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# Efficient C–N coupling for urea electrosynthesis on defective $\text{Co}_3\text{O}_4$ with dual-functional sites†

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Urea electrosynthesis under ambient conditions is emerging as a promising alternative to conventional synthetic protocols. However, the weak binding of reactants/intermediates on the catalyst surface induces multiple competing pathways, hindering efficient urea production. Herein, we report the synthesis of defective  $\text{Co}_3\text{O}_4$  catalysts that integrate dual-functional sites for urea production from  $\text{CO}_2$  and nitrite. Regulating the reactant adsorption capacity on defective  $\text{Co}_3\text{O}_4$  catalysts can efficiently control the competing reaction pathways. The urea yield rate of  $3361 \text{ mg h}^{-1} \text{ g}_{\text{cat}}^{-1}$  was achieved with a corresponding faradaic efficiency (FE) of 26.3% and 100% carbon selectivity at a potential of  $-0.7 \text{ V}$  vs. the reversible hydrogen electrode. Both experimental and theoretical investigations reveal that the introduction of oxygen vacancies efficiently triggers the formation of well-matched adsorption/activation sites, optimizing the adsorption of reactants/intermediates while decreasing the C–N coupling reaction energy. This work offers new insights into the development of dual-functional catalysts based on non-noble transition metal oxides with oxygen vacancies, enabling the efficient electrosynthesis of essential C–N fine chemicals.

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## Introduction

Urea is widely used as a fertilizer, as well as a chemical raw material for medicines and pesticides. However, traditional industrial urea synthesis, relying on the reaction of  $\text{CO}_2$  and  $\text{NH}_3$ , involves harsh conditions and leaves a significant carbon footprint,<sup>1,2</sup> necessitating the pursuit of greener, more energy-efficient, and cost-effective alternatives.<sup>3,4</sup> Electrochemical urea synthesis from N-integrated electrocatalytic  $\text{CO}_2$  reduction is gradually emerging as a promising alternative to reform the conventional urea industry.<sup>5–7</sup> Most strikingly, emerging electrocatalytic co-reduction of  $\text{N}_2$  and  $\text{CO}_2$  in an aqueous solution has been demonstrated as a feasible method for urea synthesis.<sup>8–12</sup> However, the inherent chemical inertness of  $\text{N}_2$ , such as high triple bond dissociation energy ( $941 \text{ kJ mol}^{-1}$ ) and low solubility ( $\sim 0.02 \text{ v/v}$ , 1 atm,  $25^\circ\text{C}$ ), poses a considerable limitation on urea yield rates.

Alternatively, N-integrated electrochemical  $\text{CO}_2$  reduction can be performed with nitrate/nitrite, which is a widely existing contaminant in the wastewater and poses severe risks to both public health and the environment. In particular, using nitrite as a N-containing reactant for urea electrosynthesis is highly desirable due to its merits of superior solubility (e.g.  $\text{KNO}_2$ ,  $2.81 \text{ g mL}^{-1}$ ,  $0^\circ\text{C}$ ) and lower N=O bond dissociation energy ( $204 \text{ kJ mol}^{-1}$ ), which could potentially result in higher performance for urea synthesis.<sup>13</sup> Some pioneer works have shown that urea can be synthesized from electrochemically coupling  $\text{CO}_2$  and nitrate/nitrite.<sup>14–20</sup> For example, the  $\text{In}(\text{OH})_3$ –S catalyst could promote co-electrolysis of  $\text{CO}_2$  and nitrate to urea with an average yield rate of  $533.1 \text{ mg h}^{-1} \text{ g}_{\text{cat}}^{-1}$  and a Faradaic efficiency (FE) of 53.4% at the potential of  $-0.6 \text{ V}$  vs. the reversible hydrogen electrode (RHE).<sup>21</sup> Over a  $\text{Cu}_1\text{–CeO}_2$  catalyst, the yield rate of urea could reach  $52.84 \text{ mmol h}^{-1} \text{ g}_{\text{cat}}^{-1}$  at  $-1.6 \text{ V}$  vs. RHE.<sup>22</sup> Low-valence  $\text{Cu}^{\delta+}$ -doped anatase  $\text{TiO}_2$  was employed as an electrocatalyst for the co-electrolysis of  $\text{CO}_2$  and nitrite ions to urea with a yield rate of  $20.8 \text{ }\mu\text{mol h}^{-1}$  at a low potential of  $-0.4 \text{ V}$  vs. RHE.<sup>23</sup> In another study, a Te-doped Pd nanocrystal was utilized to enhance the electrochemical urea production by coupling  $\text{CO}_2$  and nitrite reduction with nearly 12.2% FE and 88.7% N atom efficiency.<sup>24</sup> Despite substantial progress having been made in urea synthesis, there still remains a significant challenge in improving urea FE and yield rate. This is because the parallel  $\text{CO}_2$  reduction reaction, nitrate/nitrite reduction reaction and inescapable hydrogen evolution reaction (HER) strongly compete with the desirable C–

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N coupling reaction, leading to low selectivity of the urea product. The primary obstacle lies in the suppression of selective hydrogenation of intermediate species to unwanted by-products and the hindrance of the unfavorable HER to facilitate C–N coupling for urea production.

To address this, we proposed a two-step thermal annealing strategy to synthesize defective  $\text{Co}_3\text{O}_4$  catalysts, integrating dual-functional sites to favor urea electrosynthesis. We discovered that the presence of oxygen vacancies ( $\text{V}_\text{o}$ ) could construct well-matched adsorption/activation sites, which strengthened the adsorption of  $\text{CO}_2$  and  $\text{NO}_2^-$  reactants. The modulated local environment on Co atoms near the oxygen vacancies, serving as the dual-functional sites, not only strengthened the adsorption of intermediates but also decreased the C–N coupling resistance in urea synthesis. Notably, when using defective  $\text{Co}_3\text{O}_4$  obtained under suitable treatment conditions as an electrocatalyst, an impressive urea yield rate of  $3361 \text{ mg h}^{-1} \text{ g}_{\text{cat}}^{-1}$  was achieved with a corresponding FE of 26.3% and 100% carbon selectivity at the potential of  $-0.7 \text{ V vs. RHE}$ .

## Results and discussion

We used a two-step thermal annealing strategy to synthesize defective  $\text{Co}_3\text{O}_4$  catalysts with tunable  $\text{V}_\text{o}$  concentrations. Briefly,  $\text{Co}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$  and  $\text{NaCl}$  were first mixed in an agate mortar, followed by adding  $\text{NaOH}$  solution for sufficient grinding and solvent removal. The resulting solid solution was then directly pyrolysed under an Ar atmosphere to get the pristine  $\text{Co}_3\text{O}_4$ , in which the  $\text{V}_\text{o}$  concentrations could be tuned by thermal treatment under a reducing atmosphere ( $\text{H}_2/\text{Ar}$ ) for different times (0.5, 1.0 and 1.5 h). The as-synthesized catalysts were denoted as  $\text{Co}_3\text{O}_4\text{-0.5}$ ,  $\text{Co}_3\text{O}_4\text{-1.0}$ , and  $\text{Co}_3\text{O}_4\text{-1.5}$ , respectively. We commenced our characterization using scanning electron microscopy (SEM). As shown in Fig. 1a and b and S1,† both the pristine and reduced thermally treated  $\text{Co}_3\text{O}_4$  had nanosized and octahedral structures, indicating that the reducing thermal annealing process did not affect the initial morphology. The morphological and structural information was further studied by transmission electron microscopy (TEM) and high-resolution transmission electron microscopy (HRTEM). Typically, the TEM images in Fig. 1c and d demonstrated that  $\text{Co}_3\text{O}_4\text{-1.0}$  had a rougher surface than pristine  $\text{Co}_3\text{O}_4$ , which could be attributed to the formation of high  $\text{V}_\text{o}$  concentration on the catalyst surface. The HRTEM images in Fig. 1e and f showed that the lattice fringe of the  $\text{Co}_3\text{O}_4$  nanoparticle was 0.24 nm, which is ascribed to the lattice plane distance of the (311) plane of  $\text{Co}_3\text{O}_4$ . The main difference was that the pristine  $\text{Co}_3\text{O}_4$  featured a flatter crystal surface (Fig. 1e) than  $\text{Co}_3\text{O}_4\text{-1.0}$  (Fig. 1f). The normalized Co K-edge X-ray absorption near edge structure (XANES) spectra (Fig. 1g) revealed a slightly reduced average valence state of the Co species in  $\text{Co}_3\text{O}_4\text{-1.0}$ , indicating that  $\text{V}_\text{o}$  can manipulate the electronic state of the nearby Co atoms. Additionally, the Co K-edge extended X-ray absorption fine structure (EXAFS) spectra of pristine  $\text{Co}_3\text{O}_4$  and  $\text{Co}_3\text{O}_4\text{-1.0}$  demonstrated three distinct peaks (Fig. 1h), which were identified as Co–O, tetrahedral Co–Co, and octahedral Co–Co bonds, respectively. Comparably, the

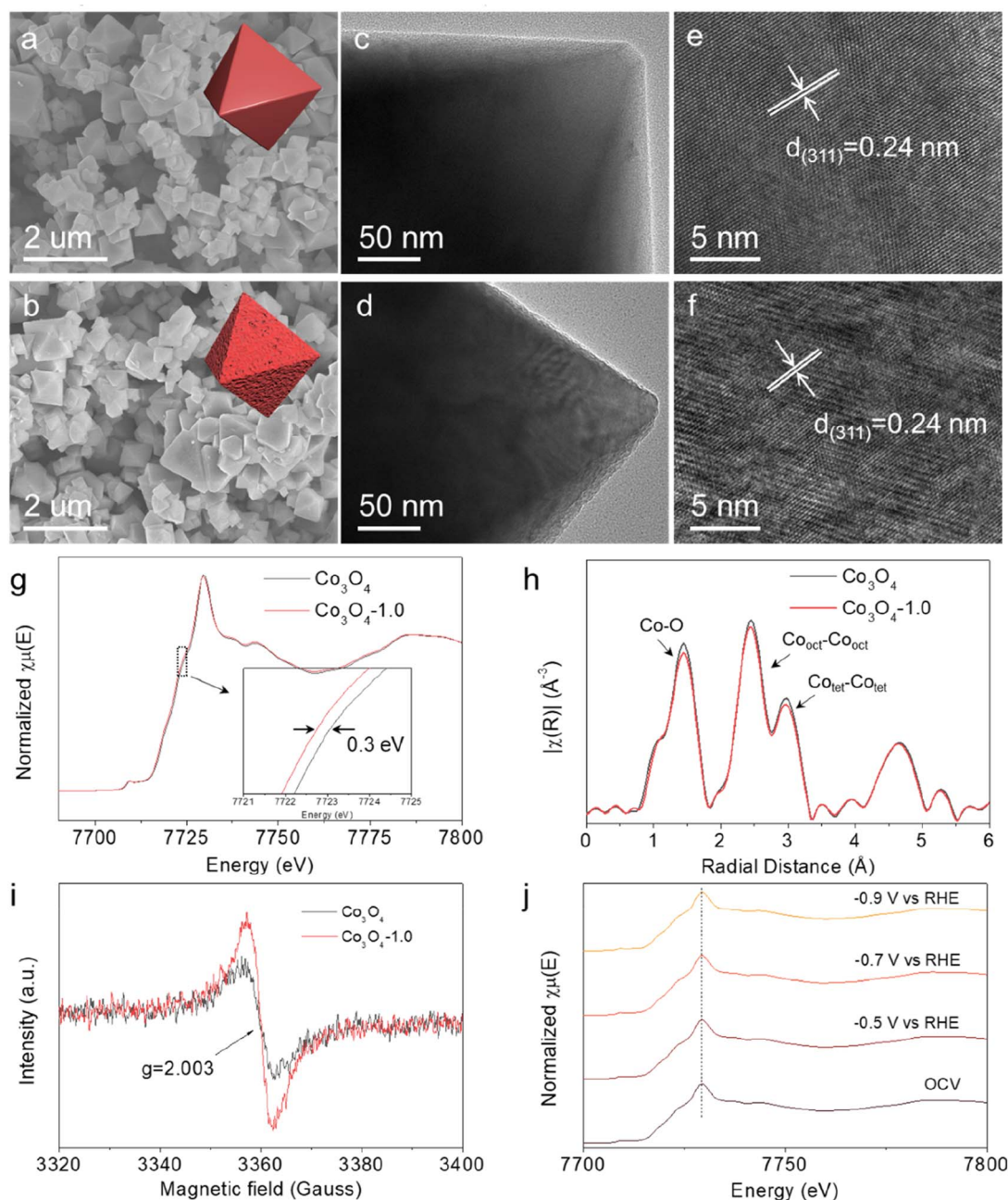
intensity of the Co–O bond in  $\text{Co}_3\text{O}_4\text{-1.0}$  was lower than that in the initial  $\text{Co}_3\text{O}_4$ , suggesting a reduced oxygen coordination number for the Co atoms in  $\text{Co}_3\text{O}_4\text{-1.0}$ . This reduction is attributed to the introduction of additional  $\text{V}_\text{o}$  in  $\text{Co}_3\text{O}_4\text{-1.0}$ .<sup>25,26</sup>

The X-ray diffraction (XRD) patterns of the catalysts are shown in Fig. S2.† In detail, the peaks located at 31, 37, 44, 59 and  $65^\circ$  can be indexed to the (220), (311), (400), (511) and (440) crystalline planes of  $\text{Co}_3\text{O}_4$  (JCPDS: PDF#43-1003), respectively, confirming the successful synthesis of  $\text{Co}_3\text{O}_4$ . The electron paramagnetic resonance (EPR) spectra were recorded to identify  $\text{V}_\text{o}$ . The electron deficient oxygen species exhibited an EPR signal at  $g = 2.003$  (Fig. 1i and S3†).<sup>27,28</sup> The intensity increased with increasing annealing time, revealing a higher  $\text{V}_\text{o}$  concentration in the  $\text{Co}_3\text{O}_4\text{-1.0}$  catalyst than in the pristine  $\text{Co}_3\text{O}_4$  catalyst. Furthermore, X-ray photoelectron spectroscopy (XPS) measurement analysis was performed to investigate the chemical states and compositions of the catalysts. The XPS spectra of O 1s (Fig. S4†) verified that the lattice oxygen ( $\text{O}_1$ ) and oxygen vacancies ( $\text{O}_2$ ) existed in all catalysts.<sup>29,30</sup> The  $\text{O}_2/\text{O}_1$  ratio means the ratio of  $\text{O}_2$  and  $\text{O}_1$  area in the O 1s spectra, and followed the sequence of pristine  $\text{Co}_3\text{O}_4 < \text{Co}_3\text{O}_4\text{-0.5} < \text{Co}_3\text{O}_4\text{-1.0} < \text{Co}_3\text{O}_4\text{-1.5}$ , suggesting the presence of more  $\text{V}_\text{o}$  species in the  $\text{Co}_3\text{O}_4\text{-1.5}$  catalyst, which was consistent with the EPR results. Fig. S5† displayed the impact of  $\text{V}_\text{o}$  on the electronic structure of  $\text{Co}_3\text{O}_4$ . The peaks at around 780 eV (Co  $2\text{p}_{3/2}$ ) and 795 eV (Co  $2\text{p}_{1/2}$ ) retained the characteristic feature of Co species.<sup>25</sup> Apparently, the  $\text{Co}^{2+}/\text{Co}^{3+}$  ratio increased from 0.85 to 1.54 with increasing  $\text{V}_\text{o}$  concentration, suggesting a decrease in Co valence state on the  $\text{Co}_3\text{O}_4$  surface after thermal treatment under the reducing atmosphere. Interestingly, we found a positive linear correlation between the ratio of  $\text{Co}^{2+}/\text{Co}^{3+}$  and the ratio of  $\text{O}_2/\text{O}_1$  (Fig. S6†), indicating that the  $\text{V}_\text{o}$  level in the catalysts was directly correlated with the electron cloud density of Co atoms, which is critical for the electrocatalytic activity.

The structural stability of electrocatalyst is a crucial indicator when assessing its performance. Therefore, we first carried out operando X-ray absorption spectroscopy (XAS) to investigate the structural and valence state changes of Co element under the electrochemical reaction conditions. In the operando XAS measurement, the potential was decreased from open-circuit voltage (OCV) to  $-0.5 \text{ V}$ ,  $-0.7 \text{ V}$ , and  $-0.9 \text{ V vs. RHE}$ . Fig. 1j and S7† demonstrate that there was no energy shift on the Co K-edge in the XANES spectra as the applied potential decreased. Similarly, no changes were observed in the coordination environment of Co atoms in the EXAFS spectra (Fig. S8†). These results collectively suggest that the local structure and electronic state of Co species in  $\text{Co}_3\text{O}_4\text{-1.0}$  remained stable during the electrochemical reaction process.

The electrocatalytic activity of the as-prepared catalysts for application in urea synthesis was evaluated in an H-type cell. Cyclic voltammetry (CV) was carried out to preliminarily assess the performance of the  $\text{Co}_3\text{O}_4\text{-1.0}$  in Ar or  $\text{CO}_2$  saturated  $0.2 \text{ M KHCO}_3 + 0.02 \text{ M KNO}_2$  electrolyte. As shown in Fig. 2a, in the presence of  $\text{CO}_2$ ,  $\text{Co}_3\text{O}_4\text{-1.0}$  exhibited larger current density responses during the forward scan from  $-0.2 \text{ V}$  to  $-0.8 \text{ V vs. RHE}$ , resulting from simultaneous electroreduction of  $\text{CO}_2$  and  $\text{NO}_2^-$ . Subsequently, chronoamperometry tests were conducted





**Fig. 1** SEM images of (a) pristine  $\text{Co}_3\text{O}_4$  and (b)  $\text{Co}_3\text{O}_4$ -1.0. TEM images of (c) pristine  $\text{Co}_3\text{O}_4$  and (d)  $\text{Co}_3\text{O}_4$ -1.0. HRTEM images of (e) pristine  $\text{Co}_3\text{O}_4$  and (f)  $\text{Co}_3\text{O}_4$ -1.0. (g) XANES and (h) Fourier-transformed EXAFS spectra of the Co K-edge over pristine  $\text{Co}_3\text{O}_4$  and  $\text{Co}_3\text{O}_4$ -1.0. (i) EPR spectra of pristine  $\text{Co}_3\text{O}_4$  and  $\text{Co}_3\text{O}_4$ -1.0. (j) Operando XANES under the electrochemical conditions of Co K-edge over  $\text{Co}_3\text{O}_4$ -1.0. OCV stands for open-circuit voltage.

at applied potentials ranging from  $-0.5$  to  $-0.9$  V vs. RHE to analyze the FE and the yield of products. The yield of urea was quantified using the well-established urease decomposition and indophenol blue spectrophotometric method (Fig. S9 and S10<sup>†</sup>).<sup>15,25,31</sup> The performances of urea electrosynthesis at different applied potentials are displayed in Fig. 2b and S11.<sup>†</sup> It can be found that  $\text{Co}_3\text{O}_4$ -1.0 mainly yielded  $\text{H}_2$ ,  $\text{NH}_3$ , and urea with a combined FE of around 100%. With increasing applied potential, the current density and the FE of urea are increased. At the potential of  $-0.7$  V vs. RHE, the FE of urea could reach

26.3% with the current density of  $34.1 \text{ mA cm}^{-2}$  over the  $\text{Co}_3\text{O}_4$ -1.0 catalyst. When the potential was beyond  $-0.7$  V vs. RHE, the FE of urea decreased, mainly due to the highly competitive HER. It is noteworthy that no other reduction products from  $\text{CO}_2$  were detected across the potential range, indicating that the catalyst demonstrated a 100% carbon selectivity for C–N coupling. Fig. S12–S14<sup>†</sup> show the FE of urea and current density of comparative catalysts (pristine  $\text{Co}_3\text{O}_4$ ,  $\text{Co}_3\text{O}_4$ -0.5, and  $\text{Co}_3\text{O}_4$ -1.5) within the potential range from  $-0.5$  V to  $-0.9$  V vs. RHE. This indicates that the product distributions of  $\text{NH}_3$ , urea, and



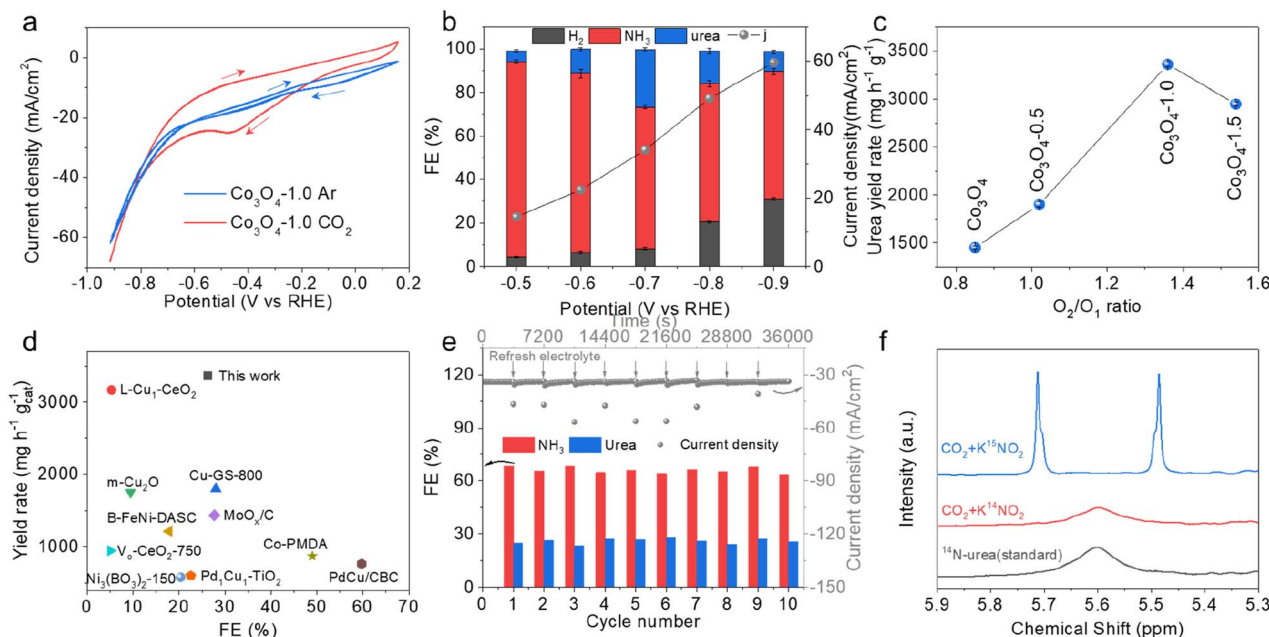


Fig. 2 (a) CV curves of  $\text{Co}_3\text{O}_4$ -1.0 at the scan rate of  $50 \text{ mV s}^{-1}$  in Ar or  $\text{CO}_2$  saturated electrolyte. (b) The FE of major products and total current densities over the  $\text{Co}_3\text{O}_4$ -1.0 electrocatalyst in  $\text{CO}_2$  saturated  $0.2 \text{ M KHCO}_3 + 0.02 \text{ M KNO}_2$  electrolyte. (c) The urea yield rate vs. the ratio of  $\text{O}_2/\text{O}_1$  at the potential of  $-0.7 \text{ V}$  (vs. RHE). (d) Comparison of the FE (%) of urea and yield rate ( $\text{mg h}^{-1} \text{ g}_{\text{cat}}^{-1}$ ) of urea of  $\text{Co}_3\text{O}_4$ -1.0 with those of state-of-the-art catalysts for application in urea synthesis. (e) Stability tests of  $\text{Co}_3\text{O}_4$ -1.0 at the potential of  $-0.7 \text{ V}$  (vs. RHE). (f)  $^1\text{H}$  NMR spectra obtained by using  $\text{K}^{14}\text{NO}_2$  and  $\text{K}^{15}\text{NO}_2$  as the reactants with  $\text{CO}_2$  and a standard  $^{14}\text{N}$ -urea sample.

$\text{H}_2$  were correlated with the  $\text{V}_\text{o}$  concentration in the catalysts, in which  $\text{Co}_3\text{O}_4$ -1.0 with an appropriate  $\text{V}_\text{o}$  concentration exhibited the best performance for urea formation among all the catalysts. At the optimized condition, the urea yield rate over the  $\text{Co}_3\text{O}_4$ -1.0 catalyst could reach  $3361 \text{ mg h}^{-1} \text{ g}_{\text{cat}}^{-1}$  (Fig. 2c), which was nearly 2.3 times that of pristine  $\text{Co}_3\text{O}_4$ . For the pristine  $\text{Co}_3\text{O}_4$  catalyst, the FE of urea was only 13.7% with a limited urea yield rate of  $1450 \text{ mg h}^{-1} \text{ g}_{\text{cat}}^{-1}$ . Comparison with the state-of-the-art catalysts for urea synthesis is presented in Fig. 2d and Table S1†. It shows that  $\text{Co}_3\text{O}_4$ -1.0 was highly efficient as an electrocatalyst for C–N coupling to produce urea and it could reach high FE and yield rate at a low reduction potential.

Remarkably, in the ten consecutive cycles at  $-0.7 \text{ V}$  vs. RHE, the  $\text{Co}_3\text{O}_4$ -1.0 achieved long-term stability with no obvious decays in current density, and the FE of urea and  $\text{NH}_3$  could be held around 25% and 65%, respectively, during the whole electrolysis process (Fig. 2e). Moreover, the XRD, XPS, and TEM measurements showed that the crystal structure, valence states,  $\text{V}_\text{o}$ , and morphology were well preserved after the long-term electrolysis (Fig. S15–S17†), indicating the excellent electrochemical and structural stability of the  $\text{Co}_3\text{O}_4$ -1.0 catalyst.

To identify the source of urea production, we performed the isotope-labelling experiments in  $\text{CO}_2$ -saturated  $0.2 \text{ M KHCO}_3 + 0.02 \text{ M K}^{15}\text{NO}_2$  electrolytes. Fig. 2f illustrates the  $^1\text{H}$  NMR spectrum of the electrolyte with  $\text{K}^{15}\text{NO}_2$ , displaying the typical double peaks of  $^{15}\text{N}$ -urea at 5.48 and 5.71 ppm, while the  $^1\text{H}$  NMR spectrum of the electrolyte with  $\text{K}^{14}\text{NO}_2$  only exhibits a single peak of  $^{14}\text{N}$ -urea at 5.60 ppm. The  $^1\text{H}$  NMR results confirm that there is urea production in the coupling

electrolysis of  $\text{CO}_2$  and  $^{14}\text{NO}_2^-$  on  $\text{Co}_3\text{O}_4$ -1.0. Furthermore, we employed the  $^1\text{H}$  NMR technique (Fig. S18 and S19†) to quantitatively measure the generated urea and concurrently validate the accuracy of the UV-vis method. The obtained results (Fig. S20†) demonstrate a consistent urea concentration measurement between the  $^1\text{H}$  NMR and UV-vis methods, confirming the accuracy of the employed quantitative method.

Considering the experimental observations above, we think that the high electrocatalytic activity of the  $\text{Co}_3\text{O}_4$ -1.0 catalyst could be attributed to the generation of abundant active sites in defective  $\text{Co}_3\text{O}_4$ , which was kinetically favorable for the reaction. To verify the hypothesis, we estimated the electrochemical active surface area (ECSA) of various catalysts through electrochemical double-layer capacitance ( $C_{\text{dl}}$ ) measurements. Fig. S21† shows the cyclic voltammetry (CV) curves with various scan rates at the non-faradaic region for the catalysts. The linear slopes in Fig. S22† show that the  $\text{Co}_3\text{O}_4$ -1.0 had a larger ECSA, suggesting that the higher  $\text{V}_\text{o}$  concentration was responsible for the generation of more active sites. After normalizing the urea yield rate to ECSA (Fig. S23†),  $\text{Co}_3\text{O}_4$ -1.0 still exhibited the largest urea yield rate ( $1832 \text{ mg h}^{-1} \text{ g}_{\text{cat}}^{-1}$ ) at the potential of  $-0.7 \text{ V}$  vs. RHE, which indicates that the  $\text{V}_\text{o}$  concentration regulation could also improve the intrinsic activity of urea production over the catalysts. In addition, the electrochemical impedance spectrum (EIS, Fig. S24†) was recorded to probe the effect of defects on the charge transport kinetics. It showed that charge resistance ( $R_{\text{ct}}$ ) on  $\text{Co}_3\text{O}_4$ -1.0 was much lower than that on pristine  $\text{Co}_3\text{O}_4$ , which is favorable for enhancing the reaction rate.

The defect-engineering would serve as a promising strategy for simultaneously enhancing the chemisorption capability of

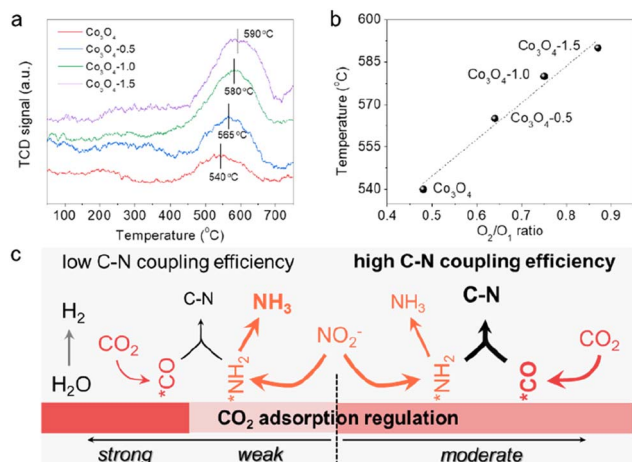


Fig. 3 (a) CO<sub>2</sub>-TPD plots for different catalysts normalized to ECSA. (b) The temperature of CO<sub>2</sub> desorption peak vs. the ratio of O<sub>2</sub>/O<sub>1</sub> over different catalysts. (c) Schematic diagram of CO<sub>2</sub> adsorption regulation for efficient C–N coupling.

these inert molecules and regulating the electronic structure of the catalyst to construct dual-functional sites. As effective chemisorption of reactants is essential for the initial stage of urea electrosynthesis, we turn to screen the effect of defects on the CO<sub>2</sub> chemical adsorption properties. The improved adsorption capacity of CO<sub>2</sub> molecules on V<sub>o</sub>-enriched Co<sub>3</sub>O<sub>4</sub> can be verified *via* the CO<sub>2</sub> temperature programmed desorption (CO<sub>2</sub>-TPD) spectra (Fig. 3a) normalized to ECSA. Comparably, all the defective Co<sub>3</sub>O<sub>4</sub> catalysts exhibited stronger binding strength and larger adsorption peaks than the pristine Co<sub>3</sub>O<sub>4</sub>. Typically, it was exhibited that CO<sub>2</sub> desorption occurred at around 580 °C on the Co<sub>3</sub>O<sub>4</sub>-1.0 catalyst, indicating that it had a strong CO<sub>2</sub> adsorption site with excellent CO<sub>2</sub> adsorption ability. In contrast, the CO<sub>2</sub> desorption peak of the pristine Co<sub>3</sub>O<sub>4</sub> catalyst was at around 540 °C, which was significantly weaker than that of the Co<sub>3</sub>O<sub>4</sub>-1.0 catalyst. We then correlated the temperature of the CO<sub>2</sub> desorption peak with the concentration of V<sub>o</sub> (the ratio of O<sub>2</sub>/O<sub>1</sub>) to investigate the correlation between V<sub>o</sub> and CO<sub>2</sub> adsorption capacity (Fig. 3b). This indicates that an increase in the concentration of V<sub>o</sub> in the catalyst leads to an increase in the CO<sub>2</sub> adsorption capacity. Combining the previous analysis of electrochemical performance over the as-prepared catalysts (Fig. 2), we can conclude that regulating the CO<sub>2</sub> adsorption capacity can control the competing reaction pathways (Fig. 3c). Moderating CO<sub>2</sub> adsorption is conducive to the adsorption and activation of CO<sub>2</sub> molecules to participate in the desirable C–N coupling process with 100% carbon efficiency.

To gain an in-depth understanding of the C–N coupling mechanism, *in situ* Raman spectroscopy and attenuated total reflection-surface-enhanced Fourier transformed infrared (ATR-FTIR) spectroscopy were carried out. Considering the potential window for urea production, potentials ranging from –0.5 to –0.9 V *vs.* RHE with 0.1 V intervals were selected for spectral acquisition during the *in situ* studies. As shown in Fig. 4a, five distinct peaks were observed in Raman spectra that correspond

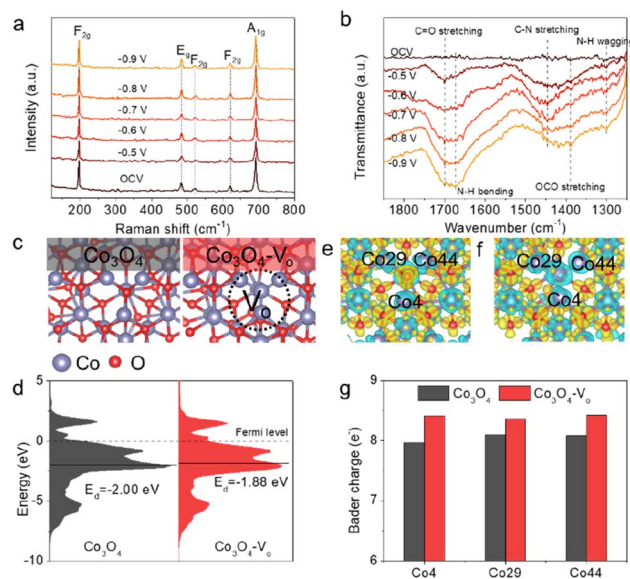


Fig. 4 *In situ* Raman (a) and ATR-FTIR (b) spectra recorded at different potentials for Co<sub>3</sub>O<sub>4</sub>-1.0 in CO<sub>2</sub> saturated 0.2 M KHCO<sub>3</sub> + 0.02 M KNO<sub>2</sub> electrolyte. (c) Schematic structures of Co<sub>3</sub>O<sub>4</sub> and Co<sub>3</sub>O<sub>4</sub>-V<sub>o</sub>. (d) The PDOS of the Co atoms in Co<sub>3</sub>O<sub>4</sub> and Co<sub>3</sub>O<sub>4</sub>-V<sub>o</sub> with the d-band center position (E<sub>d</sub>) marked by the black line and the Fermi level set as zero. Differential charge densities of (e) Co<sub>3</sub>O<sub>4</sub> and (f) Co<sub>3</sub>O<sub>4</sub>-V<sub>o</sub> structures. Yellow and blue contours represent electron accumulation and depletion, respectively. (g) Bader charge comparison of the Co atom around the V<sub>o</sub> in Co<sub>3</sub>O<sub>4</sub> and Co<sub>3</sub>O<sub>4</sub>-V<sub>o</sub>.

to the characteristic vibration peaks of Co<sub>3</sub>O<sub>4</sub>.<sup>32</sup> The intensity of the vibration peak did not change obviously at different applied potentials, suggesting that the Co<sub>3</sub>O<sub>4</sub> catalyst is stable during electrolysis. However, no reaction intermediate signal was recorded in the Raman spectra, which could be attributed to the strong Raman signal of Co<sub>3</sub>O<sub>4</sub> that overshadows the faint reaction intermediate signals. Alternatively, some intermediate signals were observed in ATR-FTIR spectra (Fig. 4b). The stretching vibrations observed at around 1700 and 1390 cm<sup>-1</sup> are attributed to the formation of \*CO and \*OCO intermediates, respectively,<sup>8,21</sup> indicating the CO<sub>2</sub> molecules can adsorb on the surface-active sites of the Co<sub>3</sub>O<sub>4</sub>-1.0 catalyst and further be converted into \*CO intermediates. The observation of peaks located at around 1675 and 1300 cm<sup>-1</sup> corresponded to the bending and wagging modes of the N–H bond, respectively, suggesting the reduction of NO<sub>2</sub><sup>-</sup> in the electrolysis.<sup>33</sup> Specifically, a typical peak at around 1445 cm<sup>-1</sup> was observed across all tested potentials, demonstrating that the C–N coupling was successfully achieved for urea production.<sup>8,21</sup> The enhanced peak intensity of the C–N bond at the potential of –0.7 V *vs.* RHE was consistent with the aforementioned experimental results, suggesting that the catalyst exhibited higher activity at this potential.

To strengthen our conclusion, we further conducted density functional theory (DFT) calculations. We built the correlative theoretical models, including Co<sub>3</sub>O<sub>4</sub> and Co<sub>3</sub>O<sub>4</sub>-V<sub>o</sub> (Fig. 4c and S25†), based on the above experimental characterizations. The partial density of states (PDOS) calculations on the Co atoms in

$\text{Co}_3\text{O}_4$  and  $\text{Co}_3\text{O}_4\text{-V}_\text{o}$  are shown in Fig. 4d. Notably, the d-band center position of Co atoms in the  $\text{Co}_3\text{O}_4\text{-V}_\text{o}$  structure ( $-1.88$  eV, relative to the Fermi level) was found to be higher than that in the  $\text{Co}_3\text{O}_4$  structure ( $-2.00$  eV). According to the d-band theory, metal-based catalysts with a higher d-band center (closer to the Fermi level) exhibit stronger interaction between the reaction intermediates and active sites, resulting in an enhanced C–N coupling process. To confirm this assumption, we conducted the simulations on the intermediates involved in  $\text{CO}_2$  and  $\text{NO}_2^-$  adsorption over the  $\text{Co}_3\text{O}_4$  and  $\text{Co}_3\text{O}_4\text{-V}_\text{o}$  structures. The adsorption structures of  $^*\text{CO}_2$  and  $^*\text{NO}_2$  on  $\text{Co}_3\text{O}_4$  (111) and  $\text{Co}_3\text{O}_4\text{-V}_\text{o}$  (111) surfaces are shown in Fig. S26 and S27.† Apparently, the adsorption free energy of  $^*\text{CO}_2$  on  $\text{Co}_3\text{O}_4\text{-V}_\text{o}$  ( $-0.31$  eV) is lower compared to that on  $\text{Co}_3\text{O}_4$  ( $-0.17$  eV), and the adsorption free energy of  $^*\text{NO}_2$  on  $\text{Co}_3\text{O}_4\text{-V}_\text{o}$  ( $-3.39$  eV) is also less than that on  $\text{Co}_3\text{O}_4$  ( $-2.88$  eV), indicating that the  $\text{CO}_2$  molecule and  $\text{NO}_2^-$  ion are more likely to be adsorbed and activated on the  $\text{Co}_3\text{O}_4\text{-V}_\text{o}$  surface. In addition, we pursued theoretical insights into the relationship between electronic structure and the electrocatalytic properties to deeply understand the impact of  $\text{V}_\text{o}$ . To gain further insights into the electronic properties of  $\text{Co}_3\text{O}_4$  and  $\text{Co}_3\text{O}_4\text{-V}_\text{o}$ , we analyzed the charge transfer within the catalyst structure. As depicted in Fig. 4e and f, we can find that the electrons of Co atoms were consumed in both  $\text{Co}_3\text{O}_4$  and  $\text{Co}_3\text{O}_4\text{-V}_\text{o}$ , owing to the lower electronegativity of the Co atom compared to the O atom. The detailed Bader charge analysis further revealed that in the  $\text{Co}_3\text{O}_4\text{-V}_\text{o}$  structure, the Co4, Co29 and Co44 atoms, which are located in close proximity to the  $\text{V}_\text{o}$ , had Bader charges of  $8.41$ ,  $8.37$  and  $8.43e^-$ , respectively. These values are higher than those observed in the  $\text{Co}_3\text{O}_4$  structure (Fig. 4g). This indicates that the presence of  $\text{V}_\text{o}$  in the  $\text{Co}_3\text{O}_4$  structure leads to the formation of an electron-rich local environment on the Co atoms, which is consistent with the XAS and XPS results. Our computational analysis highlights that the introduction of  $\text{V}_\text{o}$  in the catalysts can effectively regulate the electronic structure of the catalyst, leading to the enhancement of adsorption behaviors of intermediates and electrochemical activity for urea synthesis.

## Conclusions

In summary, we achieved urea electrosynthesis by co-reduction of  $\text{CO}_2$  and nitrite using the  $\text{V}_\text{o}$ -enriched  $\text{Co}_3\text{O}_4$  as the electrocatalyst. The as-prepared  $\text{Co}_3\text{O}_4\text{-1.0}$  catalyst demonstrated an outstanding urea yield rate of  $3361 \text{ mg h}^{-1} \text{ g}_{\text{cat}}^{-1}$  with a corresponding FE of 26.1% and 100% carbon selectivity at the potential of  $-0.7 \text{ V vs. RHE}$ . The remarkable catalytic performance was attributed to the integration of dual-functional sites in defective  $\text{Co}_3\text{O}_4$ , which can be regulated by the amount of  $\text{V}_\text{o}$ . Experimental measurements and theoretical calculations corroborated the role of  $\text{V}_\text{o}$  in constructing the well-matched adsorption/activation sites, which is beneficial for promoting the chemical adsorption of reactants, redistributing the electronic structure of the catalyst. The modulated local environment on Co atoms further enhanced the adsorption of intermediates and decreased the C–N coupling resistance in urea synthesis. Our study demonstrates a route for highly

efficient urea synthesis, and the methodology can be used for designing other electrocatalysts for the C–N coupling process.

## Data availability

The data that support the findings of this study are available in the ESI† of this article.

## Author contributions

P. S. L., Q. G. Z., and B. X. H. proposed the project, designed the experiments, and wrote the manuscript. P. S. L. performed all the experiments. P. S. L., J. Y. L., T. B. W., X. N. S., Q. L. M., X. C. K., and X. F. S. performed the analysis of experimental data. Q. G. Z. and B. X. H. co-supervised the whole project. All authors discussed the results and commented on the manuscript.

## Conflicts of interest

There are no conflicts to declare.

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