

Supporting information

The effects of SO₂ impurities on CO₂ electroreduction on bare silver and SiO₂ coated silver in different cell geometries

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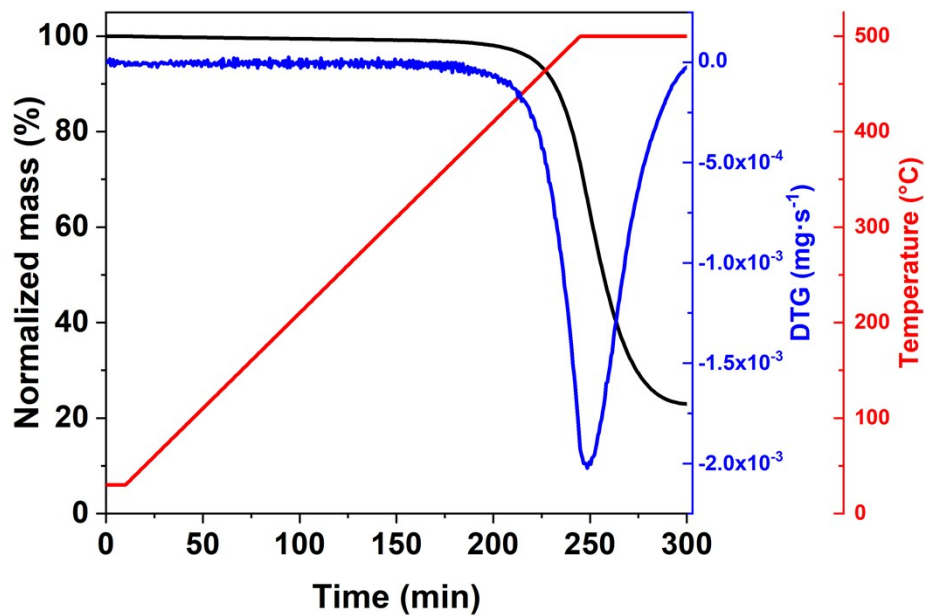


Figure S1. TG/DTA analysis of the original Ag/CB catalyst.

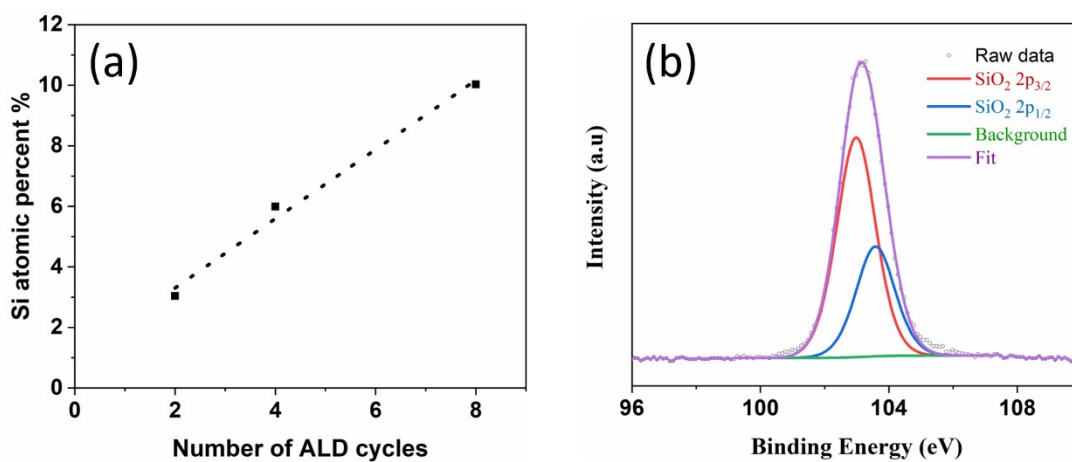


Figure S2. (a) Growth per ALD cycle of SiO₂ deposition on the surface of Ag/CB catalyst. (b) Si 2p XPS spectra of the Ag/CB catalyst with 8 cycles of SiO₂ coating.

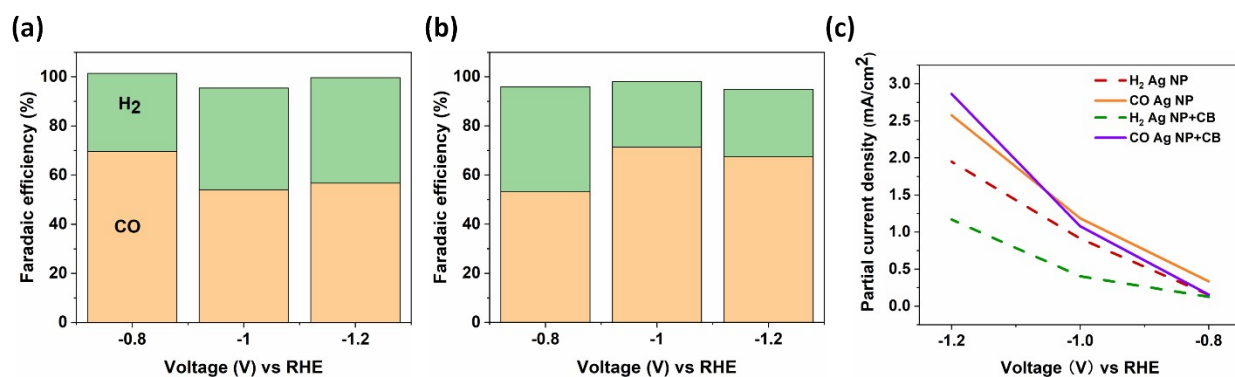


Figure S3. (a) Pure Ag nanoparticles drop casted on glassy carbon electrode tested at H-cell. (b) Ag nanoparticles mixed with carbon black to mimic the Ag/CB catalyst drop casted on glassy carbon electrode tested at H-cell. (c) Partial current density of CO and H₂ from pure Ag nanoparticles and Ag nanoparticles mixed with carbon black.

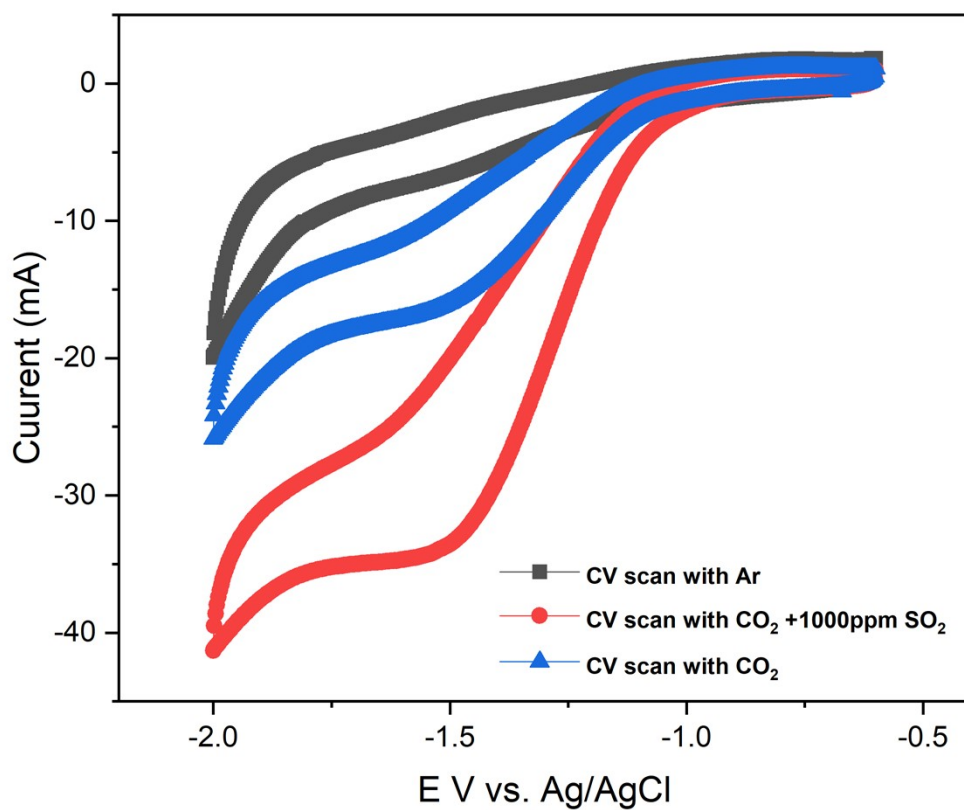


Figure S4. Cyclic voltammetry tests under various gas supplies: pure Ar, CO₂ with 1000ppm SO₂ and pure CO₂

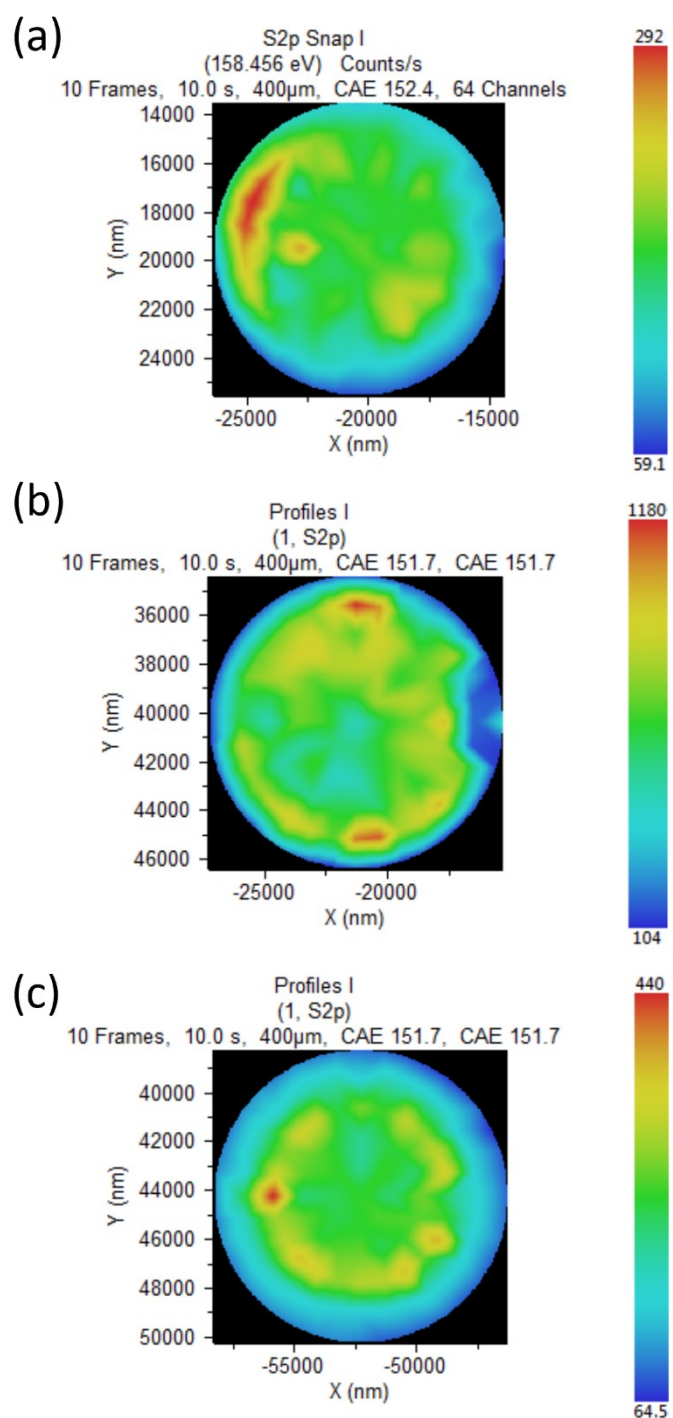


Figure S5. XPS area scan of the Ag/CB catalyst tested at H-cell. (a) uncoated Ag/CB catalyst after 1h CO₂ electrolysis at H-cell with pure CO₂. (b) uncoated Ag/CB catalyst after 1h CO₂ electrolysis at H-cell with 1000 ppm SO₂. (c) 8 cycles SiO₂ coated Ag/CB catalyst after 1h CO₂ electrolysis at H-cell with 1000 ppm SO₂.

Method to calculate the SO₂ concentration required to get SO₂ reduction

$$FE_{missing} = FE_{in\ total\ (Ag/CB\ with\ pure\ CO_2\ feed)} - FE_{in\ total\ (Ag/CB\ with\ CO_2 + SO_2\ feed)}$$

$$FE_{in\ total} = FE_{H_2} + FE_{CO}$$

$$I_{missing} = FE_{missing} \times I_{in\ total}$$

$$I_{SO_2\ RR\ partial\ current} = I_{missing}$$

$$Required\ flow\ rate_{SO_2} = \frac{I_{SO_2\ RR\ partial\ current} \times 6.242 \times 10^{18}}{N_{number\ of\ electrons} \times N_A} \times V_{25^\circ C}$$

$$V_{25^\circ C} = \frac{nRT_{25^\circ C}}{P} = 24.465L/mol$$

$$N_A = 6.02 \times 10^{23} mol^{-1}$$

$$N_{number\ of\ electrons} = 6$$

$$Required\ C_{concentration\ SO_2} = \frac{Required\ flow\ rate_{SO_2}}{Flow\ rate_{in\ total}} \times 100\% \times \frac{1 \times 10^6\ ppm}{\%}$$

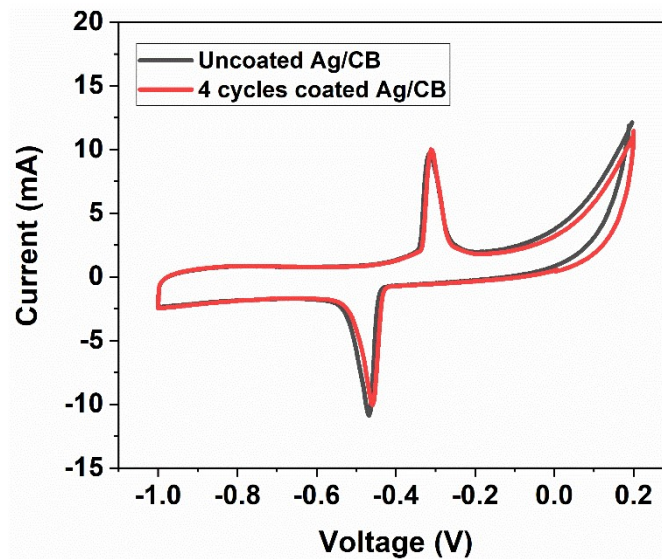


Figure S6. CV testing of the uncoated Ag/CB catalyst and the catalyst with 4 cycles of SiO₂ coating, tested at MEA reactor.

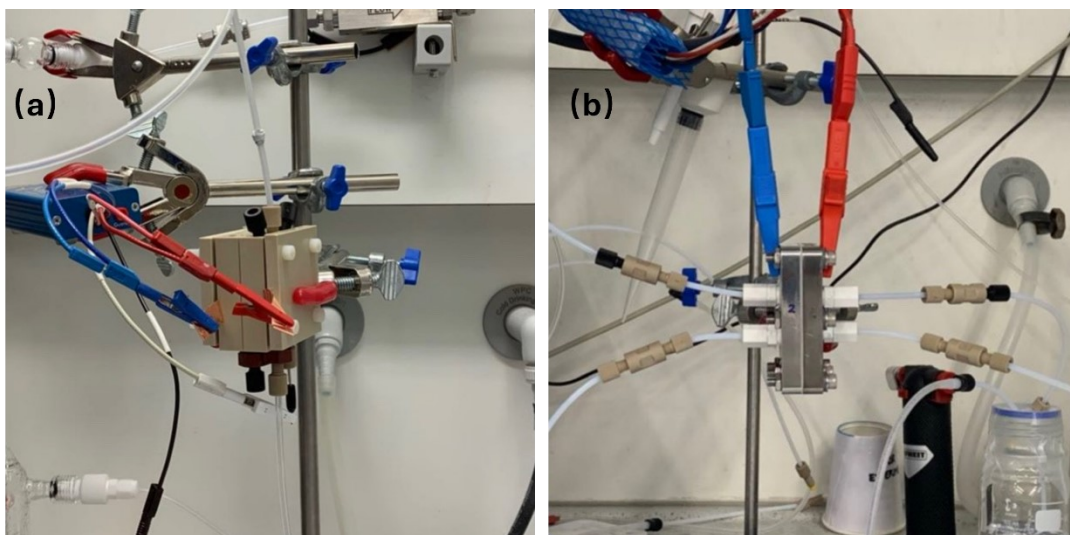


Figure S7. A visual representation of both cell geometries. (a) H-cell (b) MEA

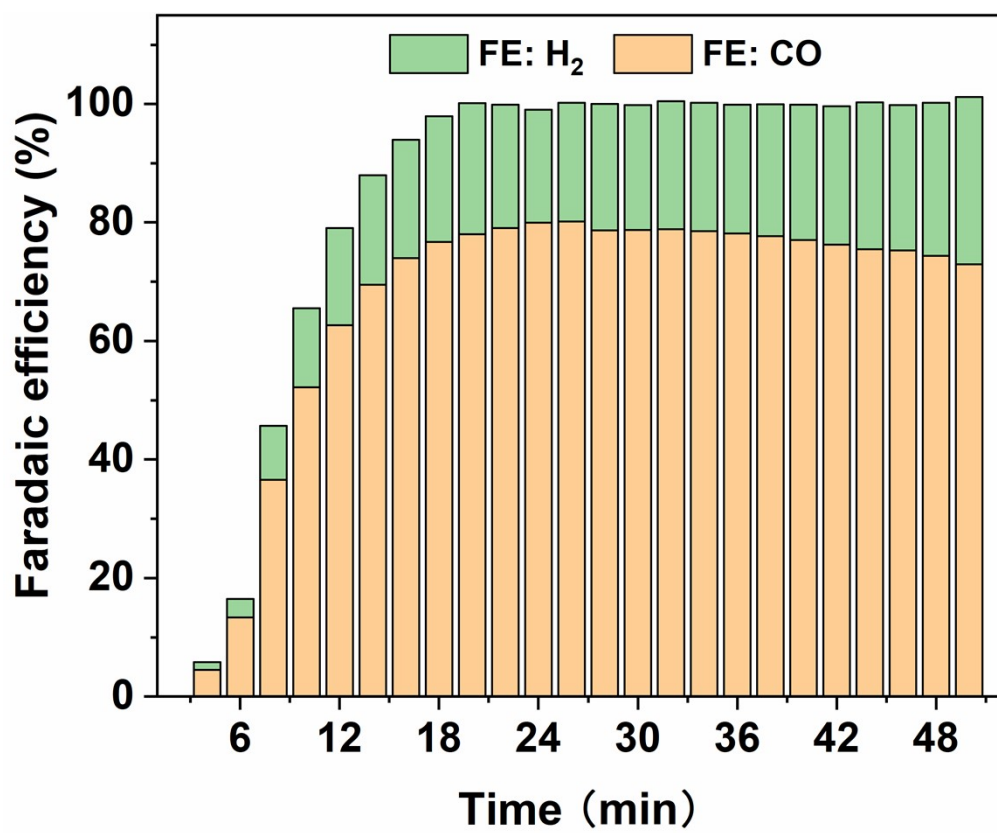


Figure S8. Faradaic efficiency of the original Ag/CB catalyst tested at MEA reactor at 100 mA/cm² current density in a 1 M KHCO₃ electrolyte with pure CO₂ feed for 50 minutes.

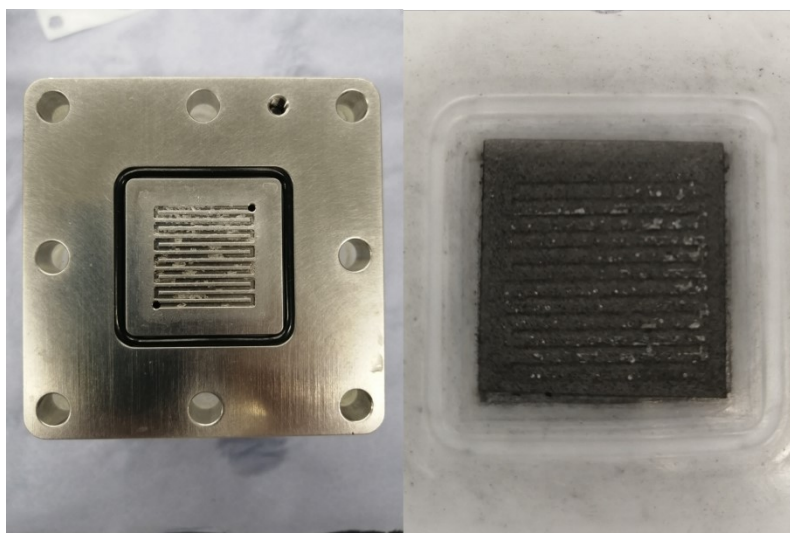


Figure S9. Salt formation after 50 minutes CP testing of the uncoated Ag/CB catalyst at 100 mA/cm^2 .

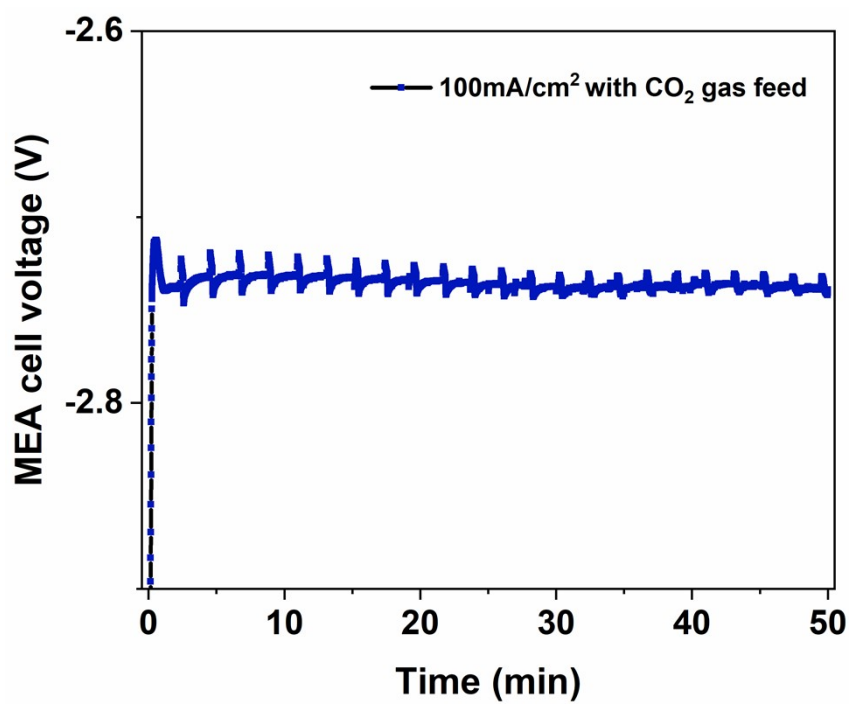


Figure S10. CP testing of the uncoated Ag/CB catalyst with pure CO_2 gas feed at 100 mA/cm^2 .

Table S1. ALD experimental conditions. (RT indicates room temperature)

Material	Reactant (T,°C)	Co-reactant (T,°C)	Type of reactor	T _{reaction} (°C)	Pulse and Purge Time (s) (SiCl ₄ -N ₂ -H ₂ O- N ₂)	Flow rate(L/min)
SiO ₂	SiCl ₄ (RT)	H ₂ O vapor (RT)	Flat substrate reactor	100	15-60-30-60	0.5-0.5-0.5- 0.5
SiO ₂	SiCl ₄ (RT)	H ₂ O vapor (RT)	Fluidized bed reactor	100	60-150-60-150	0.5-2-2-2

Table S2. Gas recipes for H-cell testing and MEA testing

Gas composition	Types of reactors	Gas flow rate (sccm)		
		CO ₂	CO ₂ + 100 ppm SO ₂	CO ₂ + 10000 ppm SO ₂
CO ₂	H-cell	8	0	0
CO ₂ + 10 ppm SO ₂	H-cell	7.2	0	0.8
CO ₂ + 100 ppm SO ₂	H-cell	0	8	0
CO ₂ + 1000 ppm SO ₂	H-cell	7.2	0	0.8
CO ₂	MEA cell	20	0	0
CO ₂ + 1000 ppm SO ₂	MEA cell	18	0	2