Electronic Supplementary Information



Fig. S1. (a-d) TEM images of Au@Co-Ce prepared at different reaction times: a) 120 s, b) 125 s, c) 4 min, d) 10 min; The insets show STEM-EDX elemental maps of corresponding samples.

The addition of ammonia can trigger the redox self-assembly process to result in formation of the Au@Co₃O₄ multicore-shell nanospheres (Fig. S1a) after reacting for 120s. When Ce(NO₃)₃ is added at 120 s, Ce³⁺ rapidly reacts with the Co₃O₄ on the surface to form CeO₂ on the outermost shell. After etching with Ce³⁺ for 5 s, the core-shell structure remain unchanged (Fig. S1b). However, the EDX element mapping (Fig. S1b inset) reveals the CeO₂ is already form on the outermost shell. As the reaction proceeds, the electrons released from Ce³⁺ migrate onto the Au surface to etch the Co₃O₄ around Au NPs.¹ Accompanied by the dissolution of Co₃O₄ and driven by thermodynamic equilibrium, the Au NPs begin to aggregate gradually, and the small void spaces around Au form slowly (Fig. S1c). In addition, because the shell is a stack of metal oxide particles, many of the pinholes should be present in the

interstice of the metal oxide particles. These pinholes not only help Ce^{3+} to diffuse into the interior of the nanosphere but also improve the dissolution of the internal Co_3O_4 .¹ After etching for 10 min, the apparent yolk-shell structure and large Au core (Fig. S1d) are formed due to the massive dissolution of Co_3O_4 .



Fig. S2. A) SEM image, B) TEM image, C) XRD pattern and D) HRTEM image of Au@Co₃O₄ multicore-shell nanosphere.



Fig. S3. XRD pattern of Au@Co-Ce MOYSNs.



Fig. S4. (a) XPS, (b), (c) high-resolution XPS for (b) Co 2p and (c) Fe 2p, (d) XRD pattern of Au@Co-Fe MOYSNs.



Fig. S5. A) SEM image, B) TEM image, C) STEM-EDX profile, the inset is the result of ICP, D) STEM-EDX elemental maps, E) XRD pattern of Au@Co₃O₄/CeO₂/Fe₂O₃ sample.



Fig. S6. TEM image of sample 2 (Au@Co-Ce multicore@shell nanosphere)



Fig. S7. a) TEM image, b) Au size distribution histogram, c) nanosphere size distribution histogram, d) N_2 adsorption-desorption isotherms of sample 4

The results show that the BET surface area of sample 4 is 59.8 m2 g-1 and pore diameter is 19.53 nm.



Fig. S8. TEM image of sample 5.



Fig. S9. TEM images of (A) Au nanoparticle, (B) Co_3O_4 , (C) CeO_2 nanoparticles.



Fig. S10. TEM images of Co-Ce mixture prepared by co-precipitation process.

ICP /wt%	Sample 1	Sample 2	Sample 3	Sample 4	Sample 5	Sample 7
Au	33.2%	32.3%	32.8%	33.3%	31.6%	0%
Co	65.8%	32.6%	19.8%	13.2%	2.4%	19.4%
Ce	0.0%	35.1%	47.4%	53.5%	66.0%	80.6%

 Table S1. ICP results of Sample 1-5 and 7.



Figure S11. The H_2 -TPR curves of sample 1-5 and sample 7.



Fig. S12. TEM image of sample 4 after catalytic test

Catalyst	Amount (mg)	Size of Au (nm)	Temperature for 100 % CO conversion(°C)	space velocity (cm ³ ·h ⁻¹ ·g ⁻¹)	reference
Au@CeO2core-shell	200	17	155	15000	ref. 2
Au@ZrO2 yolk-shell	50	15-17	240	80000	ref. 3
Au@SnO2 yolk-shell	50	40	230 (for 50%)	84000	ref. 4
Au@CeO2/ZrO2yolk-shell	none	3-5	120	120000	ref. 5
Au@SiO2 core-shell	30	1.5	180	30000	ref. 6
Au@MnO ₂ yolk-shell	30	10-15	130	60000	ref. 7
Au@Co-Ce yolk-shell	30	35	125	60000	this work

Table S2. Comparison of the Size of Au, temperature for 100 % CO conversion and space velocity for CO oxidation with other oxideencapsulated Au catalysts.

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