Towards High Solar to Fuel Efficiency Wen-Hui Cheng, Matthias H. Richter, Matthias M. May, Jens Ohlmann, David Lackner, Ian Sullivan, David M. Larson, Chengxiang Xiang, Hans-Joachim Lewerenz, Frank Dimroth, Thomas Hannappel, Bruce S. Brunschwig, Harry A. Atwater



Abstract:

We demonstrated high solar-to-H₂ efficiency in PEC devices, consisting of a III-V based tandem light absorber and RuOx/Rh NP catalysts for OER and HER. Minimizing parasitic light absorption and reflection losses with favorable band alignment further reduces the efficiency gap to the theoretical limit. We also developed a solar-driven CO2 reduction device using a gas diffusion electrode (GDE) with Ag nanoparticle catalyst directly powered by a III-V based triple junction solar cell. Device geometry was studied to extend the operation stability.

Introduction

Results, Highlights, and Accomplishments

To realize high solar-to-fuel efficiency in PEC devices, it is necessary to maintain a catalytic current density the light limiting close to photocurrent density for a solardriven light absorber, which can be fulfilled when catalyst ensembles are highly transparent. Solar-driven reduction of carbon dioxide represents a carbon neutral pathway for the synthesis of fuels and chemicals. A gas diffusion electrode (GDE) directly powered by photovoltaic cell is a promising path to producing chemical fuels from CO2 and sunlight.

Monolithic Photoelectrochemical Device for Direct Water Splitting with 19% Efficiency

CO₂ Reduction to CO with 19% Efficiency in a **Solar-Driven Gas Diffusion Electrode Flow Cell** under Outdoor Solar Illumination

JCAP Team







Outlook

Solar-to-hydrogen efficiencies of 19.3% and 18.5% are obtained in acidic and neutral electrolytes. The system reaches a value of o.85 of the theoretical limit for photoelectrochemical water splitting for the energy gap combination employed in the tandem-junction photoelectrode structure.

A solar-to-CO energy conversion efficiency of 19.1% is also under simulated AM achieved 1.5G illumination at 1 Sun. The use of a reverse-assembled GDE prevented transition from a wetted to a flooded catalyst bed allowed the device to and operate stably for >150 h with no loss in efficiency.



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