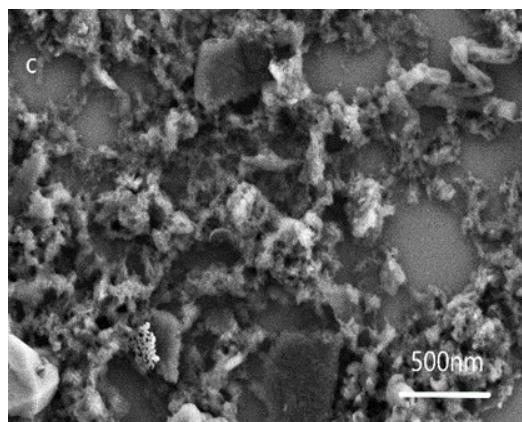
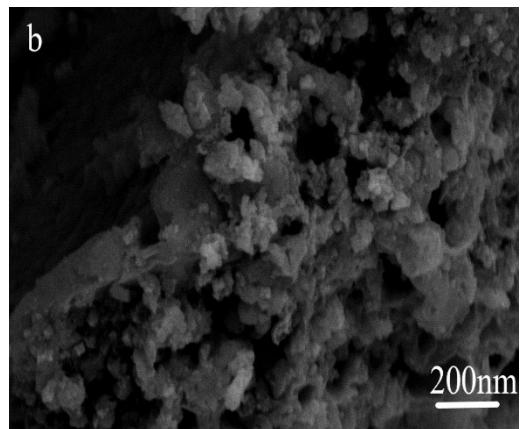
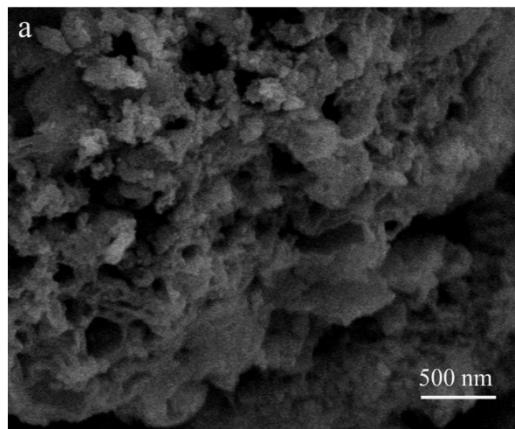


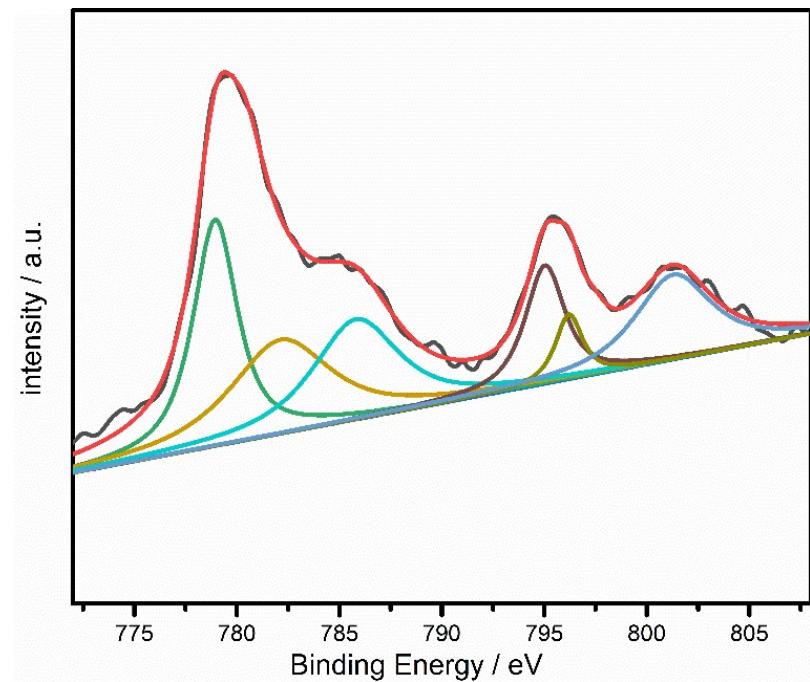
Fig. S3 (a) The field emission scanning electron microscopy (FESEM) image of freshly prepared Co-L/C<sub>3</sub>N<sub>4</sub>-10; (b) The field emission scanning electron microscopy (FESEM) image of Co-L/C<sub>3</sub>N<sub>4</sub>-10 after using; (c) The transmission electron microscopy (TEM) image of freshly prepared Co-L/C<sub>3</sub>N<sub>4</sub>-10; (d) The transmission electron microscopy (TEM) image of Co-L/C<sub>3</sub>N<sub>4</sub>-10 after using; (e) X-ray Diffraction of Co-L/C<sub>3</sub>N<sub>4</sub>-10 before after using; (f) The IR spectra of Co-L/C<sub>3</sub>N<sub>4</sub>-10 before after using.



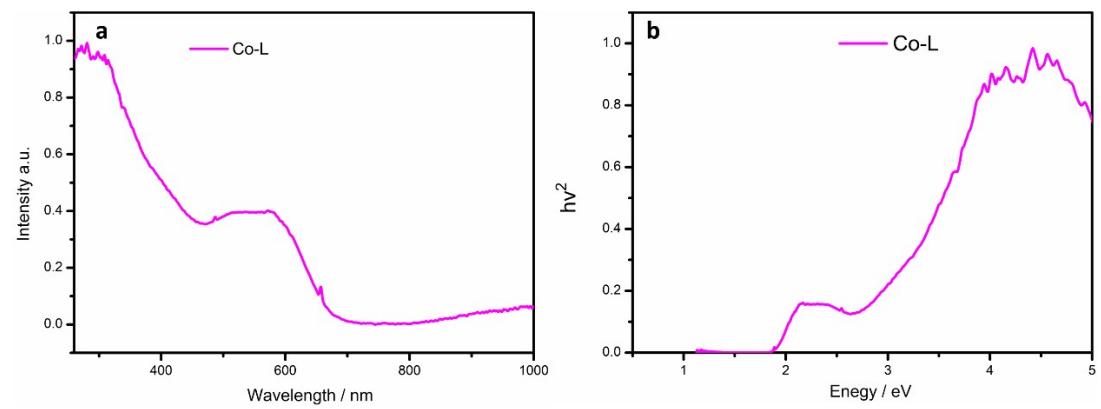
**Fig. S4** (ab) The field emission scanning electron microscopy (FESEM) image of Co-L/C<sub>3</sub>N<sub>4</sub>; (cd) The field emission scanning electron microscopy (FESEM) image of C<sub>3</sub>N<sub>4</sub>.



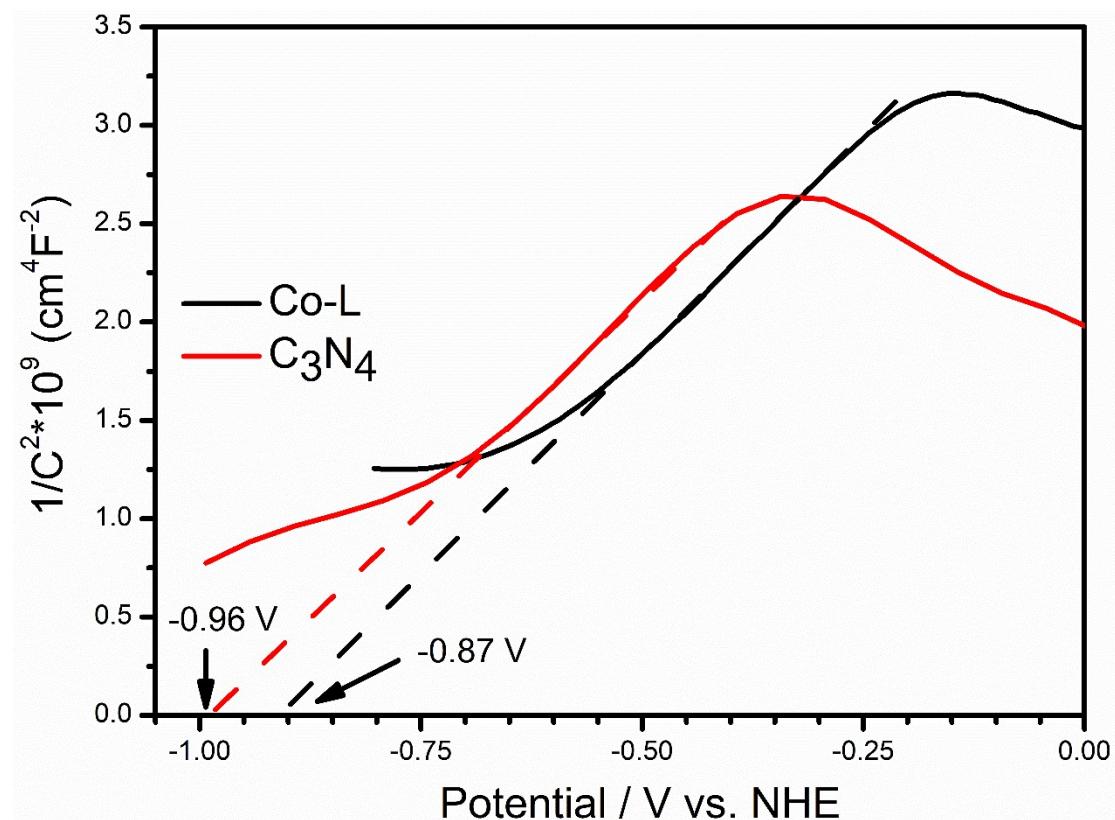
**Fig. S5** X-ray photoelectron spectroscopy (XPS) of Co 2p in Co-L/C<sub>3</sub>N<sub>4</sub>-10 after 3 circles.

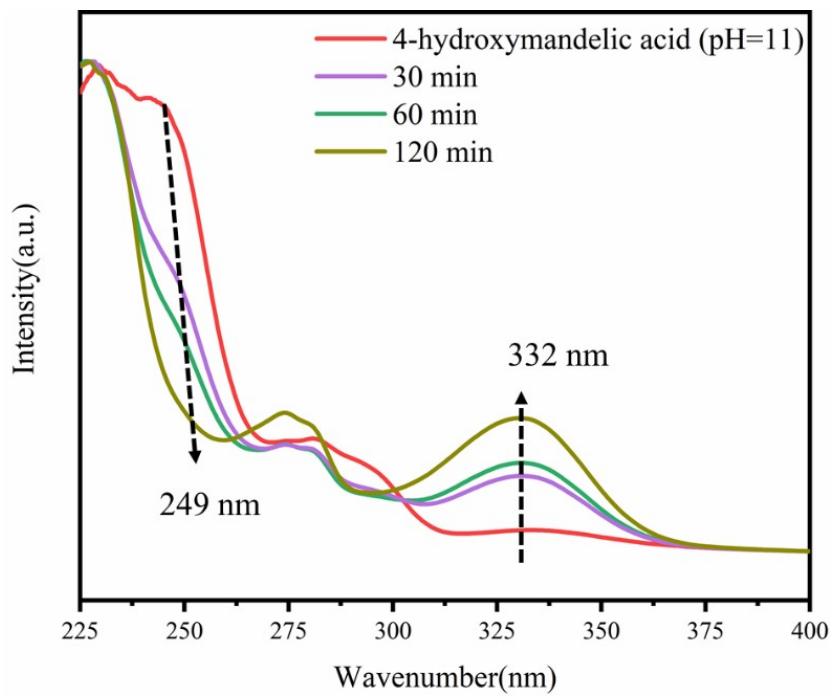


**Fig. S6** (a)The UV-vis diffuse reflectance spectroscopy of Co-L. (b) The estimation of band gap energy of Co-L.



**Fig. S7** The Mott-Schottky plot of  $\text{C}_3\text{N}_4$  and Co-L at a frequency of 962 Hz in the dark, 0.1 M KCl aqueous solution as a supporting electrolyte, 50 mV s<sup>-1</sup> scan rate, Co-L or  $\text{C}_3\text{N}_4$  loaded FTO electrode as a working electrode (area: 0.25 cm<sup>2</sup>), Ag/AgCl (filled with saturated KCl aqueous solution) as a reference electrode, Pt wire as a counter electrode.





**Fig. S8** The UV-vis spectra of 4-hydroxymandelic acid under irradiation.