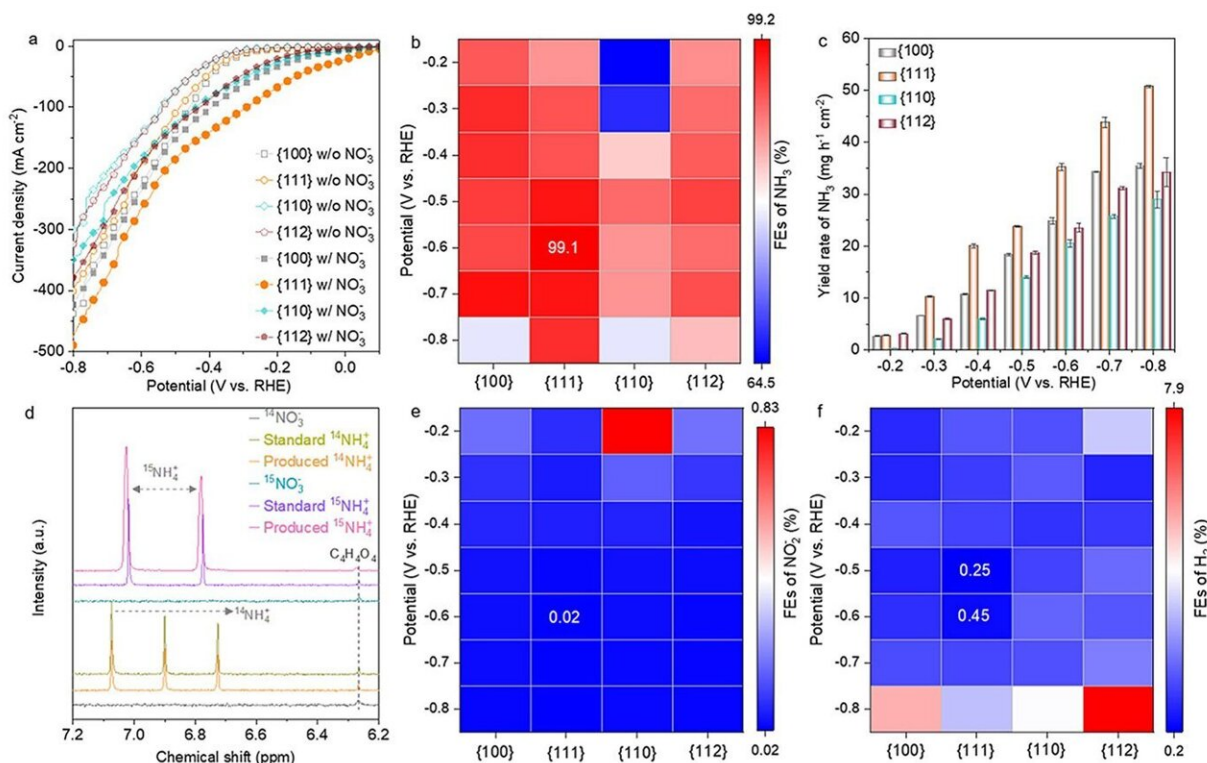


Insights into spinel cobalt oxides may lead to efficient ammonia synthesis

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eNO₃RR performance of Co₃O₄ {100}, {111}, {110}, and {112} catalysts. Credit: *ACS Nano* (2024). DOI: 10.1021/acsnano.4c06637

Researchers have made a significant breakthrough in the development of catalysts for the electrochemical nitrate reduction reaction (eNO₃RR) to ammonia, a process that has broad implications for sustainable energy,

agriculture, and industrial applications. The study is [published](#) in the journal *ACS Nano*.

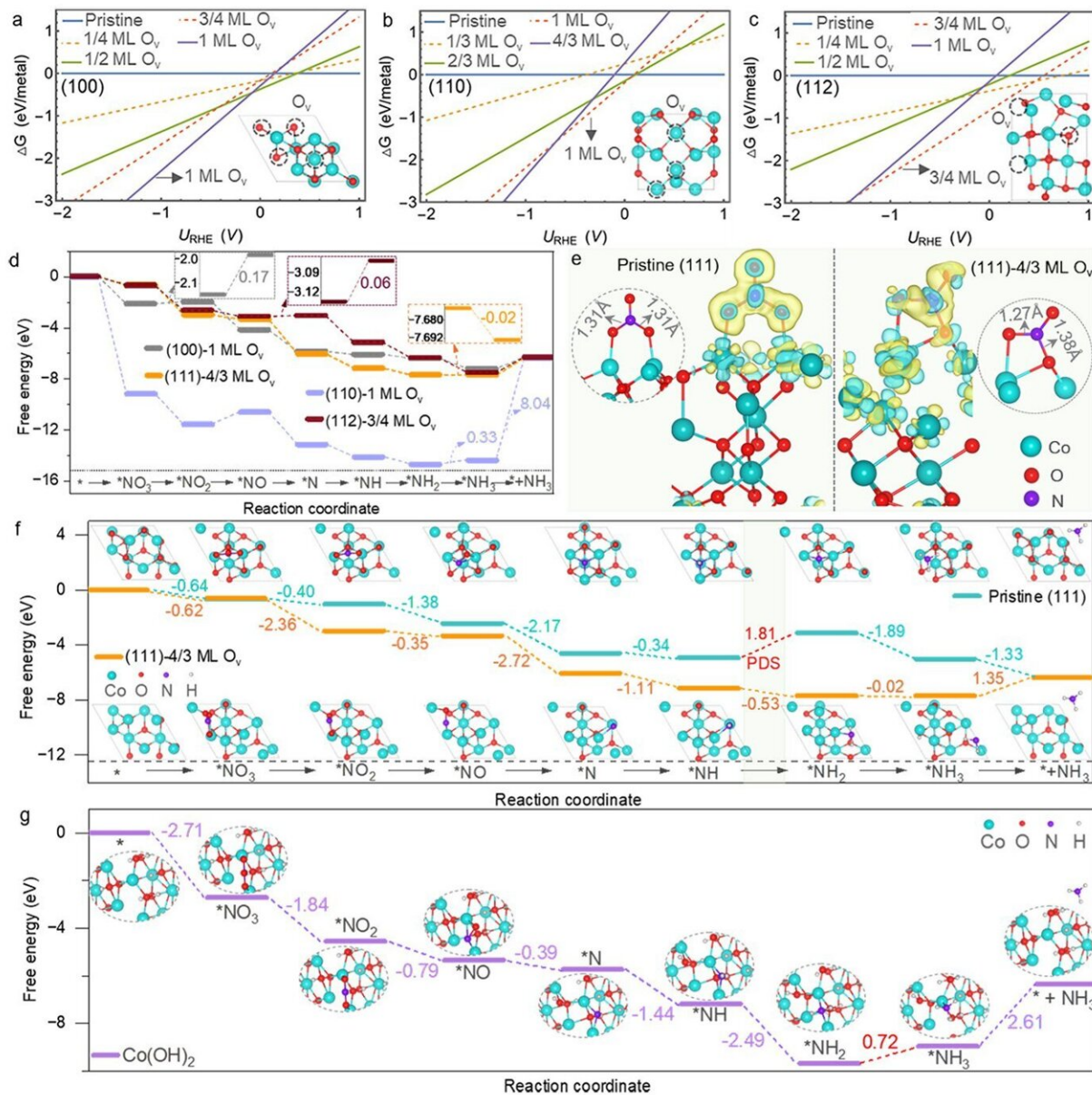
Ammonia, a critical component in global food production, also holds promise as a zero-carbon fuel due to its high energy density, clean combustion products, and established infrastructure for storage and transportation. However, the current method of producing [ammonia](#), the Haber-Bosch process, is energy-intensive and accounts for about 1.8% of global CO₂ emissions.

In their recent study, the research team focused on spinel cobalt oxides (Co₃O₄), a promising class of catalysts for eNO₃RR due to their low cost, high activity, and selectivity.

The team synthesized various Co₃O₄ nanostructures with different crystallographic facets—{100}, {111}, {110}, and {112}—to investigate how these facets influence the [catalyst](#)'s performance in ammonia production. The study revealed that the {111} facet of Co₃O₄ exhibited superior performance, achieving an impressive ammonia Faradaic efficiency of 99.1% and a yield rate of 35.2 mg h⁻¹ cm⁻².

"Our findings show that the {111} facet of Co₃O₄ is effective in transforming nitrate to ammonia," said Dr. Heng Liu, the co-first author of the paper and a Specially Appointed Assistant Professor at the Advanced Institute for Materials Research (WPI-AIMR), Tohoku University.

"This is due to the rapid formation of oxygen vacancies and Co(OH)₂ on this facet, which significantly enhances the catalyst's performance."

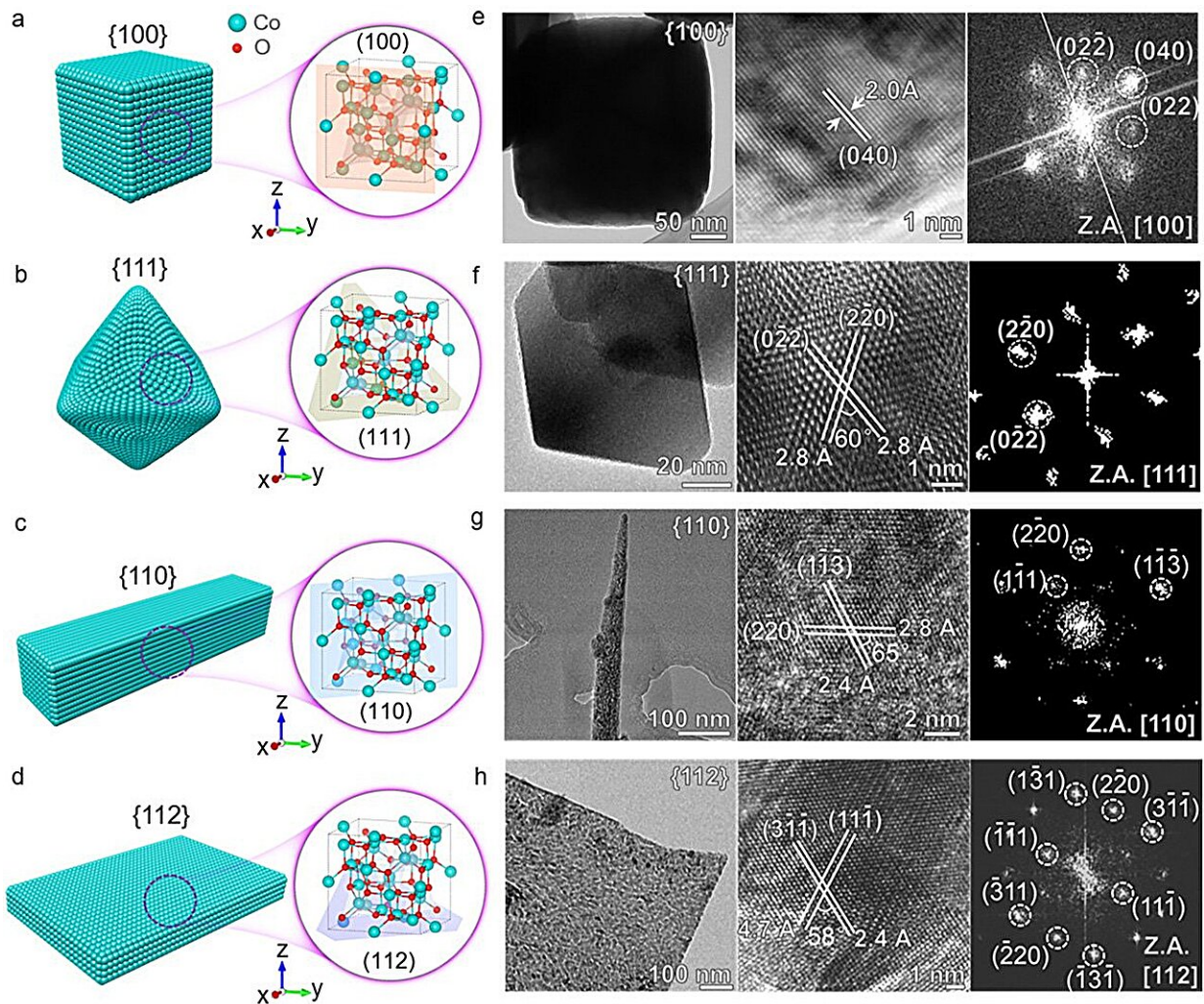


Theoretical calculations: 1D surface Pourbaix diagrams for Co₃O₄ (100), (110), and (112) surfaces, and reaction free energy diagrams for various intermediates on Co₃O₄ and Co(OH)₂ surfaces during eNO₃RR. Insets show charge density differences and N=O bond lengths. Credit: *ACS Nano* (2024). DOI: 10.1021/acsnano.4c06637

In addition, the researchers discovered that the catalyst went through a transformation process during the reaction, evolving from Co_3O_4 to a structure with oxygen vacancies, then to a $\text{Co}_3\text{O}_{4-x}\text{-Ov}/\text{Co}(\text{OH})_2$ hybrid, and finally stabilizing as $\text{Co}(\text{OH})_2$. This process was most pronounced on the {111} facet, contributing to its superior performance.

"The structural changes we observed are crucial for understanding the catalyst's activity," added Professor Hao Li, corresponding author of the paper and an associate professor at WPI-AIMR. "These insights will help us design more efficient catalysts by optimizing the exposed facets."

Ammonia's importance extends beyond agriculture, as it is a potential zero-carbon fuel and a key player in energy conversion and storage technologies. The eNO_3RR offers a sustainable alternative to the Haber-Bosch process, transforming nitrate waste into valuable ammonia while aiding environmental remediation.



Structural characterization of Co_3O_4 nanostructure with different facets. Credit: *ACS Nano* (2024). DOI: 10.1021/acsnano.4c06637

"This research lays a solid foundation for the development of more efficient, sustainable catalysts," states Li. "As we move forward, our goal is to control the final phases of the catalyst's transformation to further enhance its activity, selectivity, and stability."

This breakthrough in understanding and optimizing Co_3O_4 catalysts could pave the way for cleaner and more sustainable industrial processes,

contributing to the global efforts to achieve carbon neutrality by the 2050s.

More information: Anquan Zhu et al, Facet-Dependent Evolution of Active Components on Spinel Co₃O₄ for Electrochemical Ammonia Synthesis, *ACS Nano* (2024). [DOI: 10.1021/acsnano.4c06637](https://doi.org/10.1021/acsnano.4c06637)

Provided by Tohoku University

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