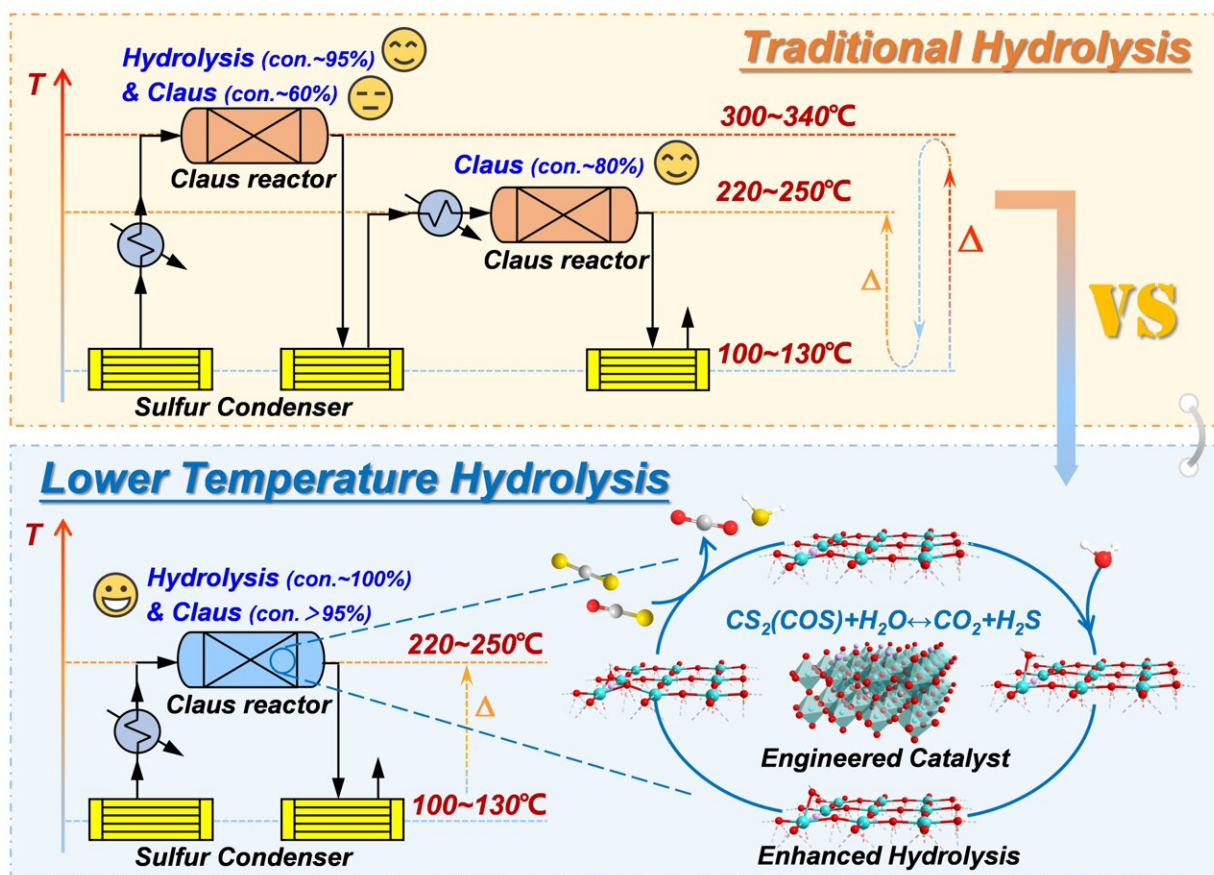


# Researchers propose titanium-based perovskite for water activation and lower-temperature hydrolysis of organic sulfur

February 14 2023, by Zhang Nannan



A schematic illustration for the comparison of traditional and novel operation process. Credit: UCAS

According to a study published in *Proceedings of the National Academy of Sciences* on Jan. 11, researchers led by Prof. Hao Zhengping from the University of Chinese Academy of Sciences (UCAS) have proposed facile oxygen vacancy ( $V_O$ ) engineering on titanium-based perovskite to motivate water ( $H_2O$ ) activation, achieving enhanced organic sulfur hydrolysis and efficient sulfur recovery at lower temperatures.

Water activation is involved in many reactions and processes. Organic [sulfur](#) ( $COS$  and  $CS_2$ ) hydrolysis is one such typical reaction using the  $H_2O$  molecule as a reactant. In industrial sulfur recovery processes, an additional hydrolysis catalyst is installed in the first stage of the Claus reactor to convert  $COS$  and  $CS_2$ . Limited by the effective activation of water, relatively higher temperatures are required for the hydrolysis of  $CS_2$  in a complex environment, which is not conducive to the sulfur recovery reaction, and thus severely affects the sulfur recovery efficiency and pollution emission control of the Claus process.

In this study, the researchers fabricated titanium-based perovskites with different  $V_O$  contents, and found that there was a linear correlation between the  $V_O$  content and the degree of  $H_2O$  dissociation and hydrolysis performance. The low-coordinated Ti ions adjacent to  $V_O$  were [active sites](#) for  $H_2O$  activation.

The introduction of  $V_O$ , especially  $V_O$  clusters, resulted in the reduction of the energy barrier for  $H_2O$  dissociation, which contributed to  $H_2O$  activation and dissociation, according to the researchers.

Furthermore, their in-depth mechanism study revealed that  $CS_2$  hydrolysis was initiated from the reaction between dissociatively adsorbed  $-OH$  and gaseous  $CS_2$  (Eley-Rideal mechanism), which was the origin of the enhanced hydrolysis activity from the enhanced  $H_2O$  activation by  $V_O$ .

Consequently, complete conversion of COS and CS<sub>2</sub> was achieved over SrTiO<sub>3</sub> after H<sub>2</sub> reduction treatment at 225°C, a favorable temperature for the Claus conversion. Surprisingly, the catalyst also showed excellent Claus catalytic activity.

Thus, both efficient organic sulfur [hydrolysis](#) performance and improved sulfur recovery efficiency can be achieved simultaneously. The application of this dual-functional catalyst can significantly improve the sulfur recovery efficiency, simplify the operation process and reduce the investment and operational cost.

Prof. Hao's team has long been engaged in the research and development of acid gas pollution control and resource recovery and utilization. They have an in-depth understanding of acid gas emission reduction and control. The production and publication of the results are based on the team's work over many years and are of great significance to the emission reduction control, resource recovery and utilization of acid gas.

**More information:** Zheng Wei et al, Oxygen vacancy-engineered titanium-based perovskite for boosting H<sub>2</sub>O activation and lower-temperature hydrolysis of organic sulfur, *Proceedings of the National Academy of Sciences* (2023). [DOI: 10.1073/pnas.2217148120](https://doi.org/10.1073/pnas.2217148120)

Provided by Chinese Academy of Sciences

Citation: Researchers propose titanium-based perovskite for water activation and lower-temperature hydrolysis of organic sulfur (2023, February 14) retrieved 27 April 2024 from <https://phys.org/news/2023-02-titanium-based-perovskite-lower-temperature-hydrolysis-sulfur.html>

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