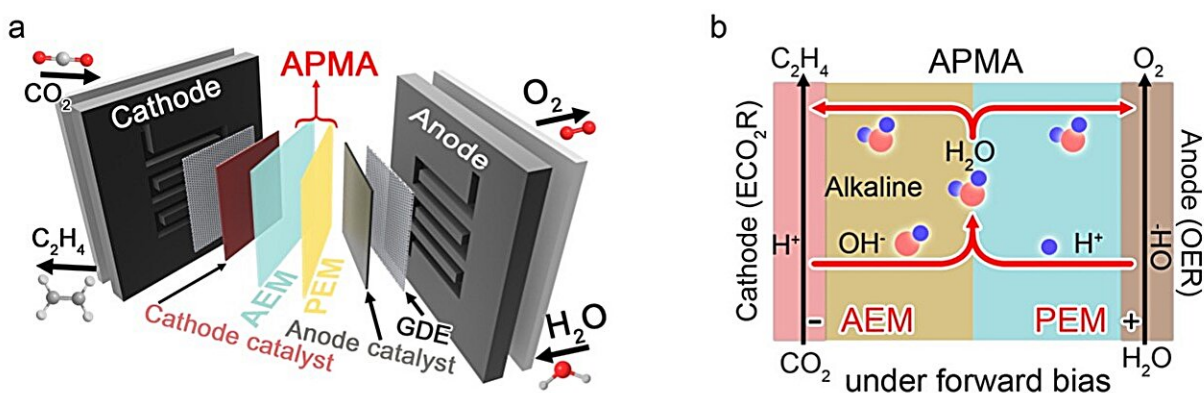


# A pure water-fed membrane-electrode-assembly system for electrocatalytic reduction of carbon dioxide

January 22 2024, by Ingrid Fadelli



(a) The schematic of the APMA MEA system architecture for ECO<sub>2</sub>R. (b) The schematic diagram of the ECO<sub>2</sub>R reaction mechanism in the APMA system with an alkaline cathode environment under forward bias mode. Credit: She et al

The sustainably powered, electrochemical reduction of carbon dioxide (CO<sub>2</sub>) into useful chemicals and feedstock could help to mitigate greenhouse gas emissions, allowing industries to reuse released CO<sub>2</sub> in beneficial ways. Most of the strategies for realizing this introduced so far, however, have notable limitations, including a poor stability over long periods of time.

Researchers at Hong Kong Polytechnic University, University of Oxford

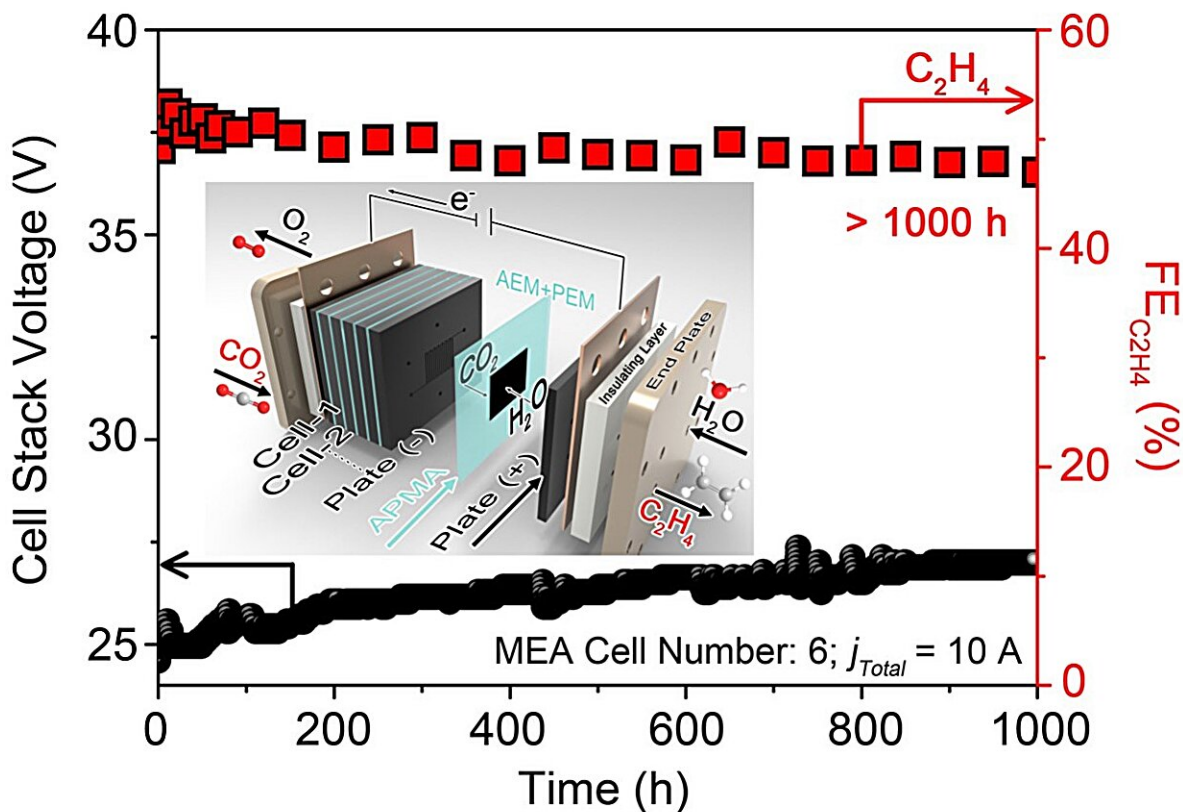
and the National Synchrotron Radiation Research Center recently introduced a new membrane-electrode-assembly system that could facilitate the stable electrocatalytic reduction of CO<sub>2</sub>.

Notably, their proposed system, which was first introduced in a paper [published](#) in *Nature Energy*, is fed by pure water (H<sub>2</sub>O) and thus does not rely on alkali-metal electrolyte.

"Our [modern society](#) heavily depends on [fossil fuels](#) to power our economy, but the resulting CO<sub>2</sub> emissions are a major threat to the climate," Shu Ping Lau, co-author of the paper, told Tech Xplore. "We are looking to harness and re-enter massive amounts of CO<sub>2</sub> into the [carbon cycle](#) by using electrocatalytic CO<sub>2</sub> reduction (ECO<sub>2</sub>R) technology to combat this. However, previous research has shown that the stability of the ECO<sub>2</sub>R system is a major challenge, with current systems lasting less than 200 hours for ECO<sub>2</sub>R-to-ethylene (C<sub>2</sub>H<sub>4</sub>)."

In their recent work, Lau and his collaborators have been trying to overcome the limitations of existing systems for electrocatalytic CO<sub>2</sub> reduction. Their objective is to create a new electrolysis architecture that suppresses carbonate formation during ECO<sub>2</sub>R and thus enables a prolonged stable operation.

"Our goal is to maintain an alkaline cathode environment without involving alkali metal cations, ultimately designing the APMA MEA (AEM+PEM assembly membrane-electrode assembly) architecture with pure H<sub>2</sub>O as the anolyte," Lau explained. "In our APMA MEA ECO<sub>2</sub>R system, we've created a way for CO<sub>2</sub> to react with H<sub>2</sub>O to produce C<sub>2</sub>H<sub>4</sub> and OH<sup>-</sup> at the cathode, while H<sub>2</sub>O is oxidized into O<sub>2</sub> and H<sup>+</sup> at the anode. The resulting OH<sup>-</sup> and H<sup>+</sup> then combine to form H<sub>2</sub>O in the middle of the membranes."



System stability performance of  $ECO_2R$ -to- $C_2H_4$  in a pure- $H_2O$ -fed APMA-MEA cell stack containing 6 APMA-MEA cells at a constant current of 10 A. Inset: Schematic of the APMA-MEA cell stack containing 6 APMA-MEA cells for the  $ECO_2R$  reaction. Credit: She et al

The new system introduced by the researchers is comprised of two distinct membranes (AEM and PEM), a cathode catalyst (stepped-surface Cu), an anode catalyst (Pt/Ti) and pure water as the anolyte. One of its most remarkable advantages is that it does not require any additional chemicals to initiate reactions, and just uses pure  $H_2O$  as the electrolyte. This means that it could be easily scaled up to an industrial level.

"Even more impressive, the APMA MEA architecture overcomes the thermodynamic limitation of CO<sub>2</sub> reacting with the electrogenerated OH<sup>-</sup> into carbonate, which extends the system's stability," Lau said.

"With its durability and efficiency, our APMA MEA system has the potential to revolutionize CO<sub>2</sub> electrocatalysis technology and transform the modern fossil energy system."

In initial tests, the APMA MEA system introduced by this team of researchers achieved highly promising results. Using only pure H<sub>2</sub>O as the anolyte and under the forward-bias mode, it was found to effectively suppress carbonate formation during ECO<sub>2</sub>R, extending, the stability of CO<sub>2</sub> reduction to the hydrocarbon C<sub>2</sub>H<sub>4</sub> to reach an impressive 1,000 hours.

"Our breakthrough in creating a stable and durable ECO<sub>2</sub>R system is crucial to industrializing ECO<sub>2</sub>R," Lau said. "With the potential to move towards industrial-level rates, the APMA MEA system could pave the way for significant reductions in CO<sub>2</sub> emissions on an industrial scale."

The promising methods and technology introduced by Lau and his colleagues could soon be further improved and evaluated, both in laboratory and real-world industrial settings. Ultimately, it could contribute to ongoing global efforts aimed at reducing carbon emissions, by facilitating the electrocatalytic reduction of CO<sub>2</sub>.

"In our initial APMA MEA system, we encountered high operation voltage and low current density, resulting in a low yield of the desired product (C<sub>2</sub>H<sub>4</sub>) and overall low energy efficiency," Lau added. "Our next step is to concentrate on enhancing the [current density](#) and energy efficiency of the APMA system while reducing the system's overpotential."

**More information:** Xiaojie She et al, Pure-water-fed, electrocatalytic

CO<sub>2</sub> reduction to ethylene beyond 1,000 h stability at 10 A, *Nature Energy* (2024). [DOI: 10.1038/s41560-023-01415-4](https://doi.org/10.1038/s41560-023-01415-4)

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