

# Ni exsolved nanoparticles from a $La_{1-\alpha}Ni_{1-x}Cr_xO_3$ perovskite for Methane Steam Reforming

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## Significance and Relevance

Exsolution is promising in catalysis due to its ability to produce uniformly distributed, anchored metallic nanoparticles, enhancing resistance to coking and agglomeration. ABO<sub>3</sub> perovskites, such as La<sub>1-α</sub>Ni<sub>1-x</sub>Cr<sub>x</sub>O<sub>3</sub>, provide ideal structural stability and flexibility for this process. The reducibility of the B-site cation, like nickel, is vital for nanoparticle formation in applications like methane steam reforming. Additionally, A-site vacancies induce distortions that facilitate cation migration, partial substitution at the B-site is necessary to preserve the integrity of the perovskite structure. The La<sub>1-α</sub>Ni<sub>1-x</sub>Cr<sub>x</sub>O<sub>3</sub> design integrates these attributes, offering a highly efficient and stable catalyst for advanced catalytic applications.

Preferred and  $2^{nd}$  choice for the topic:  $H_2$  storage and transportation, green  $H_2$  production, hydrogen vectors

Preferred presentation: Oral only

## **Introduction and Motivations**

The Methane Steam Reforming (MSR) is the most commonly used process for hydrogen production from fossil fuels due to its high hydrogen-to-carbon ratio and low by-product formation compared to other methods<sup>1,2</sup>. Although this conventional reforming method is well-established, it comes with several drawbacks. One major issue is the deactivation of catalysts due to sintering or coke formation at high temperatures, which leads to a decline in catalytic activity<sup>3, 4</sup>.

This study focuses on utilizing exsolution from an ABO<sub>3</sub> perovskite as a method for synthesizing <u>well-dispersed metallic nanoparticles that are anchored to their matrix</u>. This approach <u>enhances stability</u> <u>and resistance to carbon deposition</u>. Specifically, the ongoing research evaluates the catalytic activity and stability in MSR using exsolved Ni nanoparticles from a La<sub>1-α</sub>Ni<sub>1-x</sub>Cr<sub>x</sub>O<sub>3</sub> perovskite. The investigation includes examining stoichiometric factors, A-site deficiencies, and partial substitution of the B-site.

#### **Results and Discussion**

The La<sub>1- $\alpha$ </sub>Ni<sub>1-x</sub>Cr<sub>x</sub>O<sub>3</sub> perovskites were prepared using the Pechini sol-gel method, as illustrated in Fig. 1A. The reactions conducted thus far (see Fig. 1B) have demonstrated enhanced performance during the second run for stoichiometric perovskites ( $\alpha = 0$ ), suggesting <u>exsolution and reconstruction</u> <u>processes driven by the feed gas</u>. Notably, the sample with no partial substitution (S1) achieved 100% conversion of CH<sub>4</sub> at approximately 600°C in the second run, showing higher conversion rates at lower temperatures compared to traditional catalysts. <u>Furthermore, the exsolved nanoparticles from A-site deficient perovskites displayed initial catalytic activity at lower temperatures than their stoichiometric counterparts.</u>

These findings, supported by characterization reports, imply successful production of exsolved Ni nanoparticles from A-site deficient perovskites. They also demonstrate higher crystallinity or an increased surface quantity of Ni<sup>0</sup> under exsolution conditions (H<sub>2</sub> atmosphere). Significantly, these conditions create active sites that facilitate reactions at lower temperatures than conventional industrial catalysts, highlighting their potential for further stability assessments.





**Fig. 1.** Schematic summary showing A. the synthesis route and experimental conditions for the preparation of stoichiometric (S) and A-site deficiency (D) perovskites and exsolved nanoparticles (E), and B. light-offs for methane steam reforming reaction.

## References

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