



2D/2D Graphitic Carbon Nitride (g-C₃N₄) Heterojunction Nanocomposites for Photocatalysis: Why Does Face-to-Face Interface Matter?

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In recent years, two-dimensional (2D) graphitic carbon nitride (g-C₃N₄) has elicited interdisciplinary research fascination among the scientific communities due to its attractive properties such as appropriate band structures, visible-light absorption, and high chemical and thermal stability. At present, research aiming at engineering 2D g-C₃N₄ photocatalysts at an atomic and molecular level in conquering the global energy demand and environmental pollution has been thriving. In this review, the cutting-edge research progress on the 2D/2D g-C₃N₄-based hybrid nanoarchitectures will be systematically highlighted with a specific emphasis on a multitude of photocatalytic applications, not only in waste degradation for pollution alleviation, but also in renewable energy production [e.g., water splitting and carbon dioxide (CO₂) reduction]. By reviewing the substantial developments on this hot research platform, it is envisioned that the review will shed light and pave a new prospect for constructing high photocatalytic performance of 2D/2D g-C₃N₄-based system, which could also be extended to other related energy fields, namely solar cells, supercapacitors, and electrocatalysis.

Keywords: graphitic carbon nitride (g-C₃N₄), photocatalysis, energy conversion, environmental remediation, 2D/2D heterojunction, face-to-face interface

INTRODUCTION

Photocatalysis is emerged as one of the Holy Grails of sustainable and green technologies for solar energy conversion, energy storage, and environmental remediation, which has been intensively examined over the past few decades worldwide to search for novel photocatalysts (Inoue et al., 1979; Linsebigler et al., 1995; Ma et al., 2014; Ong et al., 2014a, 2016b; He and Que, 2016; Li et al., 2016a; Wenderich and Mul, 2016; Zhang et al., 2016a; Eftekhari, 2017; Liu et al., 2017; Osterloh, 2017; Roger et al., 2017). By harvesting solar energy as the source of renewable energy, photocatalysis will make significant impacts in the areas of (1) light-driven water splitting to hydrogen (H₂) and oxygen (O₂) (Chen et al., 2010; Bai et al., 2016; Wei et al., 2016; Putri et al., 2017; Yubin et al., 2017), (2) conversion of carbon dioxide (CO₂) to energy bearing fuels (Ong et al., 2013, 2014c; Tan et al., 2014, 2016, 2017; Gui et al., 2015; Guo et al., 2016a; Zhang et al., 2016c), (3) mineralization of waste and pollutants (Ong et al., 2014d,e; Fang et al., 2016; Liu et al., 2016c; Topcu et al., 2016; Zhao et al., 2016b), (4) selective

organic transformations (Liu et al., 2014; Zhao et al., 2016a), and (5) disinfection of bacteria (Keane et al., 2014; Bing et al., 2015) (Figure 1). Very recently, two-dimensional (2D) semiconductor photocatalysts have triggered a renaissance of interest in the field of energy, and environmental-related applications thank to the high ratio of surface-to-volume and unprecedented electronic and optical characteristics (Ong et al., 2014b; Bai et al., 2015; Liang et al., 2015d; Fang et al., 2016; Kalantar-zadeh et al., 2016; She et al., 2017; Xueting et al., 2017). Among a large array of photocatalysts, research targeting at graphitic carbon nitride (g-C₃N₄) has been flourishing in recent years. Since the first exploratory study on the use of g-C₃N₄ in photocatalytic H₂ evolution in 2009 (Wang et al., 2009), there has been an exponential increase in the scientific research on the subject of g-C₃N₄-based materials with more than 800 publications in 2016 based on Web of Science.

By and large, g-C₃N₄ can be facily prepared by nitrogen-rich precursors, namely urea, thiourea, melamine, and dicyandiamide (Han et al., 2015; Guo et al., 2016b; Zhou et al., 2016; Tong et al., 2017). Therefore, the development of g-C₃N₄-based photocatalysts is anticipated to surmount the issues of increasing concerns on fossil fuel depletion and environmental threats due to combustion of exhaustible fossil fuels. The metal-free g-C₃N₄ demonstrates distinctive attributes such as visible-light responsiveness with moderate band gap of *ca.* 2.7 eV, appealing band structures and electronic characteristic, its earth-abundant nature, non-toxicity, relative ease of synthesis, and excellent chemical stability (Lu et al., 2016; Ong et al., 2016b; Zhang et al., 2016b; Lee et al., 2017). Additionally, it has been proven that 2D semiconductor possessed improved mobility of charge carriers and reduced charge recombination as compared to the 0D and 1D nanomaterials (Meng et al., 2012; Ida and Ishihara, 2014). In spite of the fascinating properties possessed by 2D g-C₃N₄, pristine g-C₃N₄ demonstrated several shortfalls such as

sluggish separation of electron-hole pairs, limited visible-light absorption beyond 460 nm, small specific surface area, and low electrical conductivity (Liang et al., 2015b; Hou et al., 2016; Shi et al., 2016; Zhang et al., 2016g; Li et al., 2017; Xia et al., 2017). To overcome these bottlenecks, modification of bare g-C₃N₄ such as nanostructure design (Niu et al., 2012; Liang et al., 2015c; Zheng et al., 2015), intercalation with Li⁺ and Cl⁻ (Liang et al., 2015a), elemental doping (Hu et al., 2015; Huang et al., 2015; She et al., 2016), copolymerization (Fan et al., 2016; Rahman et al., 2016), coupling with metals or noble metals (Tonda et al., 2014; Ong et al., 2015b), incorporation with other semiconductors (Ong et al., 2016a; Putri et al., 2016a; Zhang et al., 2016g; Ye et al., 2017), hybridization with metal phosphides (Pan et al., 2017; Wen et al., 2017; Yi et al., 2017; Zhao et al., 2017a,b), and many more has been widely investigated to enhance the photocatalytic efficiency for practical benefits. To date, there are a number of excellent review articles highlighting on g-C₃N₄-based photocatalysts ranging from materials synthesis, functionalization, and hybridization to diversified applications (Cao et al., 2015; Dong and Cheng, 2015; Yin et al., 2015; Zhang et al., 2015a; Zhao et al., 2015; Liu et al., 2016a; Mamba and Mishra, 2016). This undoubtedly connotes the significance of this research field hitherto in the scientific community.

Recently, the incorporation of 2D g-C₃N₄ photocatalyst with other 2D nanomaterials forming 2D/2D heterojunction hybrid nanocomposites has conceivably drawn increasing attention with practical importance (Hou et al., 2014; Xing et al., 2017). As a matter of fact, the layered heterojunction comprised dissimilar 2D nanomaterials is projected to give rise to positive impacts on charge transfer and separation as a result of built-in electric field at the atomically well-defined ultrathin interface (Hou et al., 2013b; Lu et al., 2016). Thus, the research on 2D/2D heterojunction with intimate face-to-face interface is of timely significance, which will elucidate us to deeply comprehend the photocatalytic reaction mechanism at the molecular level. Therefore, in this review, this leads to my immense interest to summarize the state-of-the-art research on 2D/2D g-C₃N₄ heterojunction nanohybrids to throw light on the future research horizon of g-C₃N₄ in artificial photosynthesis and environmental remediation.

DEVELOPMENT OF 2D/2D g-C₃N₄-BASED HETEROJUNCTION

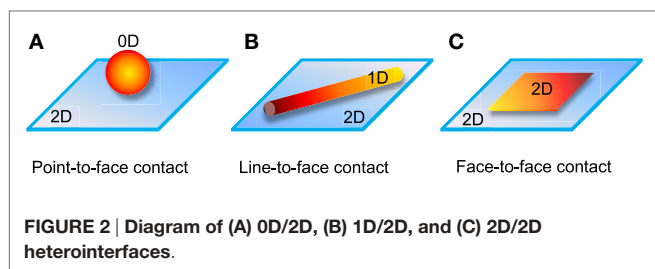
It is well documented that 2D/2D nanocomposites bestow greater electron-hole mobility across the heterojunction interface, which will in turn reduce the distance and time of charge transport to impede the electron-hole recombination rate (Hou et al., 2013a; Cheng et al., 2015; Ong et al., 2015c). This is attributed to the larger 2D/2D face-to-face contact area compared with line-to-face contact in 1D/2D heterojunction and point-to-face contact in 0D/2D heterojunction as depicted in Figure 2.

Hybridization with 2D Transition Metal Chalcogenides

In recent years, the incorporation of 2D metal sulfides has underpinned enormous interests in photocatalysis (Liu et al., 2016b; Lu



FIGURE 1 | Schematic of multifarious applications in photocatalysis research field.



et al., 2016; Yu and Sivula, 2016). In a work by Dong's research group, they reported a hierarchical sheet-on-sheet ZnIn₂S₄/g-C₃N₄ heterostructure by growing ultrathin ZnIn₂S₄ onto g-C₃N₄ nanosheets (Zhang et al., 2016e). As a result of intimate heterojunction interface formed between ZnIn₂S₄ and g-C₃N₄, the hybrid nanocomposites demonstrated whopping 17.6- and 3.9-folds enhancement of H₂ evolution compared to the single component g-C₃N₄ and ZnIn₂S₄, respectively. From the perspective of lifetime of charge carriers evidenced from time-resolved photoluminescence analysis, the average lifetime of the ZnIn₂S₄/g-C₃N₄ nanohybrids was reduced from 10.45 to 8.97 ns relative to that of pristine g-C₃N₄ nanosheets, which was attributed to rapid charge transfer and separation to hinder the electron-hole recombination. In addition to ZnIn₂S₄, MoS₂-decorated S-doped g-C₃N₄ heterojunction films were successfully developed by Chen and coworkers (Figure 3A) for enhanced photoelectrocatalysis (Ye et al., 2016). It is noted that the generation of anodic current by the MoS₂/S-doped g-C₃N₄ photoanode was markedly twice than that by the S-doped g-C₃N₄, highlighting the rational importance of a robust heterointerface with intact p-n junctions for effective charge migration (Figure 3B).

Till now, the theoretical understanding on the coupling interaction and transfer of charge carriers between 2D g-C₃N₄ and 2D MoS₂ has not been exhaustively investigated. Wang et al. (2014) elucidated the fundamental mechanism of photocatalytic improvements by systematically exploring the interface region between MoS₂ and g-C₃N₄. Based on the density functional theory (DFT) calculations, it was confirmed the presence of charge redistribution at the 2D/2D heterojunction interface of MoS₂/g-C₃N₄. It is worth mentioning that a type II heterojunction structure was successfully developed due to the well-matched band alignment as attested by the density of states results. As a result of efficient migration of charge carriers, a polarized field was formed at the contact heterointerface, prohibiting the electron-hole recombination. Therefore, this DFT finding provides new inroads into the importance of constructing 2D/2D nanocomposites for face-to-face interaction, which could certainly be extended to other binary or even ternary layered heterojunction for enhanced photochemistry applications.

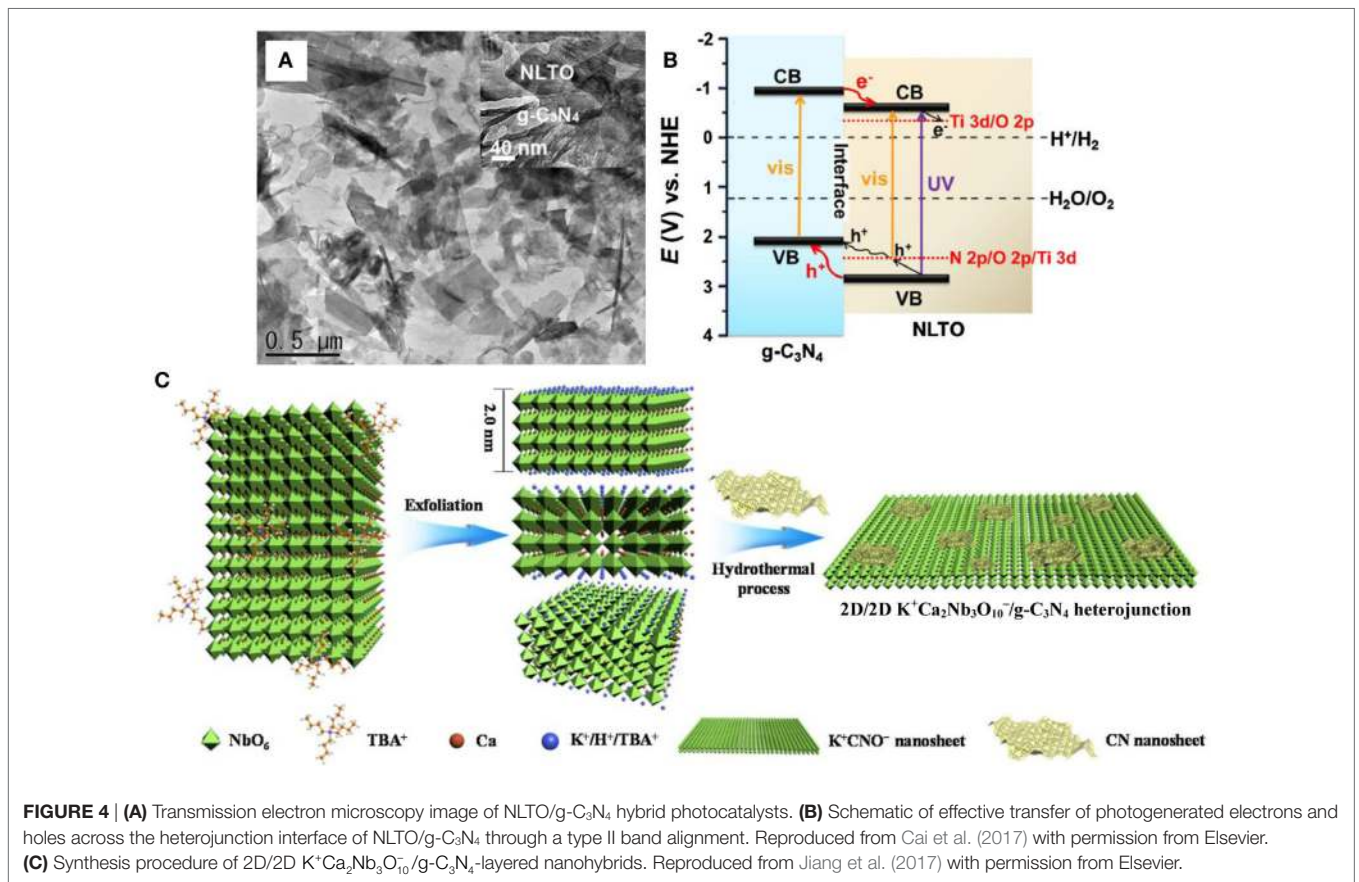
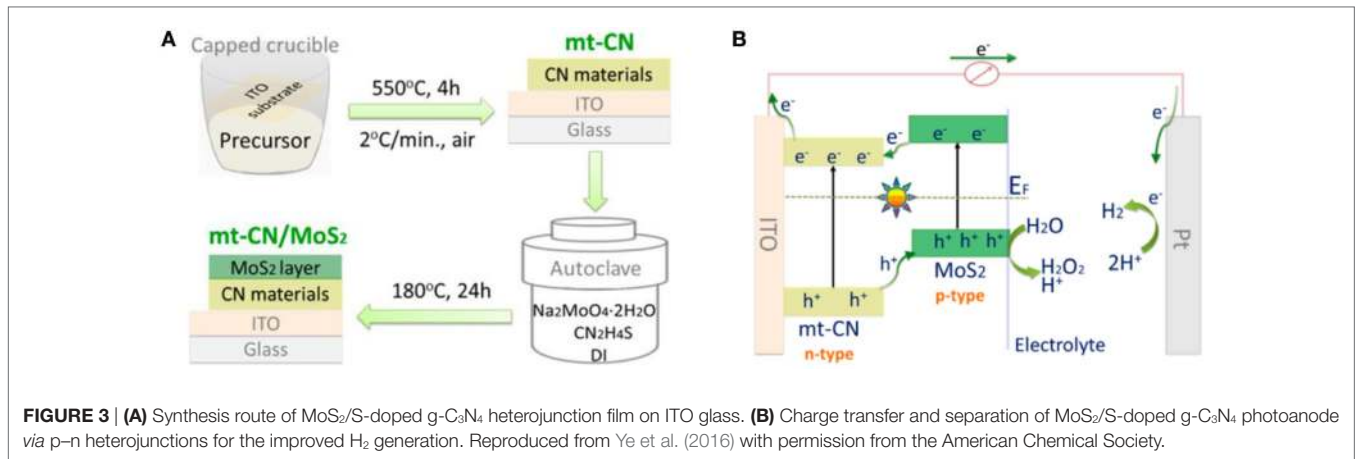
Hybridization with 2D Metal Oxides

Apart from 2D transition metal chalcogenides, coupling g-C₃N₄ nanosheets with 2D metal oxides, such as TiO₂ (Gu et al., 2014), WO₃ (Li et al., 2016b), and SnSb₂O₆ (Zhang et al., 2016d), has become the recent research focus to improve the photocatalytic performance. For instance, Li et al. (2016b)

employed a hydrothermal and deposition-heating technique to fabricate WO₃/g-C₃N₄ nanosheet arrays on the FTO substrates. The photoelectrochemical splitting of natural seawater using the WO₃/g-C₃N₄ nanostructures presented two times greater in the photocurrent density in reference to the pristine WO₃ nanosheet arrays under the simulated sunlight source. This was accredited to the well-matched band energy of WO₃ and g-C₃N₄ forming synergistic interfacial contacts for remarkably boosting the charge migration ability. In another work, King's group hybridized AgIO₃ with anisotropic g-C₃N₄ nanosheets to form a 2D/2D layered heterointerface toward increased photodegradation of Rhodamine B and methyl orange pollutants (Li et al., 2015). It is anticipated that these works will lay a pioneer groundwork and shed light for future directions in the interface engineering of 2D metal oxides and g-C₃N₄ for the advancement in solar energy conversion and environmental remediation toward practical applications.

Most recently, perovskite-type nitrogen-doped La₂Ti₂O₇ (NLTO), comprising a 2D architecture with a thickness of 7 nm, was for the first time hybridized with 2 nm thick g-C₃N₄ nanosheets by means of a facile two-step hydrothermal method and a thermal treatment process (Figure 4A) (Cai et al., 2017). The hybrid layered nanomaterials showed excellent photocatalytic H₂ evolution with a high apparent quantum efficiency of 2.1% at 400 nm. The enhanced photoactivity was ascribed to the successful development of the large 2D/2D interface between NLTO and g-C₃N₄, resulting in long lifetime and favorable transfer of charge carriers *via* type II band alignment. Upon visible-light illumination, the photoexcited electrons were facilely migrated from g-C₃N₄ to NLTO, whereas the photogenerated holes were transported from NLTO to g-C₃N₄, hampering the charge recombination process (Figure 4B). Notably, this was arisen from the band bending formed at the heterointerface, which induced a built-in electric field for the flow of charge carriers.

Furthermore, 2D niobium phase-layered perovskite Dion-Jacobson compounds have become a hot focal field in materials science and engineering for clean energy production and environmental cleaning in the past few years (Maeda et al., 2014; Oshima et al., 2016). It is well known that the layered perovskite Ca₂Nb₃O₁₀⁻ nanosheets, emerging from NbO₆ octahedra building blocks, are promising owing to their good chemical stability, high surface area, and relatively cost-effective (Sabio et al., 2010). Therefore, it is expected that by integrating the idea of Ca₂Nb₃O₁₀⁻ nanosheets into 2D g-C₃N₄ photocatalysts from the viewpoint of materials design, the photocatalytic efficiency of the composite nanomaterials will be exhilaratingly elevated. In a very recent work published in 2017, Chen's group designed a visible-light-responsive 2D/2D K⁺Ca₂Nb₃O₁₀⁻/g-C₃N₄ nanosheet heterojunction, which was fabricated by a hydrothermal process (Figure 4C), for photodegradation of tetracycline hydrochloride (Jiang et al., 2017). The hybrid photocatalyst manifested a strikingly high activity, which was 6.6 and 1.8 times greater compared with that of pure K⁺Ca₂Nb₃O₁₀⁻ and g-C₃N₄, respectively, due to the well-contacted heterointerfaces with strong interfacial coupling. Similarly, another group of researchers has synthesized the nanosheet composites by combining Ca₂Nb₂TaO₁₀ and g-C₃N₄ nanosheets through a simple solution exfoliation-reassembly



technique for solar H₂ production (Thaweesak et al., 2017). As such, it is believed that this research will cast new opportunities for engineering 2D/2D perovskite-based nanosheets coupled with g-C₃N₄ heterojunction interface for multitudinous light-driven applications.

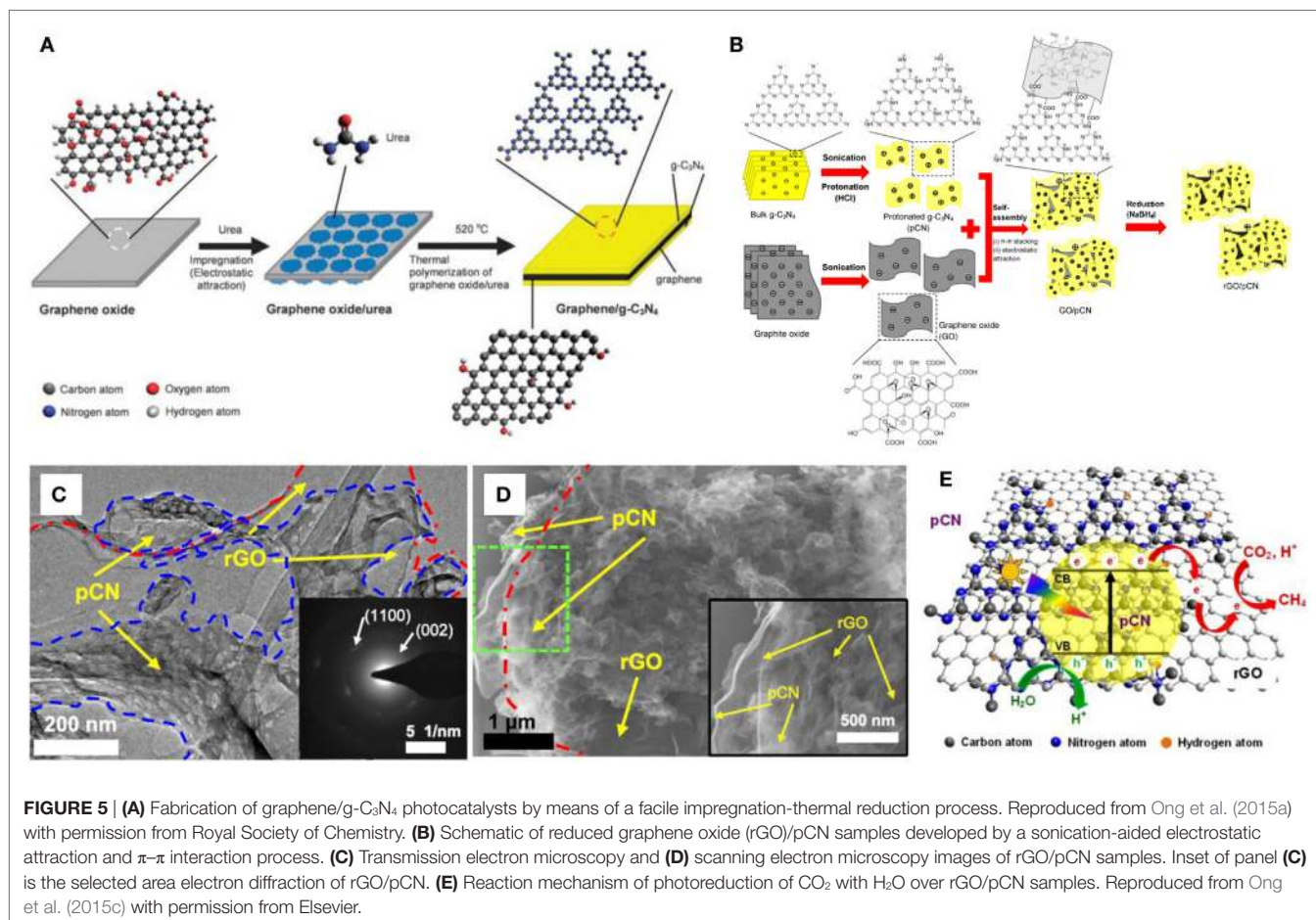
Hybridization with 2D Graphene

At present, π -conjugated carbonaceous nanomaterials, including carbon nanotubes, grapheme, and carbon nanodots has emerged as one of the most fascinating and exciting research

directions in the past 10 years especially in the arena of environmental remediation, energy conversion, and energy storage (Tan et al., 2013; Cazorla-Amorós, 2014; Li et al., 2014; Cao and Wei, 2015; Fan, 2015; Ali Tahir et al., 2016; Carmona et al., 2016; Kotal et al., 2016; Xu et al., 2016). These carbon nanostructures have been commonly utilized as excellent reduction cocatalysts by incorporating with semiconductor photocatalysts to prolong the lifetime of charge carriers to diminish the recombination of electron-hole pairs (Xu et al., 2013; Himaja et al., 2015; Liu et al., 2015; Tan et al., 2015a; Hu, 2016; Zhang et al., 2016; Ong

et al., 2017). It is discernable that the 2D graphene has directed a worldwide trend in the materials research stemming from its large surface area, remarkable electronic, optical and mechanical features, and high chemical stability (Putri et al., 2015, 2016b; Voon et al., 2016; Xiang et al., 2016). Up to now, a plethora of literature reports has been devoted to fabricate 2D/2D graphene/g-C₃N₄ nanohybrids for photoredox catalysis in H₂ evolution, pollutant degradation, and CO₂ reduction (Xiang et al., 2011; Li et al., 2013; Xu et al., 2015; Wan et al., 2016). In a work by Ong et al. (2015a), sandwich-like graphene/g-C₃N₄ nanocomposites were prepared *via* a one-step impregnation-thermal reduction process by employing graphene oxide and urea as the precursors (**Figure 5A**). Interestingly, the absorption band edge of the nanohybrids was slightly red shifted toward a longer wavelength, resulting in a reduction in the band gap energy. This phenomenon was contributed by the covalent cross linker (C–O–C) formed between g-C₃N₄ and graphene as a result of thermal heating at the high temperature. For the first time, the metal-free graphene/g-C₃N₄ photocatalyst played a prominent role in the reduction of CO₂ to CH₄ under visible light, which was 2.3 times higher than pristine g-C₃N₄. In this sense, this study incontrovertibly focuses the spotlight on the innovative design of metal-free layered photocatalysts as a new class of light-active materials for a cornucopia of catalytic applications.

In another closely related work by the similar research group, the novel 2D/2D-reduced graphene oxide (rGO)-hybridized protonated g-C₃N₄ (rGO/pCN) was rationally constructed by π - π stacking and electrostatic self-assembly between the positively charged pCN and the negatively charged rGO (**Figure 5B**) (Ong et al., 2015c). In the rGO/pCN hybrid nanoarchitectures, a well-dispersed sheet-on-sheet structure of rGO and pCN confirmed the well-intact interfacial contact (**Figures 5C,D**) as compared to the rGO/g-C₃N₄, which employed the unmodified g-C₃N₄ with a negatively charged surface. Essentially, the rGO/pCN heterojunction nanosheets endowed pronounced 5.4 and 1.7 times enhancement in the photoconversion of CO₂ to CH₄ with respect to the pCN and rGO/g-C₃N₄, respectively. Thus, this distinctly underlines the fundamental and technological importance of surface charge modification between two dissimilar 2D nanomaterials for robust interfacial interactions in unraveling the charge dynamics for enhanced photocatalysis. Benefiting from the predominant role of graphene as the electron reservoir (Zhang et al., 2012, 2015b; Tan et al., 2015b; Mateo et al., 2016; Varadwaj and Nyamori, 2016), the photoinduced electrons were transferred from pCN to rGO across the interface to overwhelmingly suppress the charge recombination rate (**Figure 5E**). All in all, the smart 2D/2D interface engineering design of graphene/g-C₃N₄ developed thus far is considered as an auspicious means, which



could be eminently extended to heteroatom-doping graphene-hybridized g-C₃N₄ for targeting superior photochemistry applications for real-life applications.

CONCLUSION AND OUTLOOK

In short, the burgeoning developments of nanoscale architecturing of g-C₃N₄-based hybrid structures over the past 8 years have witnessed a wealth of knowledge and information for the intelligent design and myriads of applications in sustainable energy conversion and environmental purification. The applications, which encompass water splitting, H₂ generation, O₂ evolution, CO₂ fixation, and pollutant degradation, have readily made full use of the intriguing features of g-C₃N₄, namely metal-free 2D nanomaterials, earth-abundant nature of the elements, visible-light optical absorption, high redox power, and excellent chemical stability. Since the advent of g-C₃N₄ photocatalysts by Wang et al. (2009) for H₂ generation, there is a rocket rise of research works on the modification of bare g-C₃N₄ to conspicuously increase the specific surface area, introduce porosity by textural modifications, extend visible-light absorption to longer wavelengths (even up to near infra-red) for the utilization of whole solar spectrum, decrease the band gap energy and bolster the charge migration and separation.

In this mini-review, a systematic discussion on the most updated advancements of engineering 2D/2D g-C₃N₄ heterojunction layered nanoarchitectures with boosted photoactivity has been reviewed. It is worth mentioning that albeit there is a large library of recent research discoveries on the 2D/2D g-C₃N₄-based photocatalysts, there exist numerous open issues, limitations, questions, and complexities of materials science, chemistry, physics, and environmental science, which require extensive research now and future. Among all, the actual mechanisms of enhanced catalytic efficiency toward the water splitting and CO₂ reduction followed by their respective reaction pathways are still up in the air yet until now. It is envisaged that the pertinent mechanism underlying the photocatalytic performance should be deeply explored by joining the experimental findings and theoretical computational simulations for the future research. In this manner, the rationale behind the profound photocatalytic enhancement especially on the rate determining steps of the reaction in the 2D/2D nanohybrids can be entirely comprehended. Apart from that, the charge carrier dynamics and transfer pathway for the Z-scheme system, p-n heterojunction, n-n heterojunction, Schottky junction, homojunction, and facet junction in the 2D/2D g-C₃N₄-based system will be facily understood. Benefiting from both experimental results and first-principles DFT calculations, this will in turn provide us a rational outline to advance the state

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of the research on photocatalysis for the next breakthrough in the field of energy conversion.

Moreover, the coupling interaction between 2D g-C₃N₄ nanosheet and another 2D semiconductor is of utmost importance for developing intact heterojunction interfaces for efficient electron-hole shuttling to prolong the lifetime of charge carriers to accelerate the photocatalytic efficiency. In-depth studies in engineering, the intimate heterointerfaces of the 2D/2D nanohybrids at a molecular level will give rise to captivating results for tuning the existing molecular structure of bare g-C₃N₄, thereby enhancing mobility of electron-hole pairs and subsequently improving the photocatalytic redox ability. Additionally, it is crucial to attain a facile and low-cost metal-free 2D/2D g-C₃N₄-based photocatalyst system without comprising metal-containing semiconductors for practical benefits. Thus, continuous efforts in exploring non-metal semiconductors to couple with g-C₃N₄ will be advantageous for industrialization and commercialization in the long run to combat the cost concern for the large-scale processes.

To cut a long story short, it is apparent that the research progress has been tremendously impressive at this juncture by viewing at the relatively short period of time and history of g-C₃N₄-based photocatalysis. Undeniably, the incessant research efforts on the 2D/2D-layered nanocomposites will open new vistas and lay a strong foundation for advanced light-driven catalysis and electrocatalysis, which undeniably warrant continuous research along this direction. Without any doubts, this will act as a new paradigm for the next generation smart artificial photocatalytic systems for practical and commercial benefits in order to bridge the gap between lab-scale research and large-scale industrial applications. All in all, with the ceaseless cooperative work from all segments and disciplines in the world, the targets of building a cleaner, greener, sustainable, and zero-energy environment will be systematically accomplished in years to come. Last but not least, it is genuinely hoped that this mini-review will paint a much clearer image to direct us for the upcoming research horizons in 2D/2D photocatalysis for momentous breakthroughs in attaining highly effective, efficient, and economical g-C₃N₄-based system in future.

AUTHOR CONTRIBUTIONS

W-JO carefully outlined the contents of the review and wrote the entire manuscript.

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Conflict of Interest Statement: The author declares that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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