

## New catalyst turns greenhouse gases into hydrogen gas

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Figure 1. Schematic comparison, SEM images, the correlation between the number of ALD cycles and the particle size/population, and x-ray photoelectron curves for the samples. (A) Conventional exsolution for LSTN and (B) corresponding SEM image of LSTN. Scale bar, 500 nm. (C) Topotactic exsolution via ALD for LSTN-20C-Fe and the corresponding SEM image of (D)



LSTN-20C-Fe after reduction. Scale bar, 500 nm. Credit: Ulsan National Institute of Science and Technology

A new nanocatalyst that recycles major greenhouse gasses, such as carbon dioxide (CO<sub>2</sub>) and methane (CH<sub>4</sub>), into highly value-added hydrogen (H<sub>2</sub>) gas has been developed. This catalyst is expected to greatly contribute to the development of various waste-to-energy conversion technologies, as it has more than twice the conversion efficiency from CH<sub>4</sub> to H<sub>2</sub>, compared to the conventional electrode catalysts.

A research team, led by Professor Gun-Tae Kim in the School of Energy and Chemical Engineering at UNIST has developed a novel method to enhance the performance and stability of catalysts, used in the reaction (i.e., dry reforming of methane, DRM) that produces  $H_2$  and <u>carbon</u> <u>monoxide</u> (CO) from well-known greenhouse gasses, such as  $CO_2$  and  $CH_4$ .

The conventional catalysts used for the dry reforming of methane (DRM) are nickel (Ni)-based metal complexes. Over time, however, the performance of catalysts degrade, so does the <u>catalyst</u> lifetime. This is because carbon accumulates on the surface of the catalysts, as the catalysts clump together or their reaction is repeated at a higher temperature.

"The uniform and quantitatively controlled layer of iron (Fe) via <u>atomic</u> <u>layer deposition</u> (ALD) facilitates the topotactic exsolution, increasing finely dispersed nanoparticles," says Sangwook Joo (Combined MS/Ph.D. in the School of Energy and Chemical Engineering, UNIST), the first author of the study.



The research team also confirmed that exsolution is promoted even with a very small amount of ALD-deposited Fe oxide (Fe<sub>2</sub>O<sub>3</sub>). "Notably, at 20 cycles of Fe oxide deposition via ALD, the particle population reaches over 400 particles (Ni-Fe alloys)," says Arim Seong of the School of Energy and Chemical Engineering, UNIST, the first co-author of the study. "As these particles are composed of Ni and Fe, they also exhibited high catalytic activity."



Figure 2. Catalytic properties for the DRM. (A) Reacted methane during the DRM reaction for LSTN, LSTN-10C-Fe, and LSTN-20C-Fe. (B) The activation energy of the methane reactivity calculated for LSTN, LSTN-10C-Fe, and LSTN-20C-Fe. (C) Time dependence of CH4 reactivity and H2/CO ratio for LSTN-20C-Fe in DRM at 700 degrees C. Credit: Ulsan National Institute of



Science and Technology

The new catalyst exhibited high catalytic activity for the DRM process with no observable degradation in performance for more than 410 hours of continuous operation. Their results also showed a high methane conversion (over 70%) at 700 degrees C. "This is more than twice the <u>power conversion efficiency</u> that of the conventional electrode catalysts," noted Professor Kim. "Overall, the abundant alloy nanocatalysts via ALD mark an important step forward in the evolution of exsolution and its application to the field of energy utilization."

## Provided by Ulsan National Institute of Science and Technology

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