

Supporting Information for

Electrochemical reduction of CO₂ using palladium modified boron-doped diamond electrodes: enhancing the production of CO

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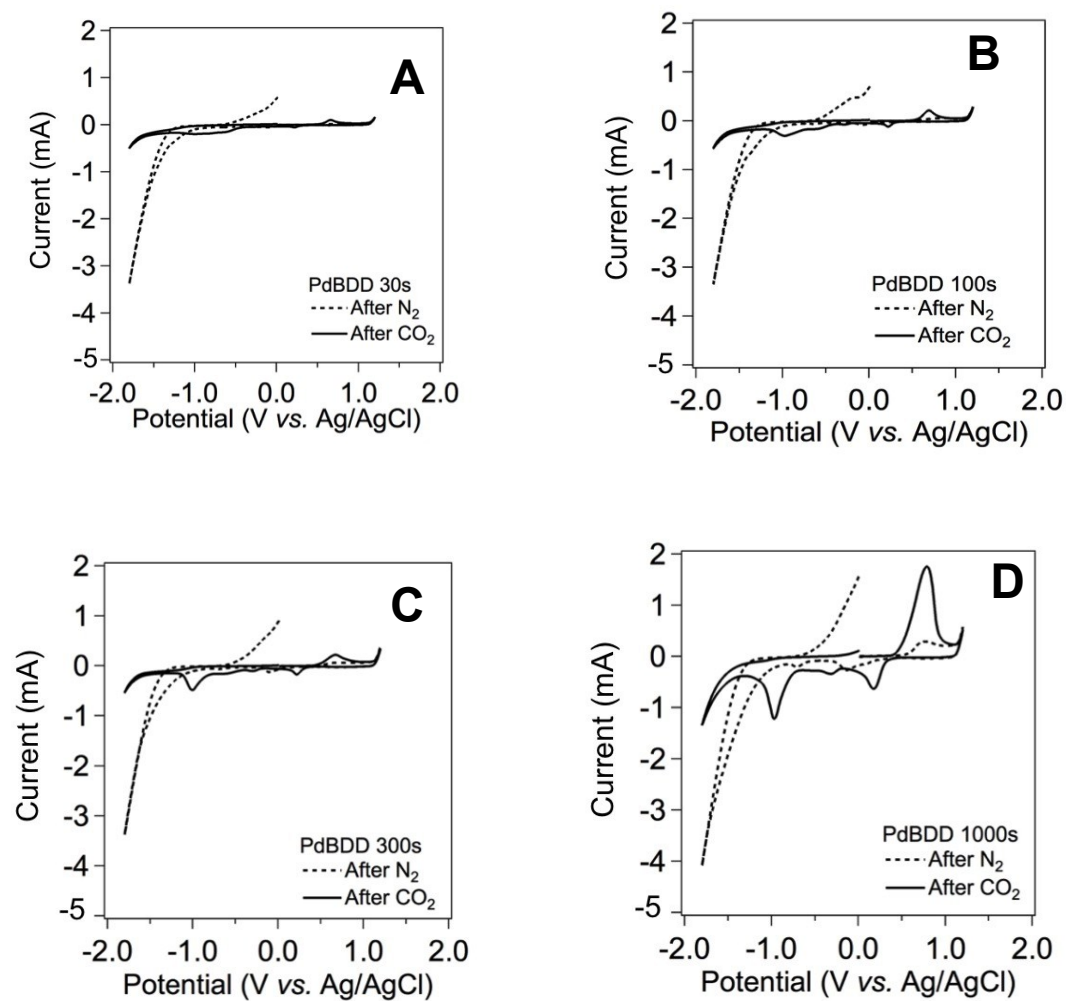


Figure S1. CVs of PdBDD30 (A), PdBDD100 (B), PdBDD300 (C), and PdBDD1000 (D) after N₂ and CO₂ bubbling, performed in 0.1 M NaCl with a scan rate of 100 mV/s, and potentials ranging from -1.8 V to 1.2 V.

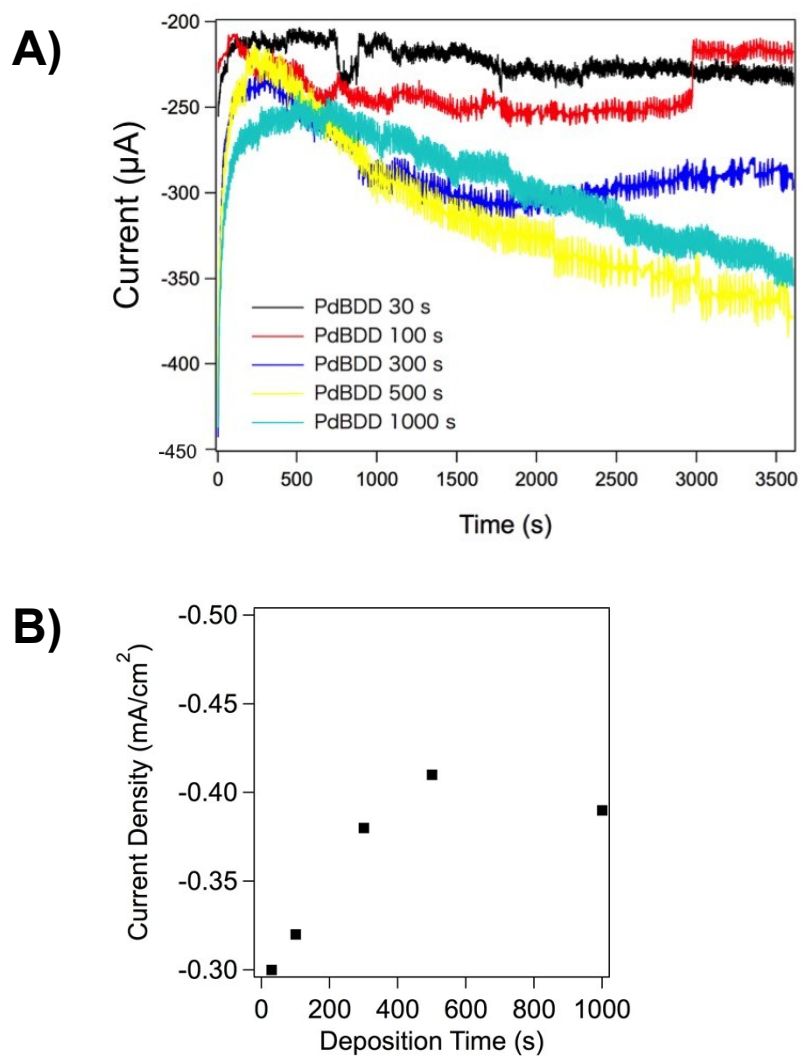


Figure S2. Chronoamperometry for the electrochemical reduction of CO_2 (A) and the current density at a potential of -1.5 V (B) using PdBDD electrodes with different Pd deposition time.

Table S1. Faradaic efficiencies for the products produced using bare BDD and Pd metal electrodes

Electrode	Potential/V	Faradaic Efficiency/%		
		HCOOH	CO	H ₂
Bare BDD	-1.6	5.76	9.95	84.92
Pd metal*	-1.6	0.70	9.34	8.53

*The total faradaic efficiency of the products with the Pd metal electrode is far from 100%. This might be due to dissolution of H₂ into the electrode [1].

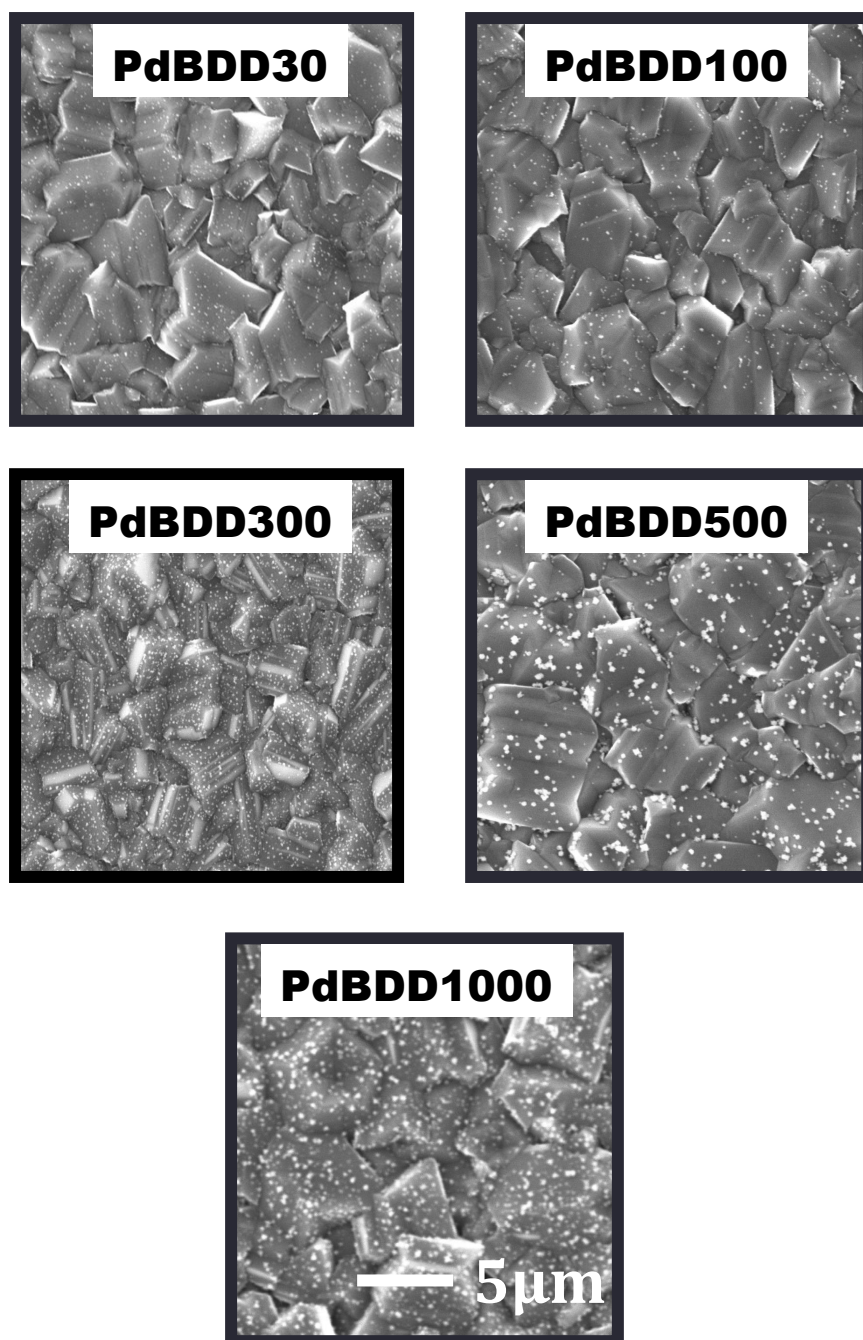


Figure S3. SEM images of PdBDD electrodes after the electrochemical reduction of CO_2 at a potential of -1.5 V for 1 hour.

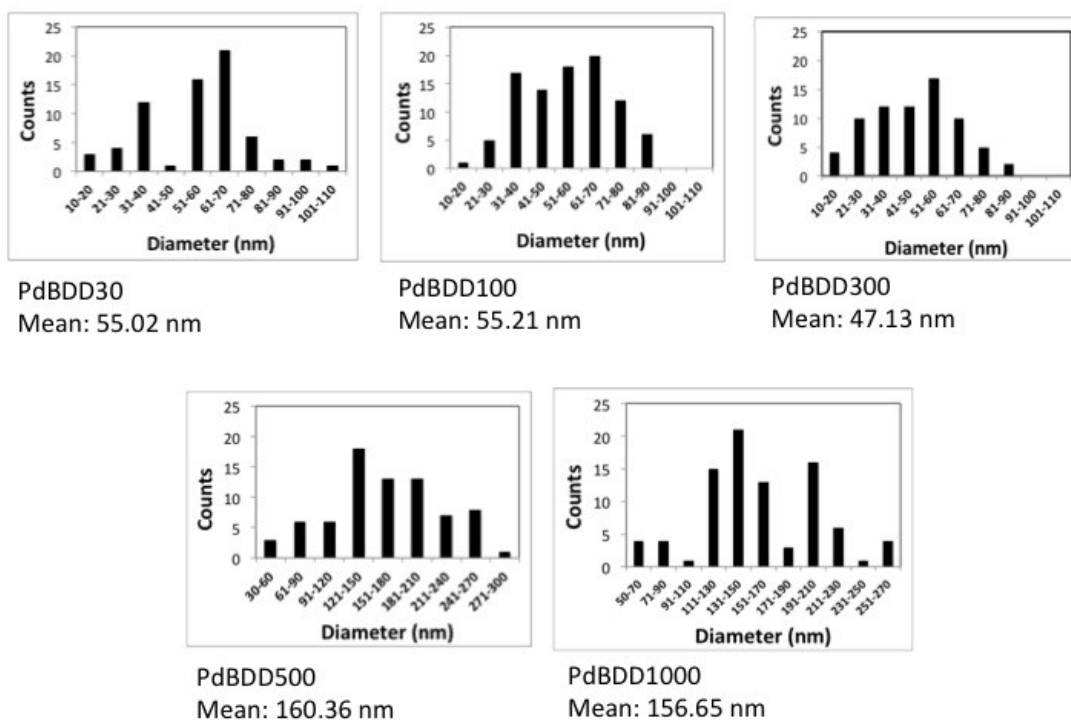


Figure S4. Pd particle size distribution after electrochemical deposition at a potential of -0.15 V using electrodes with different deposition times

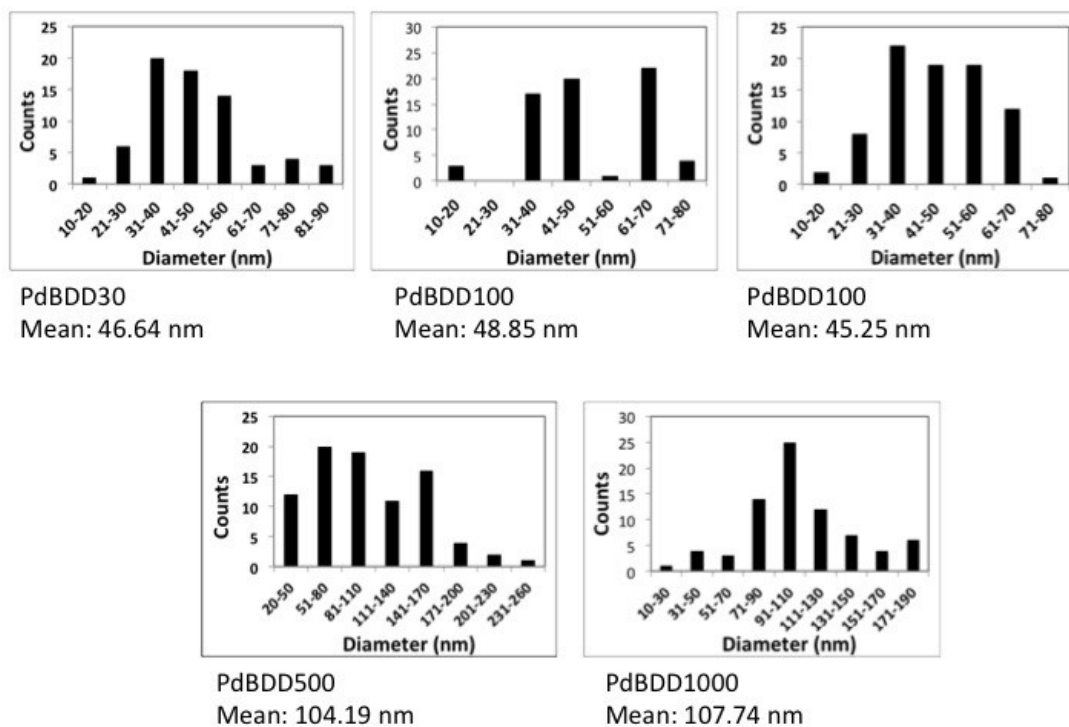


Figure S5. Pd particle size distribution on the different PdBDD electrodes, after electrochemical reduction of CO₂ at a potential of -1.5 V for 1 hour.

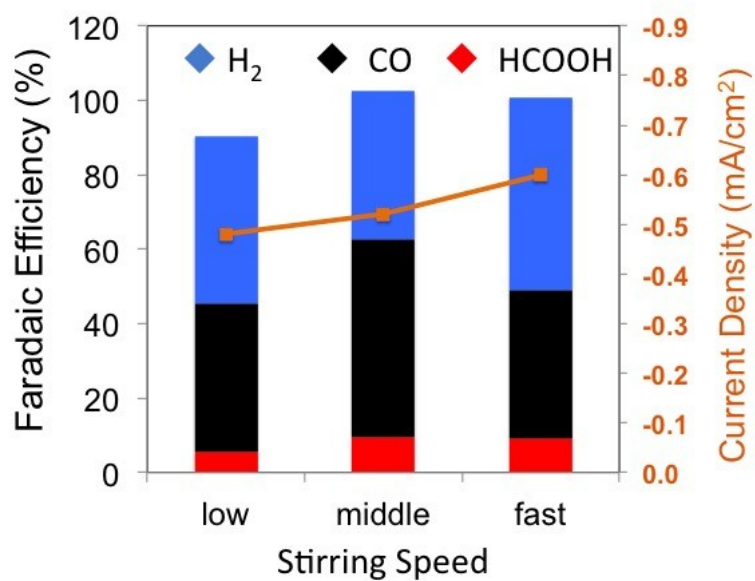


Figure S6. Stirring speed dependence of the faradaic efficiencies of the products and the current density during the electrochemical reduction of CO₂ carried out using PdBDD300 at a potential of -1.6 V for 1 hour.

Reference

- [1] Y. Hori, H. Wakebe, T. Tsukamoto, O. Koga, *Electrochem. Commun.* **1994**, *39*, 1833–1839.