

Editorial

Editorial: Special Issue Entitled “Development of g-C₃N₄-Based Photocatalysts: Environmental Purification and Energy Conversion”

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Solar photocatalysis has evolved rapidly over the past few decades and has received significant attention for its green, safe and renewable energy benefits, particularly in the current era of global crisis, being considered as a potential solution to the major problems we face today, such as the shortage of fossil fuels and the impact of human activities on the environment [1,2]. The photocatalytic process is based on the unique ability of semiconductor catalysts to generate electrons and holes when exposed to light, and these photo-generated carriers are then used to interact with substances in the system, leading to the degradation of contaminants, water splitting into H₂ production, the reduction of CO₂ and the synthesis of high-additional-value chemicals [3]. Among the numerous catalysts for realizing photocatalysis, graphitic carbon nitride (g-C₃N₄) has attracted extensive attention due to its low cost, suitable band structure, and easy structural adjustment [4].

g-C₃N₄ is a carbon-based material that can be produced via the thermal polymerization reaction using precursors such as melamine, dicyandiamide, cyanamide, urea, thiourea, ammonium thiocyanate and other similar substances [5]. Although g-C₃N₄ possesses various advantages, its fast photo-generated carriers' recombination and narrow visible-light absorption region [6,7] limit its further practical application. There is therefore an urgent need, but still a significant challenge, to rationally design and develop g-C₃N₄-based photocatalysts through various modification strategies. Adjusting the morphology and structure of carbon nitride is one of the methods of modification. A review of reaction parameters, structure design and exfoliation methods focuses on the effect of various parameters of the synthesis process of g-C₃N₄ on the photocatalytic activity and on the methods of construction of microstructures [8]. Moreover, the synthesis of g-C₃N₄ was been described by Biswas et al. [9]. In addition, Shi et al. added phloxine B to the process of preparing g-C₃N₄ to produce black g-C₃N₄ nanosheets containing cyanine defects, which not only reduces the distance of charge transfer, but also enhances the photocatalytic activity through overlying the photothermal effect, providing beneficial ideas for the design of photocatalysts to degrade antibiotic pollutants [10].

Next, the construction of heterojunctions is also one of the means to enhance the photocatalytic activity of g-C₃N₄ [11,12]. When two semiconductors are in contact, the internal electric field (IEF) can be formed because of the difference between their Fermi levels, and the separation of electrons and holes is promoted under the action of IEF, effectively solving the problem of the fast recombination of photon-generated carriers [13]. Khan's team fabricated a dual Z-scheme heterojunction photocatalyst by anchoring P and S co-doped g-C₃N₄ on Ag/AgI/WO₃, which effectively reduced the recombination of photo-generated electrons and holes due to the enhancement of IEF, and effectively improved the photodegradation efficiency [14]. Li et al. prepared an isotype heterojunction of phosphorus-doped g-C₃N₄/phosphorus-sulfur co-doped g-C₃N₄ (P-g-C₃N₄/PS-g-C₃N₄)



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using a two-step calcination method, which not only suppressed electron–hole complexation through IEF to prolong the electron lifetime, but also lowered the interfacial resistance and accelerated the electron mobility [15].

In addition, loading co-catalysts on the surface of g-C₃N₄ is also a means of improvement. For example, Al-Hartomy's group using Pt as the co-catalyst to coat on the g-C₃N₄ framework for the efficient photocatalytic reduction of CO₂ to CO and CH₄ [16]. Peng et al. loaded g-C₃N₄ onto carbon dots modified with SBA-15 mesoporous silica by means of precursor impregnation, where the carbon dots act as co-catalysts with the ability to transfer photo-generated electrons, while the mesoporous silica has a larger specific surface area and porosity [17]. Furthermore, Kim et al.'s team loaded anthraquinone (AQ) onto g-C₃N₄, endowing the conduction band with the ability to trap electrons and was able to reduce the photocatalytic decomposition of produced H₂O₂, improving the selectivity of H₂O₂ production [18]. Not only that, Sun et al. also promoted the visible-light photocatalytic reduction of carbon dioxide by loading sodium hydroxide on the g-C₃N₄ surface. In the thin layer of alkaline electrolyte formed on the carbon nitride surface, it not only acts as a hole acceptor but also maintains the cycle of carbonic acid production and decomposition [19]. Li's group synthesized a g-C₃N₄/MoS₂ composite with a hydrogen production rate 13.44 times higher than that of the pure phase under visible light due to the fact that MoS₂ acted as a co-catalyst to significantly enhance the transfer of photo-generated electrons to H⁺ [20]. In conclusion, the modification of g-C₃N₄ materials is based on the following three principles: (i) structural adjustment of the g-C₃N₄ material itself to increase the specific surface area as well as the active sites; (ii) construction of g-C₃N₄-based heterojunctions to separate the photogenerated carriers by IEF; and (iii) combining with co-catalysts to make up for the problems of g-C₃N₄ itself.

To conclude, this Special Issue entitled "Development of g-C₃N₄-Based Photocatalysts: Environmental Purification and Energy Conversion" provides a comprehensive overview of recent advances in the synthesis, modification, and energy-environmental applications of g-C₃N₄, an excellent photocatalyst. We hope that this collection of important research papers will inspire further research in this field.

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