Polymer coating accelerates fuel production

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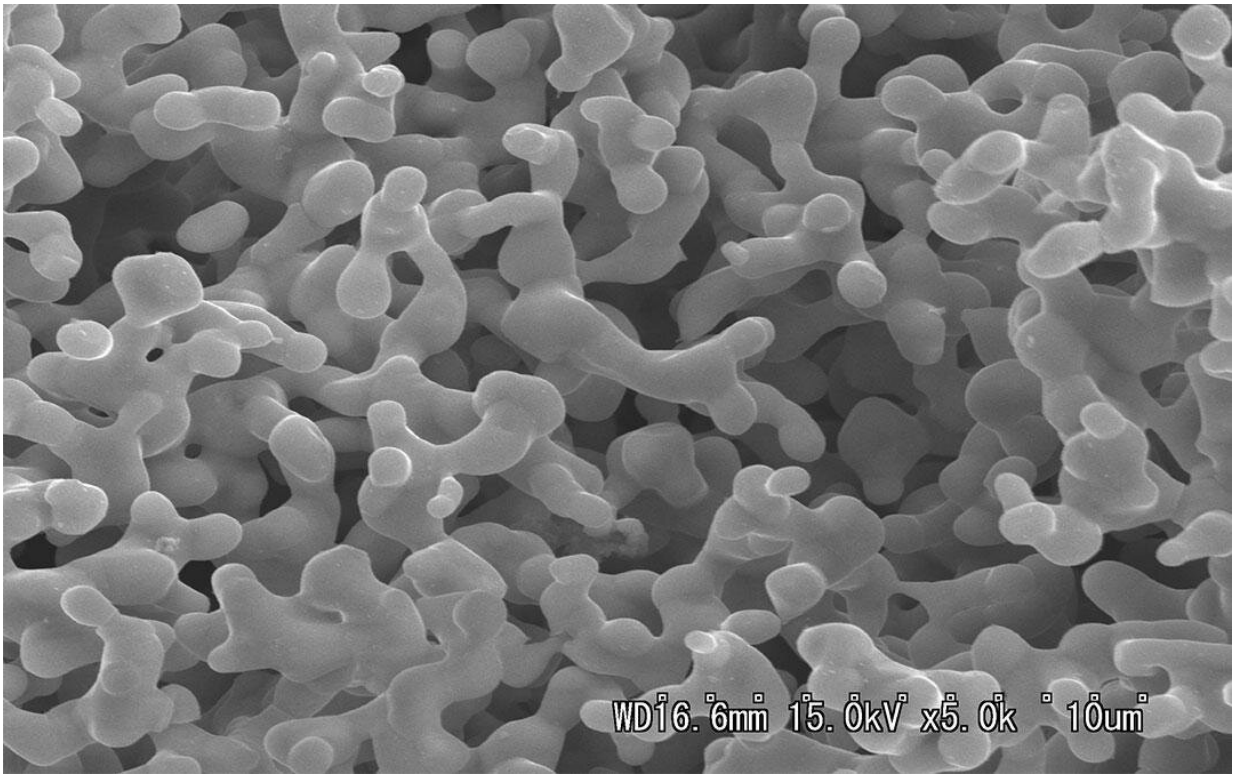

Researchers from the University of Tsukuba and Osaka University find that a polymer coating can effectively shuttle CO2 molecules to a metal catalyst, thus accelerating its product formation and offering potential greenhouse gas recycling strategies. Credit: University of Tsukuba

It is well-established that the accumulation of greenhouse gases, like carbon dioxide (CO_2), in the atmosphere contributes to climate change. Therefore, CO_2 capture and recycling are vital for mitigating detrimental

environmental effects and addressing the climate crisis. Recently, researchers from Japan designed a polymer-coated metal catalyst that accelerates CO_2 conversion and offers green energy insights.

In a study published in *ACS Catalysis*, researchers from the University of Tsukuba describe porous tin (Sn) catalysts coated with polyethylene glycol (PEG) and show how this polymer facilitates CO_2 transformation into a useful carbon-based fuel.

Various polymers can capture CO_2 molecules, and Sn catalysts are known to reduce CO_2 to other molecules, like formate (HCOO-), which can be reused to power fuel cells.

"We were interested in combining these capabilities into a single catalytic system that could scrub CO_2 from its surroundings and recycle it into formate," says research group leader, Professor Yoshikazu Ito. "However, it's difficult to obtain only the desired product, formate, at a high production rate and in high yield, so we had to fine-tune the catalyst design." The formate production rate of PEG-coated Sn was 24 times higher than that of a conventional Sn plate electrode, and no byproducts were detected (>99% yield of formate). To understand this enhanced CO_2-reduction reaction, the researchers fabricated an Sn catalyst coated with another CO_2-capturing polymer (polyethyleneimine; PEI) whose structure interacts differently with incoming CO_2. The PEG-coated Sn still outperformed the PEI-coated Sn, and considering the chemical characteristics of these polymers, the authors proposed that PEI held the CO_2 molecules too tightly, whereas PEG struck a key balance in capturing and then releasing CO_2 to the catalytic Sn core.

"Modeling this reaction using theoretical computations confirmed the favorability of PEG shuttling CO_2 to the Sn center and explained the accelerated formate production," explains Ph.D. student, Samuel Jeong. "However, we wanted to further clarify the PEG-CO_2 interactions."

More detailed computations revealed that while the absence of polymer limits the Sn catalyst's CO_2-capture ability, an excessively dense layer of PEG inhibits CO_2 transfer to the metal surface, thereby decreasing formate production. Therefore, a complete but relatively sparse layer of PEG is optimal for funneling CO_2 to Sn, while maintaining a CO_2-rich environment and preventing byproduct release.

The mantra "reduce, reuse, recycle" no longer only refers to single-use plastics. The simple catalyst-coating technique reported by Ito and co-workers can be used to develop systems that efficiently recycle CO_2 into useful compounds, like formate, which can power fuel cell devices that provide green electricity.

More information: Samuel Jeong et al, Polyethylene Glycol Covered Sn Catalysts Accelerate the Formation Rate of Formate by Carbon Dioxide Reduction, *ACS Catalysis* (2021). DOI: 10.1021/acscatal.1c02646

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