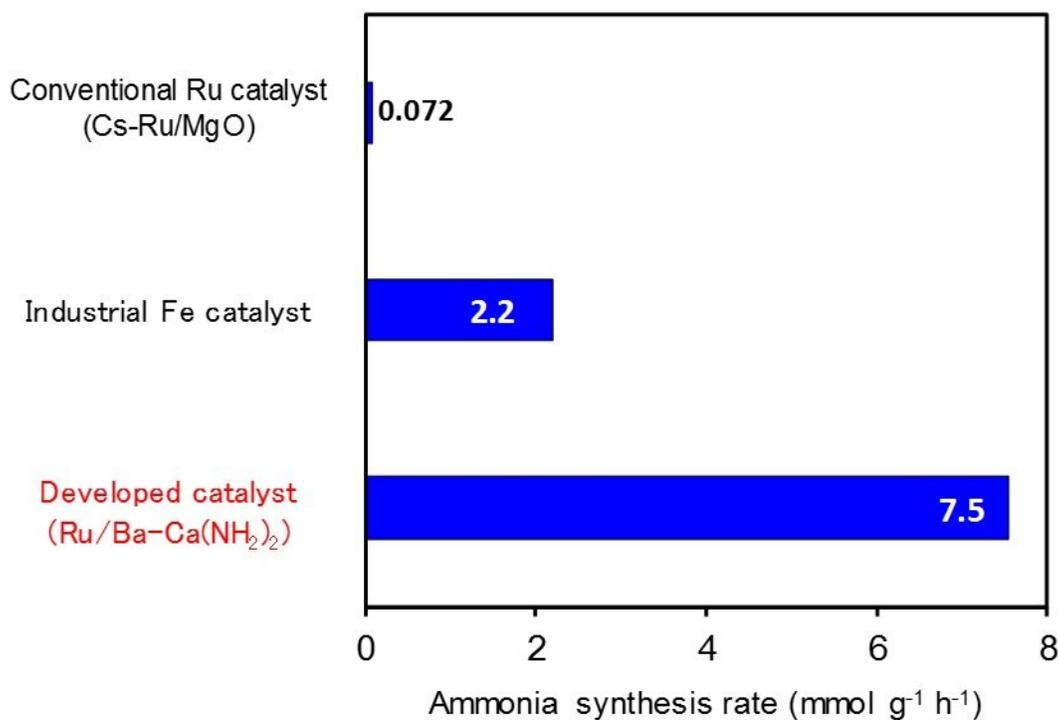


Highly efficient ammonia synthesis catalyst developed

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Comparison of ammonia synthesis activity (reaction temperature 260°C, pressure 9 atm). Credit: *Angewandte Chemie*

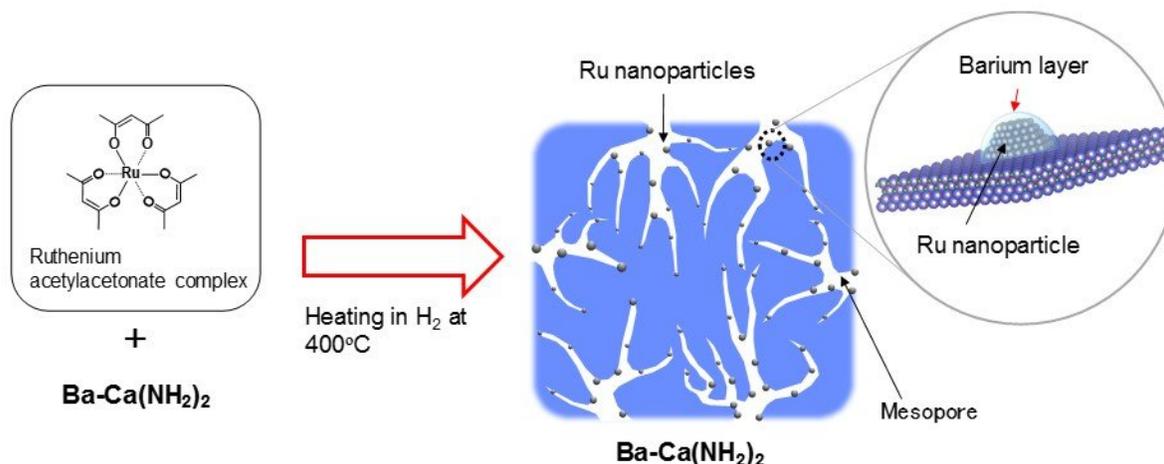
Researchers at Tokyo Institute of Technology (Tokyo Tech) have discovered that a catalyst of calcium amide with a small amount of added barium ($\text{Ba-Ca}(\text{NH}_2)_2$) with ruthenium nanoparticles immobilized onto it can synthesize ammonia at an efficiency 100 times greater than that of conventional ruthenium catalysts at low temperatures below 300°C . The performance of this catalyst is also several times higher when compared to iron catalysts currently being used industrially.

Ammonia is a raw ingredient for nitrogen fertilizers and is a key to food production. An [ammonia](#) molecule is one nitrogen atom bonded with three hydrogen atoms. As a result, ammonia is a substance with a very high hydrogen content for its mass. Because it becomes a liquid at room temperature at a pressure of 10 atmospheres, it is also an energy carrier for hydrogen, the energy source for technologies such as fuel cells.

The Haber-Bosch process, the current industrial [ammonia synthesis](#) method (established in 1913) uses a catalyst made mainly of iron and requires high temperatures (400 to 500°C) and high pressures (100 to 300 atm). To meet these conditions, ammonia is produced in large, dedicated factories and then transported to plants where it is used for industrial processes. There has long been demand for on-site production in which the required amount of ammonia can be synthesized where it is required, as opposed to the conventional large-scale process.

The Tokyo Tech research group of Professor Hideo Hosono, Professor Michikazu Hara, Associate Professor Masaaki Kitano and others discovered an ammonia synthesis catalyst that functions at high efficiency at low temperatures. They discovered that a catalyst of calcium amide with a small amount of added barium ($\text{Ba-Ca}(\text{NH}_2)_2$) with ruthenium nanoparticles immobilized onto it exhibits catalytic activity 100 times greater than that of conventional ruthenium catalysts at low temperatures below 300°C . Further, the catalytic performance of this catalyst is also several times higher when compared to iron catalysts

being used industrially (Figure 1).



The active structure of developed catalyst(Ru/Ba-ca(NH₂)₂) Credit: *Angewandte Chemie*

A ruthenium acetylacetonate complex is used as the raw material for ruthenium. By heating a powder mixed with Ba-Ca(NH₂)₂ to 400°C in a hydrogen atmosphere, a thin barium layer is formed on ruthenium nanoparticles about 3 nm in size, as porous calcium amide is formed concurrently (Figure 2). The surface area of Ba-Ca(NH₂)₂, the raw material for the catalyst, is only about 17 m²/g. However, since the

catalyst becomes porous when heated to 400°C in hydrogen with the ruthenium source, the group found that the surface area expands to approximately 100 m²/g. Further, the barium constituent added to the calcium amide moves to the surface of the catalyst during this heat processing and forms a thin layer by covering the [ruthenium](#) nanoparticles. The group discovered that this is a unique catalyst, with such active structures forming in a self-organized way and remaining stable throughout the reaction. The catalyst developed in this research exhibits the highest ammonia synthesis activity at [low temperatures](#) of any [solid catalyst](#) reported in recent years.

The [catalyst](#) developed in this research far supersedes the limits of existing catalytic materials in its ammonia synthesis activity and will contribute significantly to reducing energy used for the ammonia synthesis process. Because of this, further developing this technology is expected to lead to a new process structure for on-site synthesis of ammonia.

More information: Masaaki Kitano et al, Self-organized Ruthenium-Barium Core-Shell Nanoparticles on a Mesoporous Calcium Amide Matrix for Efficient Low-Temperature Ammonia Synthesis, *Angewandte Chemie International Edition* (2018). [DOI: 10.1002/anie.201712398](https://doi.org/10.1002/anie.201712398)

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