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Notice for the PhD Viva Voce Examination

Mr Vinay Kumar M, Registration Number: 2270152, PhD Scholar at the Department of Physics and Electronics, School of Sciences, CHRIST (Deemed to be University), will defend his PhD thesis at the public viva-voce examination on Wednesday, 02 September 2026 at 02.00 pm in Room No. 044, Ground Floor, R&D Block, CHRIST (Deemed to be University), Bengaluru - 560029, Karnataka, India.

Title of the Thesis : **Integrating Different Strategies to Develop Efficient Photocatalyst for Effective Water Purification and Splitting**

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The members of the Research Advisory Committee of the Scholar, the faculty members of the Department and the School, interested experts and research scholars of all the branches of research are cordially invited to attend this open viva-voce examination.

Place: Bengaluru
Date: 27 August 2026

Registrar (Academics)

ABSTRACT

The dual challenges of pharmaceutical contamination in water bodies and the growing demand for clean energy could be addressed with the development of highly efficient, visible-light-active, and noble-metal-free photocatalysts. Graphitic carbon nitride (gC₃N₄) has attracted considerable attention due to its suitable band gap, chemical stability, and earth-abundant composition; however, its widespread application is limited by insufficient visible-light utilization, fast charge carrier recombination, and sluggish surface reaction kinetics. This study focuses on systematic advancement of gC₃N₄ through elemental doping/co-doping and co-catalysts loading, enabling the photocatalytic activity of gC₃N₄.

The first study shows a synergistic modification strategy by doping phosphorus into gC₃N₄ and loading cobalt boride (CoB) nanoparticles onto gC₃N₄ (CoB/P-gC₃N₄). Phosphorus incorporation effectively tailors the electronic structure by narrowing the bandgap and extending visible-light absorption, while uniformly distributed CoB nanoparticles create interfacial Schottky junctions that promote directional electron transfer and suppress recombination of charge carriers. The optimized CoB/P-gC₃N₄ photocatalyst shows a 35-fold enhancement in the tetracycline (TC) degradation reaction rate compared to pristine gC₃N₄ under visible-light irradiation, highlighting the effectiveness of P-doping and CoB-loading onto gC₃N₄.

To further enhance the TC degradation, in the second work, simultaneous doping of phosphorus and potassium along with CoB-loading was done to gC₃N₄ (CoB/P-K-gC₃N₄). The P-K co-doping induces bandgap narrowing, shallow impurity states, and pronounced charge redistribution, while the ultrasmall CoB nanoparticles not only form Schottky barriers but also generate localized surface plasmon resonance (LSPR), which improves the light absorption and charge separation. As a result, CoB/P-K-gC₃N₄ shows a 21-fold increase in TC degradation kinetics, achieving 84.8% degradation of 100 ppm TC within 60 min by using only 10 mg of catalyst, corresponding to a high removal efficiency of 7.1 mg g⁻¹ min⁻¹, along with excellent structural stability and reusability. Previously, CoB/P-gC₃N₄ activity was tested in a 20 ppm TC concentration. To compare CoB/P-gC₃N₄ with CoB/P-K-gC₃N₄, CoB/P-gC₃N₄ activity was also tested under 100 ppm TC concentration. Where the reaction kinetics of CoB/P-gC₃N₄ was just ~6.5 times higher than pristine gC₃N₄, clearly indicating the superior activity of CoB/P-K-gC₃N₄ compared to CoB/P-gC₃N₄ under high concentration of TC.

The third and fourth studies focus on sustainable hydrogen production. In the third work, we have introduced a phosphorus-incorporated cobalt boride (CoPB) co-catalyst onto gC₃N₄ (CoPB/gC₃N₄). The presence of electronegative phosphorus enhances proton (H⁺) adsorption, while the CoPB co-catalyst functions as an efficient electron acceptor due to a Schottky junction formation, thereby accelerating interfacial charge transfer and surface reaction kinetics. The CoPB/gC₃N₄ photocatalyst gave a hydrogen yield of 181.6 μ -mol g⁻¹ with a hydrogen evolution rate of 102 μ -mol g⁻¹ h⁻¹, giving a five-fold improvement compared to pristine gC₃N₄, and exhibits excellent photocatalytic reusability over repeated cycles.

In the final study, a bimetallic cobalt-molybdenum phospho-boride (CoMoPB) co-catalyst is loaded onto gC₃N₄ (CoMoPB/gC₃N₄) to further enhance the hydrogen evolution. The synergistic interaction between Co and Mo centers, combined with phosphorus-induced sites, increases charge carrier density and minimizes recombination losses. The optimized CoMoPB/gC₃N₄ system achieves a hydrogen production of 294.4 μ -mol g⁻¹ with an evolution rate of 146.8 μ -mol g⁻¹ h⁻¹. Overall, this work focuses on gC₃N₄-based photocatalysts for environmental remediation and hydrogen production.

Keywords: gC₃N₄, Doping, Co-Doping, Co-Catalyst Loading, Tetracycline Degradation, Photocatalytic Hydrogen evolution.

Publications:

1. Kumar, M. V., Fendrich, M., Orlandi, M., Miotello, A., Gupta, S., Patel, R., Fernandes, R., & Patel, N. (2025). Dual strategy for enhanced photocatalytic degradation of tetracycline: Phosphorus doping and cobalt boride co-catalyst loading on g-C₃N₄. *Journal of Water Process Engineering*, 70, 107036. <https://doi.org/10.1016/j.jwpe.2025.107036>
2. Kumar, M. V., Hamsaveena, D., Samartha, M. D., Bhagat, B. R., Orlandi, M., Miotello, A., Gupta, S., Patel, R., Fernandes, R., & Patel, N. (2026). Ultrasmall cobalt boride-decorated P, K-doped gC₃N₄ for plasmon-driven degradation of high concentration tetracycline. *Applied Surface Science*. <https://doi.org/10.1016/j.apsusc.2025.165683>