

Supporting Information

Understanding CeO₂ Modified Defective Carbon as Electrocatalyst for Electrochemical Reduction of CO₂

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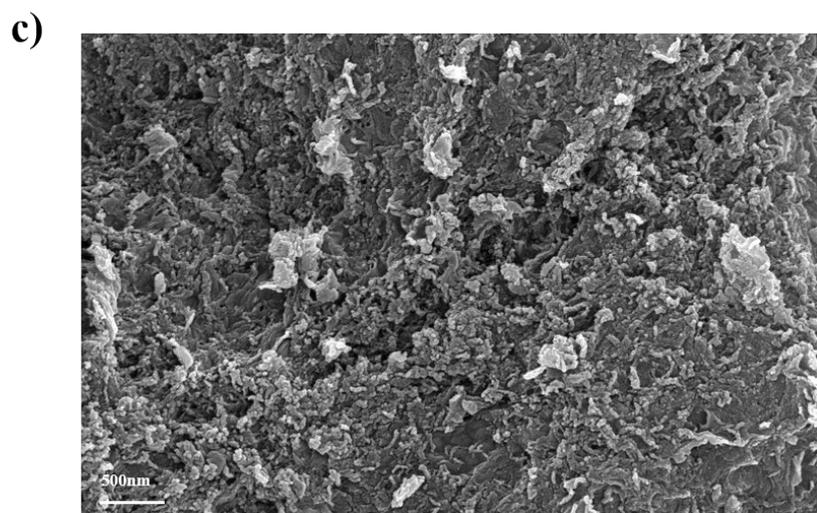
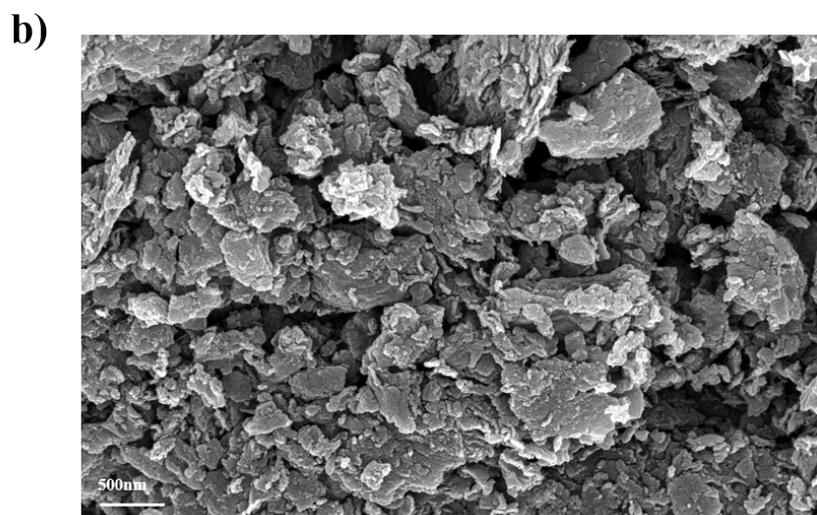
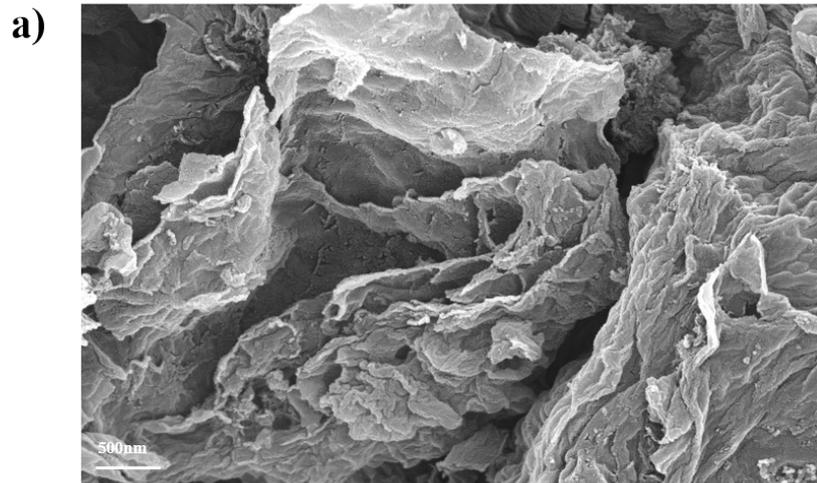


Figure S1. a) SEM image of V_0 -C-1300, b) SEM image of V_0 - CeO_2 /C-900, c) SEM image of V_0 - CeO_2 /C-1100.

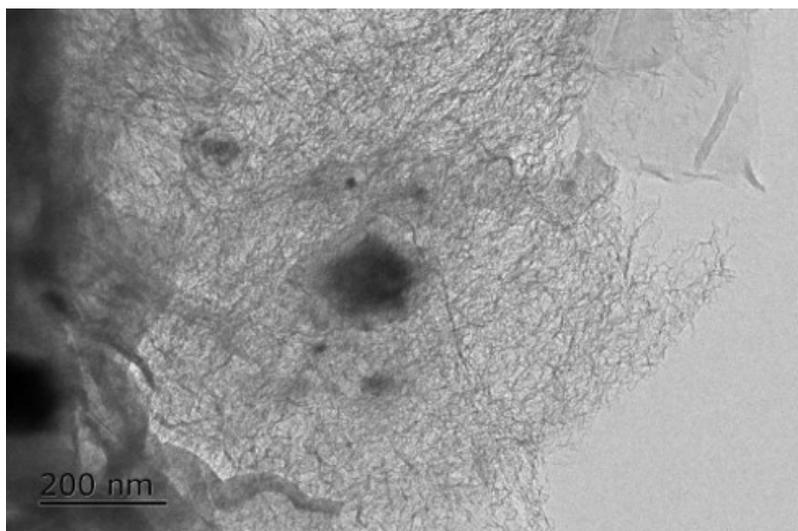


Figure S2. a) TEM image of V_0 - CeO_2 /C-1300.

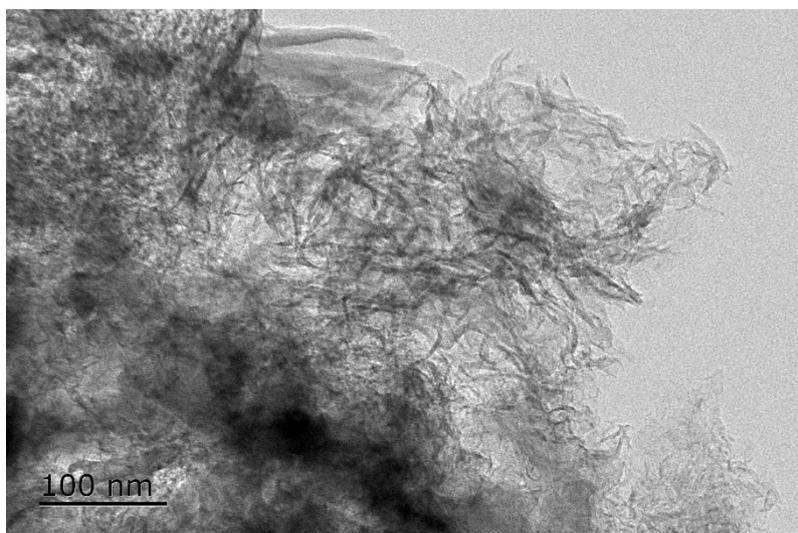


Figure S3. a) TEM image of V_0 - CeO_2 /C-900.

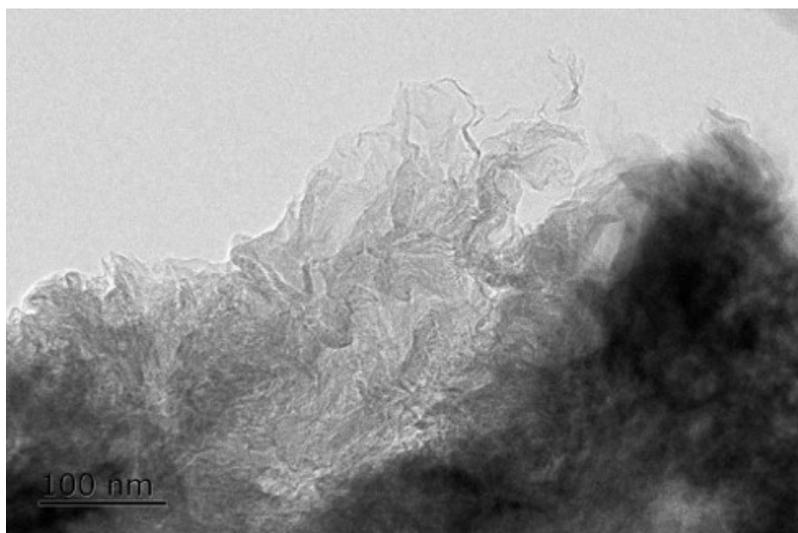


Figure S4. a) TEM image of V₀-C-1300.

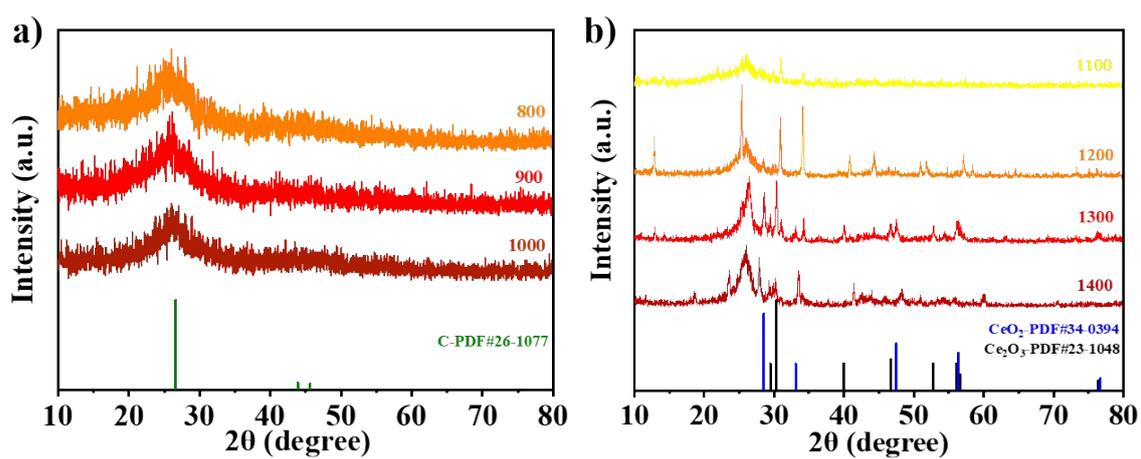


Figure S5. a-b) XRD of V₀-CeO₂/C at different temperatures.

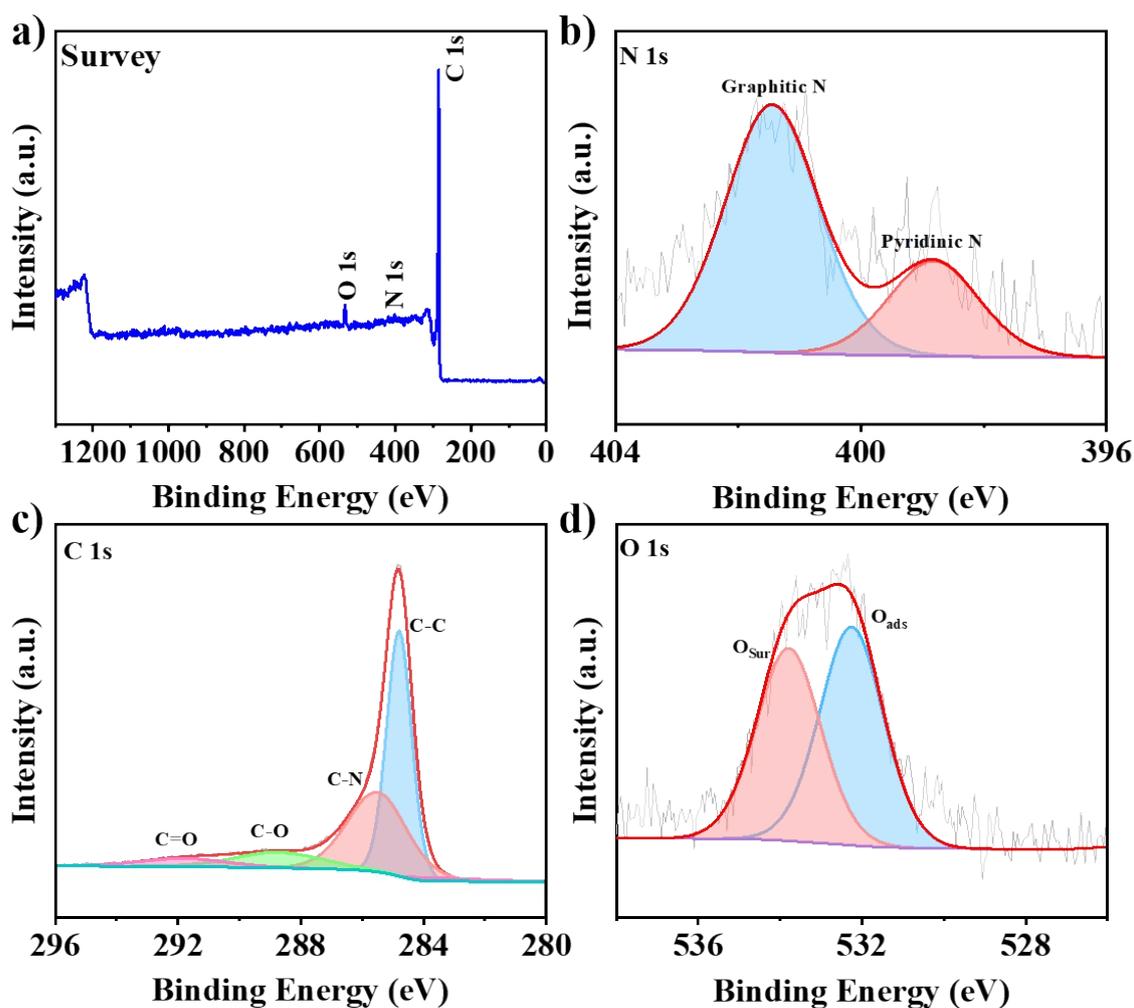


Figure S6. XPS spectra of V₀-C-1300: (a) survey spectra, (b) N1s, (c) C1s, (d) O1s.

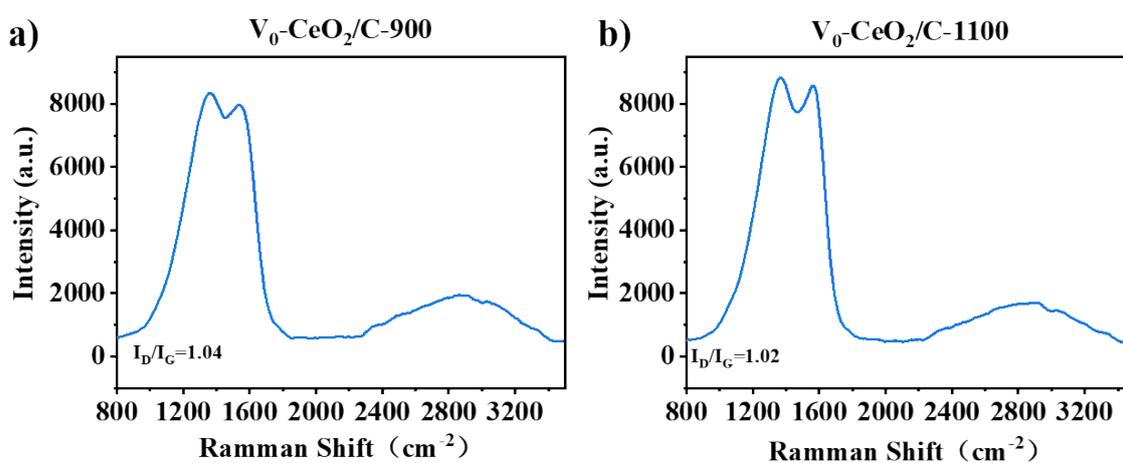


Figure S7. Raman spectrum of: a) V₀-CeO₂/C-900, b) V₀-CeO₂/C-1100.

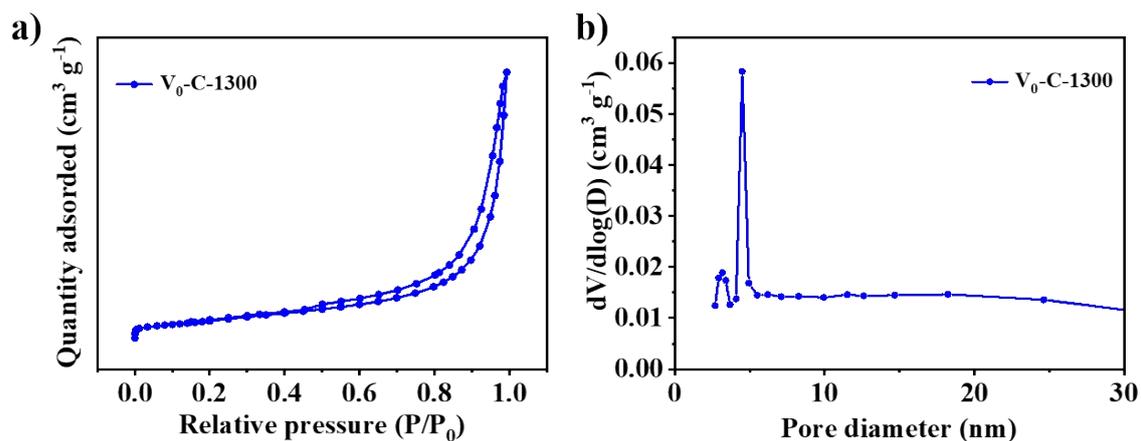


Figure S8. a) N₂ adsorption/desorption isotherms, and b) pore size distribution of V₀-C-1300.

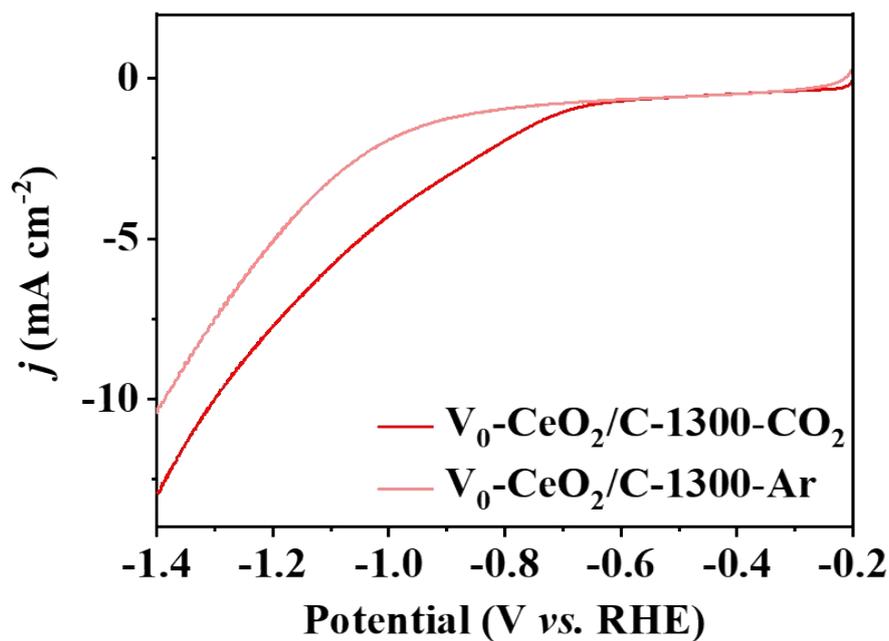


Figure S9. LSV curves of V₀-CeO₂/C-1300 in different atmospheres.

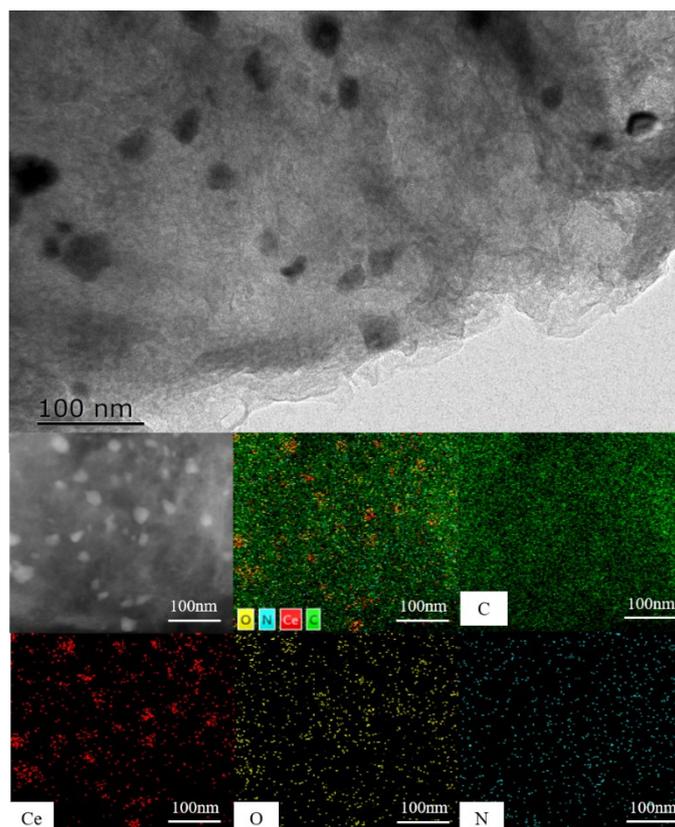


Figure S10. TEM image and EDS elemental mappings of $V_0\text{-CeO}_2/\text{C-1300}$ after stability test.

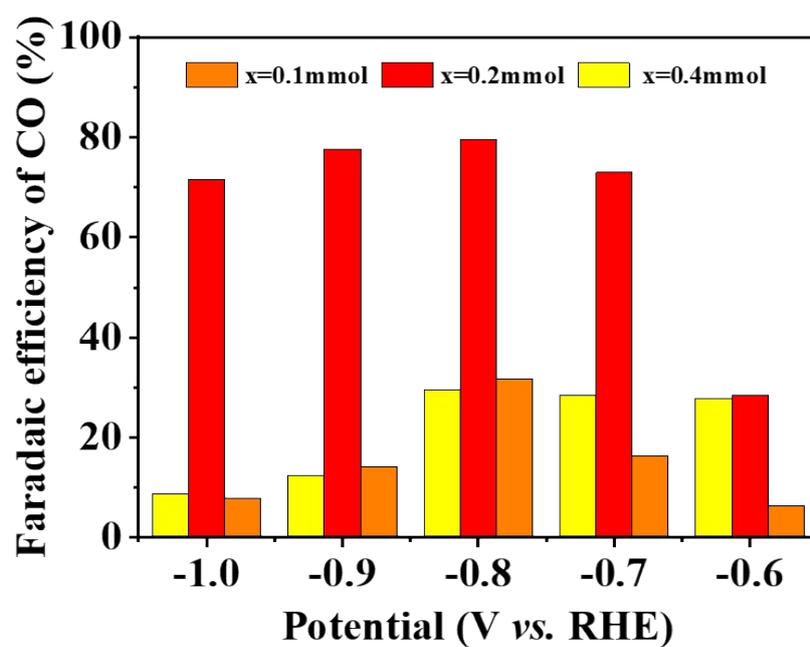


Figure S11. FE_{CO} of $V_0\text{-CeO}_2/\text{C-1300}$.

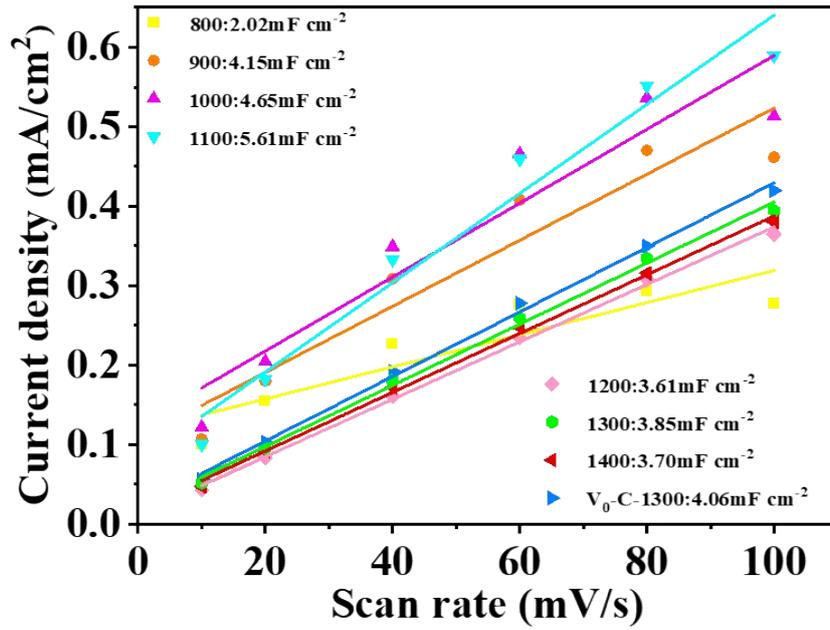


Figure S12. The electrochemical active area of $V_0\text{-CeO}_2/\text{C}$ at different temperatures and $V_0\text{-C-1300}$.

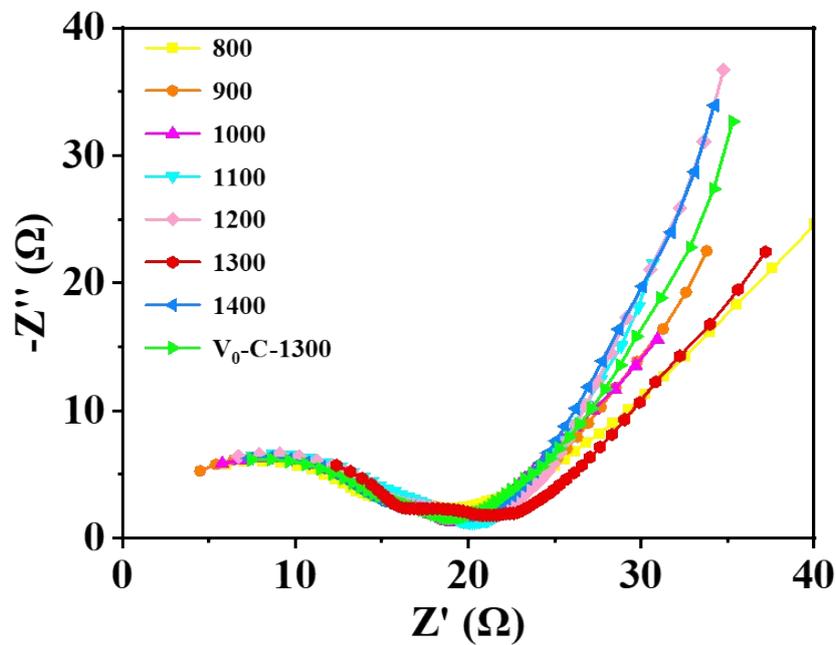


Figure S13 The electrochemical impedance spectroscopic (EIS) Nyquist plots of $V_0\text{-CeO}_2/\text{C}$ at different temperatures and $V_0\text{-C-1300}$.

Table S1. The EDS element content of V₀-C-1300.

Element	Line Type	k factor	Absorption Correction	Wt%	Wt% Sigma
C	K series	2.50675	1.00	95.14	0.45
N	K series	3.14061	1.00	0.48	0.34
Ce	L series	2.01965	1.00	4.39	0.30
Total:				100.00	

Table S2. EXAFS fitting parameters at the Ce L-edge for various samples ($S_0^2=0.75$ from CeO₂)

	shell	CN ^a	R ^b (Å)	σ^2 ^c (Å ²)	ΔE_0 ^d (eV)	R factor
CeO ₂	Ce-O	8	2.32±0.01	0.0083	6.4±1.1	0.0179
CeCl ₃	Ce-Cl	6.9±0.7	2.68±0.01	0.0107	-5.2±0.9	0.0175
	Ce-Cl	6.7±0.7	2.87±0.01	0.0107		
Ce-Sample	Ce-O	6.0±0.7	2.28±0.03	0.0202	3.3±2.1	0.0175

^aCN: coordination numbers; ^bR: bond distance; ^c σ^2 : Debye-Waller factors; ^d ΔE_0 : the inner potential correction. R factor: goodness of fit. Error bounds that characterize the structural parameters obtained by EXAFS spectroscopy were estimated as CN±20%; R ± 1%; σ^2 ± 20%.

Table S3.The catalyst contains rare earth elements and the main product of CO₂RR is CO.

Chemical formula	j/mA cm ⁻²	Product composition	Main products (FE)	Electrolyte	Electrolyzer type	Ref.
This work	12.6	CO H ₂	CO (79 %)	0.5M KHCO ₃	H-type cell	
Eu-N-C	16	CO H ₂	CO (59 %)	0.5M KHCO ₃	H-type cell	[1]
Pr-NC-8	~0.25	CO H ₂	CO (93 %)	0.5M KHCO ₃	standard three electrode system	[2]
La _{0.6} Sr _{0.4} Fe _{0.8} Ni _{0.2} O _{3-δ}	193	CO H ₂	CO (99 %)	/	symmetrical half-cell	[3]
ZnLa/CN	2.5	CO H ₂	CO (26 %)	0.1M KHCO ₃	standard three electrode system	[4]
Y/NC and Sc/NC	~2	CO H ₂	CO(88 %)/ CO(81%)	0.5M KHCO ₃	H-type cell	[5]
Ce _{0.016} Zn _{0.984} O	24	CO H ₂	CO (88 %)	0.5M KHCO ₃	H-type cell	[6]

Reference

- [1] X. Zeng, L. Liao, M. Wang, H. Wang, Rare-earth metal-N₆ centers in porous carbon for electrocatalytic CO₂ reduction, *Physical Chemistry Chemical Physics* 25 (2023) 20381-20394.
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- [5] J Liu, X Kong, L Zheng, X Guo, X Liu, J Shui, Rare earth single-atom catalysts for nitrogen and carbon dioxide reduction[J]. *ACS nano*, 2020, 14(1): 1093-1101.
- [6] X Ren, Y Gao, L Zheng, Z Wang, P Wang, Z Zheng, Y Liu, H Cheng, Y Dai, B Huang, Oxygen vacancy enhancing CO₂ electrochemical reduction to CO on Ce-doped ZnO catalysts[J]. *Surfaces and Interfaces*, 2021, 23: 100923.