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Co-catalyst design to control charge transfer and product composition for photocatalytic H₂ production and biomass reforming†

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The exploration of co-catalyst effects in photocatalytic H₂ production along with biomass reforming is still limited. This study employs S- and N-doped graphene oxide dots (SNGODs) along with Pt and Ag co-catalysts to investigate the photocatalytic reforming of xylose (C₅) into its derivatives and gaseous H₂. Our study shows that Pt-SNGODs effectively catalyze both the reduction and oxidation steps: water reduction for H₂ evolution and oxidative reforming of xylose into C₅-C₁ species through successive alternation of hydrolysis and photocatalytic oxidation. Ag-SNGODs are less effective in the H₂ evolution reaction and the accumulated photogenerated electrons facilitate cleavage of xylose, through the retro-aldol reaction, into C₃- and C₂-species, which are then reformed into C₃-C₁ species. In addition to xylose cleavage, Ag-SNGODs are effective in reducing xylose and formate into radicals, which would proceed with C-C coupling to produce a C₆ compound. The present study demonstrates how the co-catalyst of photocatalysts can influence the charge transfer dynamics and the product composition in photocatalytic biomass reforming.

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Introduction

The environmental issues caused by fossil fuel consumption across the world have urged people to find a new alternative energy source. Hydrogen with a high heating value has emerged as a promising environmentally-benign energy resource.¹ Biomass oxidation through photo-reforming, using sunlight as the energy input to produce hydrogen, offers a sustainable green energy solution.^{1,2} To improve the photocatalytic reduction process for hydrogen generation, a co-catalyst is often used.³ In addition to affecting the hydrogen evolution, the co-catalyst can also influence the organic products derived from biomass oxidation.⁴ The mechanism by which the co-catalyst affects the photo-reforming products and reaction routes is yet to be fully understood.

There has been a growing interest in the direct photocatalytic reforming of biomass into hydrogen and valuable chemicals. The photocatalytic reforming of xylose (C₅, C₅H₁₀O₅) and glucose (C₆) is particularly intriguing due to the abundance of these sugars in plant materials.⁵⁻⁸ Most research efforts have focused on glucose and only a limited number of studies reports

on the photocatalytic reforming of xylose using semiconductors,⁹⁻¹¹ overlooking the potential of xylose. This is notable, considering that hemicellulose, a polysaccharide made up of xylose, constitutes the second most abundant component in lignocellulosic biomass (5–35%).¹² Furthermore, xylose production technologies are advancing each year, making it crucial to explore applications for xylose,¹³ such as its use in biomass reforming. Research on the photocatalytic reforming of xylose into clean fuels and valuable chemicals is essential.

Many metal-oxide catalysts have been used for biomass photo-reforming, especially TiO₂,² which is the most developed photocatalyst due to its advantages of good chemical stability, non-toxicity, and a low cost.³ However, its performance is limited by its ability to only absorb the UV light, which dissipates the abundance of solar light.^{2,14} CdS photocatalysts that are sensitive to visible light have also been used, but their high charge recombination and toxic cadmium content make them less feasible for the photocatalytic reaction.^{2,15} Due to the easy method for synthesis and a great physicochemical stability, graphitic carbon nitride (g-C₃N₄) is widely used for photocatalytic processes.^{16,17} However, its drawback in visible light absorption and charge recombination needs to be tackled.¹⁸ To overcome these shortcomings, graphene-based materials were developed and have been emerged as a promising photocatalyst because of their excellent photo-conductivity, tunable electronic-structure, environmental friendliness, cheapness, and light-absorbing properties.^{19,20}

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Notably, graphene oxide dots (GODs) exhibit exceptional property-tunability owing to their quantum size effect and chemical-modulation readiness.²¹ With sizes typically below 10 nm, GODs showcase unique size-dependent electronic and optical behaviors, enabling precise control over their bandgap and versatile applications.²² The high surface area of GODs provides abundant active sites for catalytic reactions, making them highly efficient in photocatalysis.²³ Their excellent photoluminescence, biocompatibility, chemical stability, and low toxicity further enhance their wide appeal, particularly in biomedical and environmental applications.²⁴ Regarding chemical modification of GODs, the introduction of sulfur and nitrogen has been reported to narrow the bandgap, repair vacancy defects, and facilitate charge separation, thereby resulting in enhanced harvesting of solar light, promoted delocalization of photogenerated charges, and improved charge reactivity on the catalyst surface.^{8,25–27}

The challenge of low photocatalytic activity often arises due to the rapid recombination of photogenerated carriers and the inefficient use of surface charges within the photocatalyst.³ To address this issue, incorporating a co-catalyst with the photocatalyst has proven effective. The co-catalyst can enhance the separation of photogenerated electron–hole pairs and serve as an active site, facilitating crucial interfacial reactions.²⁸ Previous studies have highlighted the effectiveness of co-catalyst nanoparticles like Pt, greatly improving photocatalytic H₂ production.²⁹ Ag was seldom used as a co-catalyst for photocatalytic H₂ production due to the sluggish interfacial H₂ evolution reaction.^{29,30} Nevertheless, Ag enhanced the photocatalytic decomposition of various organic species.³¹ Regardless of the H₂ evolution efficiency, the potential impact of co-catalysts on the photocatalytic biomass reforming into valuable chemicals demands in-depth studies. Therefore, investigating the different effects of Ag and Pt as co-catalysts for the current system is expected to shed light on the transfer patterns of photogenerated charges and the correlation between reductive H₂ evolution and oxidative biomass reforming.

In this study, we used sulfur- and nitrogen-doped graphene oxide dots (SNGODs) as the model photocatalyst to examine the influence of the co-catalysts, Pt and Ag, on the photocatalytic reforming of xylose. Under simulated solar irradiation, the co-catalyst-deposited SNGODs, Pt- and Ag-SNGODs, reformed xylose into its biomass derivatives with simultaneous H₂ production. Co-catalysts Pt and Ag resulted in different biomass reforming pathways, as a consequence of different charge transfer behaviors between the catalyst and co-catalyst. Detailed reaction mechanisms of xylose conversion into its derivatives were elucidated in this study. Our results indicated that Pt-SNGODs were active in H₂ production from the photocatalytic reduction whereas the oxidative xylose reforming mainly produced formate, which is an important commodity in the chemical, agricultural, textile, leather, medicinal, and rubber sectors.^{32,33} When Ag-SNGODs were used, the xylose reforming system mainly produced gluconic acid and acetic acid, which were important chemicals in food, cleaning, medicine, and agriculture industries.^{34,35} The present work provides

a conceptual interpretation on the product selectivity in photocatalytic biomass reforming based on the selection of the co-catalyst.

Result and discussion

Characterization of the catalyst

The SNGODs were synthesized by annealing graphene oxide (GO) sheets in sulfur and NH₃ at 600 °C, oxidizing the annealed product into dots in HNO₃ solution, and finally treating the dots in NH₄OH solution at 140 °C.²⁰ SNGODs and GO sheets were mixed with H₂PtCl₆·6H₂O or AgNO₃ in an aqueous xylose solution to proceed with photo-deposition of Pt or Ag under simulated 1-sun solar irradiation, to produce Pt-SNGODs or Ag-SNGODs, respectively. The GO sheets were used to collect the photogenerated electrons from SNGODs for the deposition of Pt or Ag, which served as H₂ generation sites.³⁶ Fig. S1a† presents the transmission electron microscopy (TEM) images of SNGODs and the inset histogram illustrates that SNGODs exhibited a particle distribution around 2–5 nm, with a predominant size of 3–4 nm. A lattice *d*-spacing of 0.213 nm shown in the high-resolution image (Fig. S1b†) represents the graphene (1100) planes.³⁷ The TEM images of Pt-SNGODs and Ag-SNGODs along with the GO sheets are shown in Fig. S2 and S3,† respectively. Fig. S2a and S3a† present that both the SNGOD (with a gray color) and co-catalyst (with a darker color) particles were distributed over the GO sheets. Lattice *d*-spacing of 0.232 and 0.236 nm in the high resolution images of Fig. S2b and S3b† represent the Pt (111) and Ag (111) planes, respectively, indicating successful deposition of the co-catalysts. These TEM images also indicate the high crystallinity of the SNGOD particles, which benefits light absorption and prevents charge recombination, thereby augmenting the photocatalytic activity.²¹

The full-range X-ray photoelectron spectroscopy (XPS) spectrum of SNGODs was depicted in Fig. S4a,† which includes S_{2p} (168 eV), S_{2s} (233 eV), C_{1s} (285 eV), N_{1s} (401 eV), and O_{1s} (532 eV) peaks. Sulfur and nitrogen atoms were successfully incorporated into the graphene oxide structure as a result of the treatment. In Table S1,† the atomic ratios of O_{1s}/C_{1s}, N_{1s}/C_{1s}, and S_{2p}/C_{1s} were determined, by the XPS analysis, to be 40%, 11%, and 7%, respectively. Fig. S4b–d† were the focused regimes of Fig. S4a† to clearly show the peaks of C_{1s}, S_{2p}, and N_{1s}, respectively. Using a Gaussian function, the C_{1s}, N_{1s}, and S_{2p} spectra were deconvoluted into their constituting peaks. The C_{1s} peak in Fig. S4b† contained C–C (284.6 eV), C–N (285.9 eV), C–O (286.5 eV), C=O (288 eV), and O–C=O (288.7 eV) bonds.³⁸ A nitrogen content due to the nitrogen-doping modification increased the catalyst activity.³⁹ The S_{2p} peak (Fig. S4c†) contained –SH (162.3 eV), C–S–C (163.9 eV), C=S (165.1 eV), C–SO₂ (168.3 eV), and C–SO₃ (170.5 eV) with C–SO₂ group give the highest composition (5.7% out of 7%) among the sulfur functionalities.^{26,27} The sulfur-functionalities at the graphene periphery were highly polar, attractive to photogenerated holes, and functioned as oxidizing sites during photocatalysis.²⁶ The N_{1s} peak (Fig. S4d†) was deconvoluted into pyridinic (398.4 eV), amino (399.2 eV), pyrrolic (399.6 eV), quaternary (400.7 eV), and



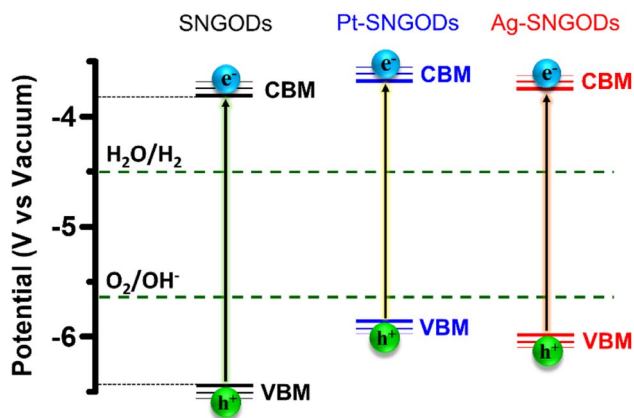


Fig. 2 Electronic-energy level diagrams of the SNGODs, Pt-SNGODs, and Ag-SNGODs in relation to the levels for H_2 and O_2 evolution reactions (HER and OER) in an aqueous solution of pH = 9.

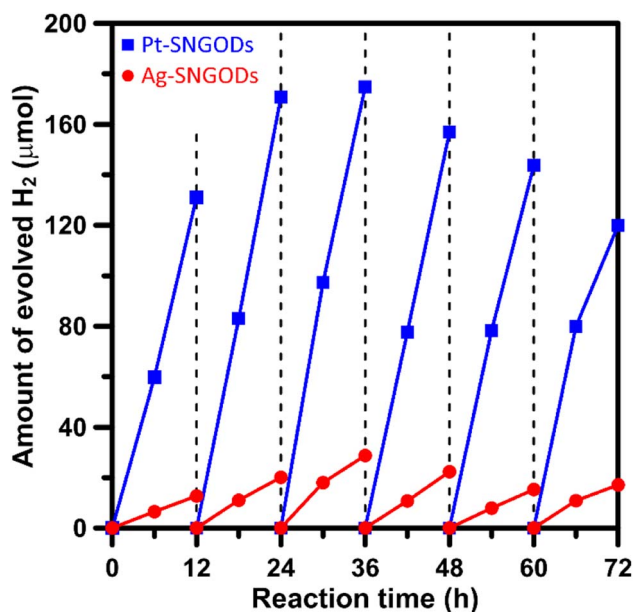


Fig. 3 Evolution of H_2 from the xylose solutions containing Pt-SNGODs and Ag-SNGODs under simulated solar irradiation (AM 1.5, 100 mW cm^{-2}).

and Table S2.† The Pt and Ag co-catalysts reduced the emission lifetime of the SNGODs because the co-catalysts quickly extracted the photogenerated electrons. Pt exhibited a more effective extraction than did Ag and resulted in a shorter lifetime.^{52,53} The presence of xylose lowered the τ value of Pt-SNGODs whereas a raised value was detected in Ag-SNGODs@xylose. The lowered value of Pt-SNGODs@xylose was due to the fact that the Pt co-catalyst effectively extracted the electrons donated by xylose (to the VB then to the CB). In contrast, the Ag co-catalyst could not effectively extract the donated electrons, thereby resulting in a prolonged PL emission.

Regarding the transfer feasibility of the photogenerated charges, the electronic structure of photocatalysts plays a vital

role to govern the redox progress during photocatalytic reactions. We subjected SNGODs, Pt-SNGODs, and Ag-SNGODs to analysis with ultraviolet photoelectron spectroscopy (UPS) to assess their valence band maximum (VBM). By deducting the width of the UPS spectra (Fig. S9†) from the excitation energy of 21.2 eV, the VBM values for SNGODs, Pt-SNGODs, and Ag-SNGODs were determined to be -6.46 eV , -5.84 eV , and -5.95 eV (*vs. vacuum*), respectively. The absorption spectra of Fig. 1 were converted to the Tauc plots (Fig. S10†) to obtain bandgap energies of 2.7, 2.1, and 2.2 eV for the SNGODs, Pt-SNGODs, and Ag-SNGODs, respectively. The conduction band minimum (CBM) level of the catalysts was determined according to the VBM levels and bandgap energies. Fig. 2 presents the summary of the electronic state levels of the catalysts in comparison with the hydrogen evolution reaction (HER) and oxygen evolution reaction (OER) levels in an aqueous solution at pH = 9. The Pt and Ag co-catalysts elevated the CBM levels and decreased the bandgap energy of SNGODs.

Photocatalytic reforming of xylose

Xylose is highly soluble in water due to its polar hydroxyl functional group.⁵⁴ In this study, an aqueous solution (250 mL) of xylose (1 g) at pH = 9 was used for the photocatalytic HER and xylose reforming.^{8,55} The photocatalytic reaction was conducted in a gas-enclosed system with side solar irradiation (AM 1.5 G at 100 mW cm^{-2}). To initiate the reaction, an equivalent amount of 5 wt% co-catalyst (Pt or Ag) precursors was charged into the solution containing SNGODs (0.2 g) and GO sheets (0.006 g). Under irradiation, the photogenerated electrons in SNGODs would be injected onto the GO sheets and therefore the co-catalyst nanoparticles were preferentially deposited on the GO sheets, as indicated in Fig. S2 and S3.† The presence of GO sheets in the system located the HER sites on the GO and protected SNGODs from the damages caused by the H_2 gas evolution.^{53,56–58} The photocatalytic reaction was undergone under a pressure of 75 mmHg.

Fig. 3 presents the amounts of H_2 produced from the system containing Pt-SNGODs and Ag-SNGODs. The reactions were conducted for 72 h with evacuating interventions every 12 h irradiation, and only H_2 was detected as the gas phase product (Fig. S11†). In the first 12 h, the H_2 produced was low because the deposition of the co-catalysts on the GO sheets was in progress in the initial period.²⁰ Both catalysts exhibited the highest HER activity after 36 h of the reaction. The total amount of H_2 generated over Pt-SNGODs was 7.7 times that over Ag-SNGODs. Although Ag-SNGODs was less active in HER, it was superior in the aspect of long-term stability. An intensive occurrence of HER on the Pt-SNGODs might have destroyed the GO sheets that were deposited with Pt, affecting the long-term stability of the Pt-SNGODs. Direct detection of the GO deterioration during the photocatalysis would be difficult. Nevertheless, we have conducted experiments to explicitly demonstrate the critical role of the GO sheets in the reaction. Fig. S12† presents the influence of the GO incorporation in the H_2 evolution from the xylose reforming over Pt-SNGOD catalysts. In the presence of the GO, the H_2 evolution was efficient and stable



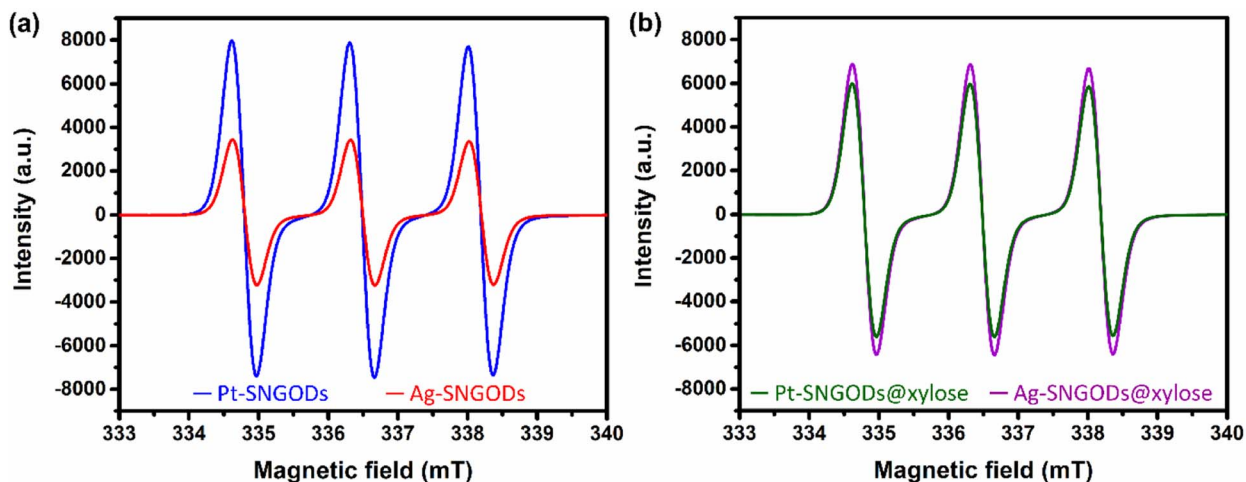


Fig. 4 The EPR signals of TEMPO contained in the Pt-SNGOD and Ag-SNGOD reacting solutions in the (a) absence or (b) presence of xylose. The solutions were irradiated with light of 462 nm in wavelength.

for at least 48 h, whereas the evolution exhibited serious decay without the GO incorporation. The incorporation of the GO sheets has resulted in the relocation of the H₂ evolution sites from SNGODs to the GO sheets, thereby minimizing the attack on the SNGODs by the reaction intermediates. This comparison indicated that the declined long-term stability of Pt-SNGODs could be attributed to the destroying GO sheet.

We have calculated the apparent quantum yields (AQYs) for H₂ production under 420 nm monochromatic illumination. The AQYs for the Pt-SNGOD and Ag-SNGOD systems were 6.8% and 2.8%, respectively. Table S3† presents the H₂ production results reported from literature for photocatalytic reforming of mono-sugar solutions. Our results were comparable to those obtained from other photocatalytic systems. Reporting

photocatalytic activity based on per gram of catalyst can be misleading, as the activity does not scale linearly with mass.^{59,60} The light harvesting efficiency depends on the catalyst dispersion situation and light scattering effects. Therefore, we have provided the AQYs for the H₂ production.

The low effectiveness of Ag in donating electrons for HER would lead to the accumulation of photogenerated electrons on the SNGODs. The electron presence can be explored using an electron-trapping agent, 4-hydroxy-2,2,6,6-tetramethylpiperidine-1-oxyl (TEMPO), which is a radical and its signal in the electron paramagnetic resonance (EPR) spectrum is quenched after trapping an electron from the environment. With the addition of TEMPO, the reacting solutions after irradiation, in the presence or absence of xylose, were analyzed

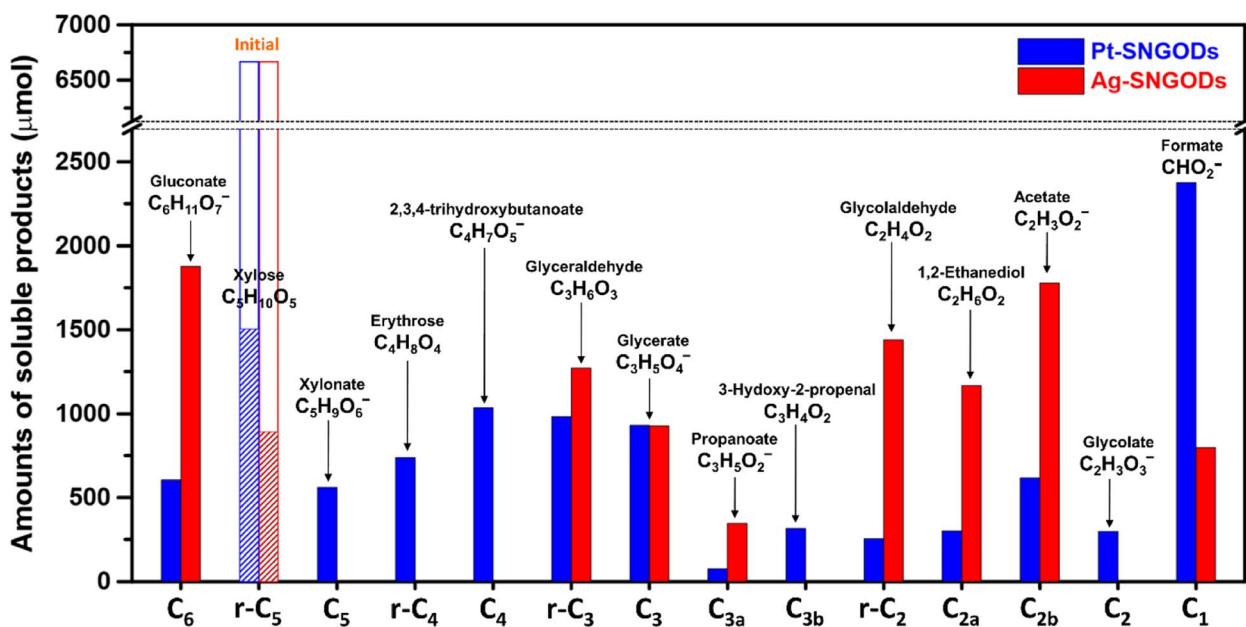


Fig. 5 Product compositions in the liquid phase after the photocatalytic reforming of xylose for 72 h.



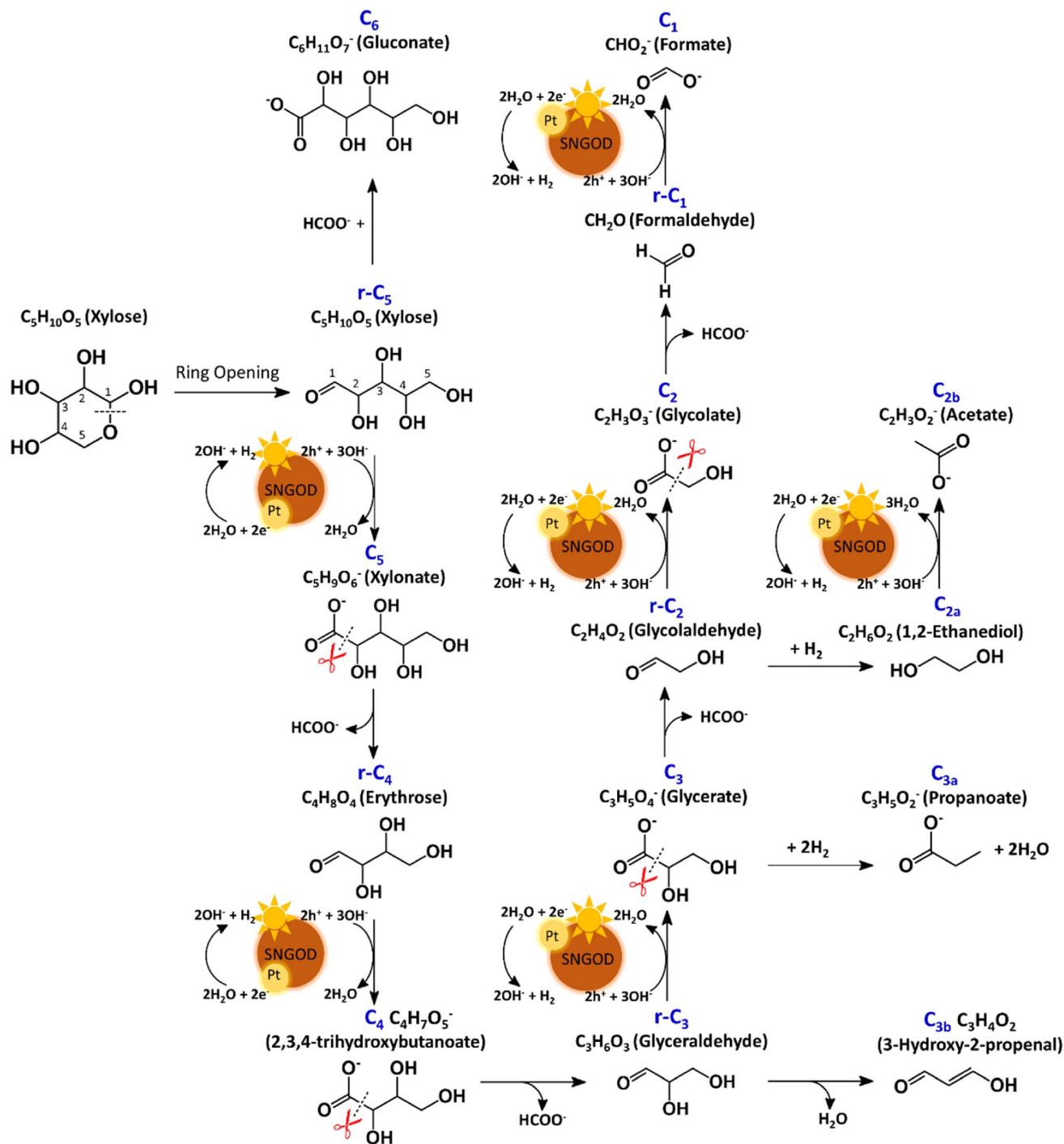


Fig. 6 Proposed mechanism of photocatalytic reforming of xylose over Pt-SNGODs.

using EPR. Fig. 4a presents the xylose-absent EPR spectra, in which the 1 : 1 : 1 signal intensity of the added TEMPO would be reduced with electron trapping.⁶⁴ Under the xylose-absent condition, the Ag-SNGOD solution exhibited a much lower EPR signal than the Pt-SNGOD solution, indicating that a large number of photogenerated electrons on Ag-SNGODs had interacted with the TEMPO radicals to minimize the EPR signal of TEMPO. Fig. 4b presents the xylose-present EPR spectra. Comparison of Fig. 4a and b depicted that the TEMPO signal

with the Ag-SNGODs was substantially increased by the xylose addition, indicating that xylose acted as an acceptor to consume the photogenerated electrons on the Ag-SNGODs and to result in the increased TEMPO signal. As to the Pt-SNGOD solution, Pt is efficient in transferring electrons for the HER. Xylose acts as a sacrificial electron donor to supply the catalyst electrons and promote the electron-trapping probability for TEMPO, thereby leading to a slight decrease in the EPR signal intensity for Pt-SNGOD solution. This EPR analysis indicated that xylose



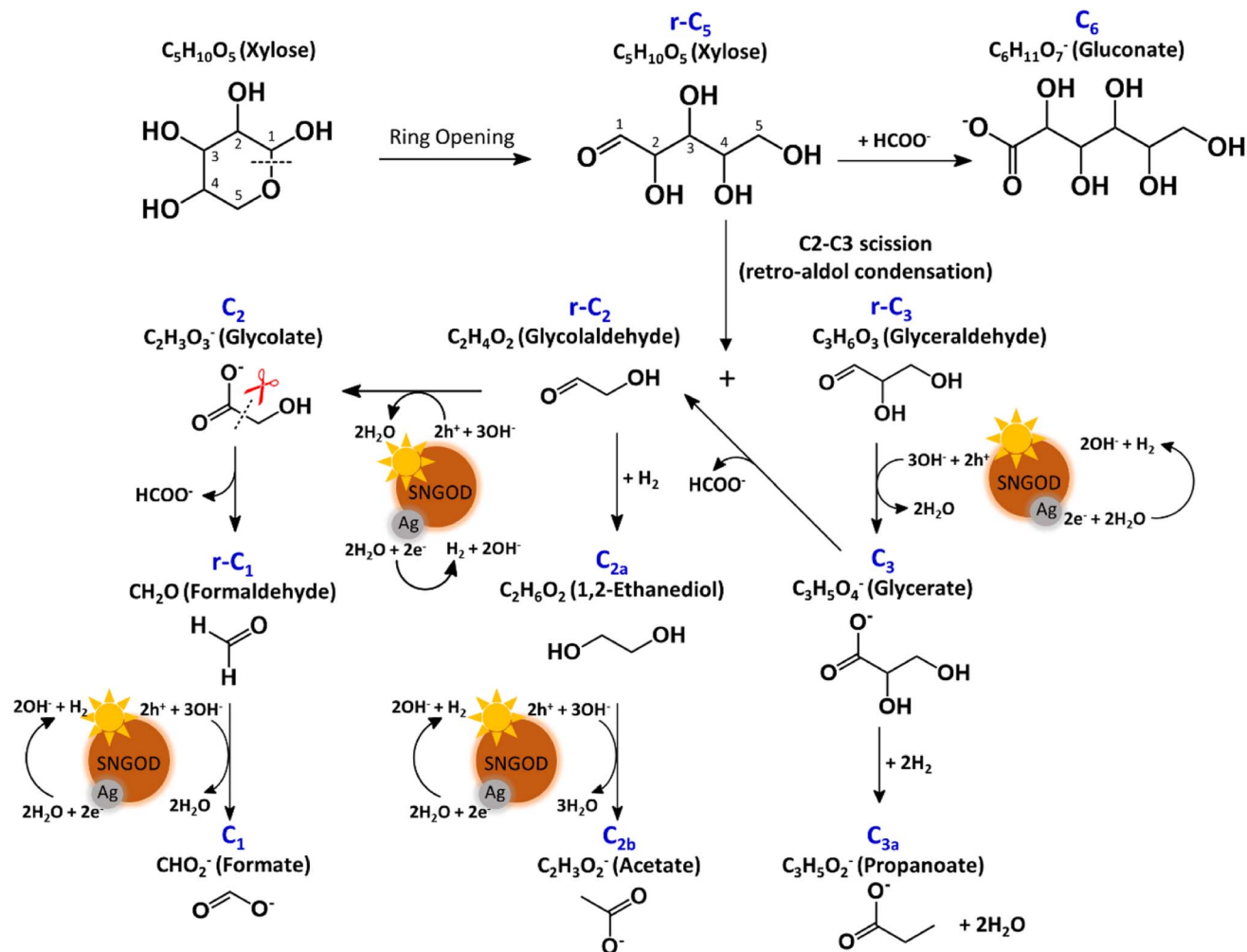


Fig. 7 Proposed mechanism of photocatalytic reforming of xylose over Ag-SNGODs.

might have accepted the accumulated electrons in the Ag-SNGODs in addition to acting as an electron donor to the VB of the SNGODs.

Besides the photocatalytic HER, the oxidative photo-reforming of xylose into other chemical compounds was investigated. In the liquid phase, ring opening of xylose, after interacting with water, would occur prior to xylose oxidation by the photogenerated holes.^{62,63} The liquid products from photo-reforming over Pt- and Ag-SNGODs were identified by the GC-MS and LC-MS analyses. GC-MS and LC-MS were used to identify products with mass-to-charge ratios (m/z) less than 50 and more than 50, respectively.²⁰ Before analysis, the samples were filtered to remove the catalyst particles. LC-MS analysis was conducted in a negative-ion mode, which created a stable deprotonated ionic molecule, namely, $[M-H]^-$.⁶⁴ Deprotonation allowed the molecular weight in solution samples to be determined. The precise chemical structure identification of these compounds was then validated using the LC-MS-MS analytical findings, in which the parent compounds were fragmented as seen in Fig. S13–S27.† Meanwhile, the products confirmed from the GC-MS analysis were shown in Fig. S28–S31,† with the MS

analysis validated the structures of the GC-MS products (Fig. S29 and S31†). For the quantitative analysis, the amounts of the products were determined based on the prepared standard samples.

After reaction for 72 h, Fig. 5 shows that Pt-SNGODs produced a greater variety of products than Ag-SNGODs, *i.e.*, 14-*vs.*-9 in the compound type. The LC-MS analyzed products from Pt-SNGODs were C_6 ($C_6H_{11}O_7^-$, gluconate), r- C_5 ($C_5H_{10}O_5$, xylose), C_5 ($C_5H_9O_6^-$, xylonate), r- C_4 ($C_4H_8O_4$, erythrose), C_4 ($C_4H_7O_5^-$, 2,3,4-trihydroxybutanoate), r- C_3 ($C_3H_6O_3$, glyceraldehyde), C_3 ($C_3H_5O_4^-$, glycerate), C_{3b} ($C_3H_3O_2^-$, 3-hydroxy-2-propenal), r- C_2 ($C_2H_4O_2$, glycolaldehyde), and C_2 ($C_2H_3O_3^-$, glycolate). For Ag-SNGODs in the LC-MS analysis, the products were C_6 , r- C_5 , r- C_3 , C_3 , and r- C_2 . Meanwhile, in the GC-MS analysis, both catalysts generated the same types of products, namely C_{2b} ($C_2H_3O_2^-$, acetate), C_{3a} ($C_3H_5O_2^-$, propanoate), C_1 (CHO_2^- , formate), and C_{2a} ($C_2H_6O_2$, 1,2-ethanediol). Fig. 5 shows that C_6 was the major product in the liquid phase for Ag-SNGODs, whereas C_1 was the main liquid product of photocatalytic reforming with the Pt co-catalyst.



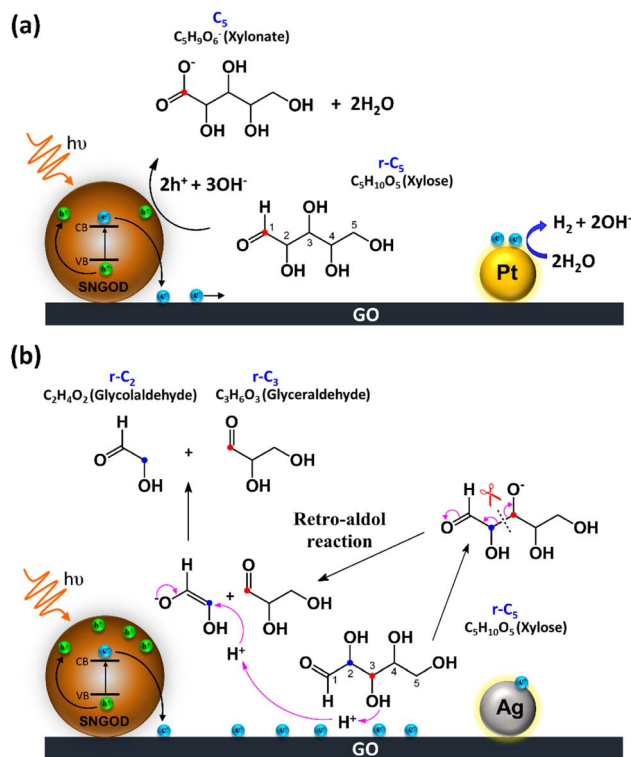


Fig. 8 The charge distributions of the SNGODs when associated with different co-catalysts. (a) Pt co-catalyst for oxidative xylose reforming (Pt-SNGODs were active in consuming photogenerated electrons for H_2 evolution). (b) Ag co-catalyst for retro-aldol reaction (Ag-SNGODs were less effective in HER).

Mechanism of xylose reforming over Pt- and Ag-SNGODs

The products from photocatalytic xylose reforming over Pt-SNGODs comprised the $r-C_x$ and C_x species with x ranging through 1–5, indicating that the reforming proceeded through successive alternation of C-eliminating hydrolysis and photocatalytic oxidation.²⁰ Fig. 6 presents the Pt-SNGODs mechanism proposed based on the products shown in Fig. 5. The ring-form xylose feed was firstly opened into linear-form xylose ($r-C_5$), which was then converted into carboxylate C_5 through photocatalytic oxidation. The downgrade reforming of $r-C_5$ to the terminal C_1 comprised successive alternating steps: photocatalytic oxidation of the aldoses/aldehydes/alcohols $r-C_x$ to carboxylates (C_x) prior to C-eliminating hydrolysis by removing one formate (C_1) molecule.⁶⁵ Along with the photocatalytic oxidation, water reduction by the photogenerated electrons was taking place to produce H_2 . Apart from the principal pathway of successive photocatalytic oxidation and hydrolysis, dehydration of $r-C_3$ might have occurred to produce $C_3H_4O_2$ (C_{3b}), which was detected in the product analysis. Because of the occurrence of photocatalytic HER in the solution, hydrogenolysis of C_3 and $r-C_2$ might have occurred to produce $C_3H_5O_2$ (C_{3a}) and $C_2H_6O_2$ (C_{2a}), respectively (Fig. 6). C_{2a} might be photocatalytically oxidized to $C_2H_3O_2^-$ (C_{2b}). The branched hydrogenolysis pathways must have played an important role in the reforming over

Ag-SNGODs because of the high contents of C_{2a} , and C_{2b} in the products (see Fig. 5).

In the photo-reforming reaction using Ag-SNGODs, Fig. 5 shows that the products in the solution comprised high proportions of C_3 - and C_2 -compounds but no C_4 -compounds, indicating that $r-C_5$ proceeded with cleavage on C2–C3 through the retro-aldol reaction mechanism to produce $r-C_2$ and $r-C_3$, as indicated in Fig. 7.⁶⁶ The retro-aldol reaction generally takes place in a basic environment and could be catalyzed by a Lewis base.⁶⁷ The $r-C_2$ and $r-C_3$ from the cleavage were further downgraded to C_1 through steps similar to those proposed for the reforming over Pt-SNGODs (see the comparison between Fig. 6 and 7). Notably, the content of C_{2b} (an acetate), derived from C_{2a} (an alcohol), was high over Ag-SNGODs relative to that over Pt-SNGODs. The hydrogenation of $r-C_2$ (an aldehyde) into C_{2a} (a carboxylate), attributable to the insufficient photocatalytic activity of Ag-SNGODs. As to the C–C coupling interaction, C_6 was profoundly produced from the photoreforming over Ag-SNGODs, with which the contents of $r-C_5$ and C_1 in the solution were substantially lower than those in the solution for the reforming over Pt-SNGODs (Fig. 5). The coupling of $r-C_5$ with C_1 must have been responsible for the formation of C_6 . A study reported that C_1 reacted with $r-C_5$ through a nucleophilic addition reaction,⁶⁸ in which the carboxyl group of C_1 attached itself to the formyl group on $r-C_5$. The coupling of $r-C_5$ with C_1 was effective in the reforming over Ag-SNGODs (Fig. 7).

To test the proposed mechanistic pathways, we randomly performed photocatalytic reforming of intermediates glycolaldehyde and glyceraldehyde (rC_2 and rC_3 , respectively) over Pt-SNGODs and Ag-SNGODs for HER. Fig. S32† shows that both reforming reactions using rC_2 and rC_3 produced H_2 , with higher HER rates over Pt-SNGODs. The results of the tests on rC_2 and rC_3 were consistent with mechanistic pathways proposed in Fig. 6 and 7.

Fig. 8 presents the charge distributions of the irradiated SNGODs when associated with different co-catalysts. The Pt-SNGODs effectively donated electrons into the aqueous solution to produce H_2 , resulting in the effective oxidation of xylose by the photogenerated holes. The oxidation occurred on the aldehyde end of xylose because the electron-rich oxygen of the formyl group attracted the hole-loaded SNGODs. In contrast, the Ag-SNGODs were less effective in donating electrons for HER, thereby resulting in the accumulation of electrons on the GO sheets and the low efficiency of hole donating to oxidize xylose, even if the holes on the SNGODs (acting as a Lewis acid) still attracted the formyl oxygen of the xylose. With the electron accumulation, the GO sheets would interact with the hydroxyl group on the C3 of xylose, converting xylose into an alkoxide intermediate where the oxygen atom at C3 carried a negative charge. This intermediate would then proceed with the retro-aldol reaction for breaking the C2–C3 bond to produce the $r-C_2$ and $r-C_3$ fragments. The interaction between xylose and the accumulated electrons has been reflected by the EPR analysis shown in Fig. 4.

The lower xylose-oxidizing ability of Ag-SNGODs, relative to that of Pt-SNGODs, also resulted in a higher rate in the C–C



coupling of radicals from reductions to produce C_6 . According to Fig. 3, Ag-SNGODs were not sufficiently effective in coupling H-radicals, derived from photocatalytic reduction of H_2O ,⁶⁹ for the occurrence of HER. The abundant xylose and formate in the reacting system would interact with the H-radicals to produce r- C_5 - and C_1 -derived radicals, which would subsequently proceed with C-C coupling to form C_6 . This explains the high C_6 content in the products of the photocatalytic xylose reforming over Ag-SNGODs. This study has demonstrated how the co-catalyst property affected the charge accumulation degree and thus the product formation mechanism.

Conclusions

This study explores the impact of co-catalysts (Pt and Ag) when combined with the SNGOD photocatalyst on the photocatalytic reforming of xylose (C_5) into H_2 and C_1 - C_6 compounds. Pt was more effective than Ag in extracting photogenerated electrons for HER. The accumulation of electrons in Ag-SNGODs was disclosed using EPR analysis. Regarding the oxidative reforming of xylose, Pt-SNGODs effectively converted xylose into C_5 - C_1 , through successive alternation of C-eliminating hydrolysis and photocatalytic oxidation, and moderately produced C_6 through C-C coupling. With Ag-SNGODs, the accumulated photo-generated electrons interacted with xylose to cleave the r- C_5 (xylose), through the retro-aldol reaction, into r- C_3 and r- C_2 , which were then successively decomposed into C_3 - C_1 . The EPR analysis evidenced the interaction between xylose and the electrons accumulated in Ag-SNGODs. A profound amount of C_6 was produced over Ag-SNGODs, indicating that the HER-inactive Ag-SNGODs provided sufficient H-radicals to generate formate- and xylose-derived radicals for subsequent coupling into C_6 . Overall, this research has shed light on the selective production of products in photocatalytic biomass reforming, emphasizing the significance of co-catalyst selection.

Experimental section

Synthesis of photocatalyst

Using a modified Hummers' method, GO was prepared from natural graphite powder (Bay carbon, SP-1, USA).⁷⁰ 5 g of graphite was mixed with 2.5 g of $NaNO_3$ ($\geq 99\%$, Merck, Germany) in 150 mL of concentrated H_2SO_4 (18 M, Wako, Japan). 15 g of $KMnO_4$ ($\geq 99\%$, J.T. Baker, USA) was gradually added with stirring while maintaining the temperature at 35 °C in an ice bath for 4 h. Then, 250 mL of deionized (D.I.) water was slowly added, and the mixture was stirred for 15 min at 98 °C. Another 800 mL of D.I. water was added and stirred for 30 min to terminate the oxidation process. The GO preparation was completed by adding 10 mL of H_2O_2 . The resulting slurry was washed and filtered six times. GO was obtained by freeze-drying the slurry for 24 h.

A ratio of 1 : 1 between GO and sulfur powder (99.999%, Acros, USA) was used to make the SNGODs by calcining the mixture in an NH_3 flow (150 mL min^{-1}) at 600 °C for 3 h. Every 0.2 g of the obtained mixture was oxidized in 60 mL of concentrated HNO_3 (60%, Showa, Japan) under sonication for

12 h at room temperature. Then, the HNO_3 was removed by evaporating the liquid phase at 140 °C, producing a dot sample. The dot samples were autoclaved in a NH_3 solution (30 wt% in water, Sigma-Aldrich, USA) for 5 h at 140 °C. A heating process at 100 °C, along with stirring, was applied to remove the residual NH_3 solution. To remove the large dots, the solution obtained was sieved with Sartoclon Slice 200 with Hydrosart membranes (Sartorius, Germany). Finally, the catalyst named SNGODs was ready to be used.

Photocatalyst characterization

Transmission electron microscopy (TEM, Jeol 2100 F, Japan) was used to determine the microstructure of the catalyst. The chemical composition of the catalysts was quantitatively analyzed using X-ray photoelectron spectroscopy (XPS) with AXIS Ultra DLD (Kratos, UK) equipped with Al $K\alpha$ radiation. The C_{1s} , N_{1s} , and S_{2p} spectra were split into component peaks using a Gaussian fitting method. The optical absorbance of the catalysts was measured using a Hitachi U-4100 (Japan) spectrophotometer. Suspensions with a concentration of approximately 0.18 mg mL^{-1} were put in a 1 cm quartz cuvette. The photoluminescence (PL) spectrum of SNGODs in aqueous solution (0.18 mg mL^{-1}) was measured at ambient temperature using a fluorescence spectrophotometer (F-700, Hitachi, Japan). Time-resolved photoluminescence (TRPL) analysis was performed at an excitation wavelength of 405 nm using a spectrofluorometer (FS5, Edinburgh Instruments, UK).

Photocatalytic activity measurements

A powder of xylose (Sigma-Aldrich, St. Louis, MO, USA) was used as a substrate or hole scavenger in all the experiments. 1 g of xylose was dissolved in 20 mL of D.I. water, and then 0.2 g of SNGODs, 0.006 g of GO, and 5 wt% of either $H_2PtCl_6 \cdot 6H_2O$ (26 mg, 99%, Alfa Aesar, USA) or $AgNO_3$ (25 mg, >99%, Acros Organics, UK) were added. The solution was diluted to 250 mL with D.I. water, and the pH was adjusted to 9. During all photocatalytic reactions, the closed glass gas circulation system was connected to a Pyrex reaction vessel with side-irradiation (469.2 mL with a flat illumination window of 4 cm in diameter) for side-irradiation. The source of the irradiation was a solar simulator (Oriol Instruments, 100 mW cm^{-2} , USA) connected to an air mass AM-1.5 G filter. A photodetector (Oriol Instruments, model 71964, USA) was used to determine the intensity of the light irradiated on the system. The gas product was detected by a gas chromatograph (GC, Hewlett-Packard 7890, USA) with a molecular sieve 5A column, thermal conductivity detector, and argon as the carrier gas. The photo-reforming reaction was run for 3 days with evacuating interventions at 75 mmHg every 12 h.

The AQY values were calculated according to: $AQY (\%) = (\text{moles of } H_2 \text{ produced} \times 2 / \text{incident photon flux}) \times 100\%$, where the incident photon flux was $49.96\ \mu\text{mol h}^{-1}$ at 420 nm.⁵⁹ The bandpass wavelengths of 420 nm correspond to the 20BPF10-420 (Newport, USA) band-pass filters used. The power intensity of monochromatic light was measured using a Si photodiode integrating sphere detector (Newport model 819D-



SL-2-CAL2) coupled with an optical power meter (Newport 1919-R). The 420 nm radiation intensity was 0.20 mW cm^{-2} over an irradiated area of 19.61 cm^2 .

Electron paramagnetic resonance analysis

We performed the electron paramagnetic resonance (EPR) measurements using a Miniscope MS 5000 EPR spectrometer (Freiberg Instruments, Germany) equipped with a 0.575 W, single wavelength of 462 nm, high-power light-emitting diode. Only 50 μL of sample solution was needed. The sample was placed in a capillary tube and then inserted into the EPR hole. 4-Hydroxy-TEMPO was used as the electron trap. The field modulation, modulation frequency, scan range, and microwave power were 0.2 mT, 100 kHz, 15 mT, and 10 mW, respectively.

GC-MS and LC-MS for compounds analysis

The liquid products from photo-reforming were analyzed using a GC system (GC-2010, Shimadzu, Japan) equipped with an HP-INNOWAX column ($60 \text{ m} \times 0.25 \mu\text{m} \times 0.32 \text{ mm}$; Agilent Technologies, USA) and a mass spectrometer (GCMS-QP2010, Shimadzu, Japan), with helium gas serving as the carrier gas. A sample volume of 1 μL was used for analysis. The operating parameters included a pressure of 91.8 kPa, a total flow rate of 24.7 mL min^{-1} , a column flow rate of 2.2 mL min^{-1} , a linear velocity of 37.9 cm s^{-1} , a purge flow of 5 mL min^{-1} , a split ratio of 10, an oven temperature of $40 \text{ }^\circ\text{C}$, and an injection temperature of $150 \text{ }^\circ\text{C}$. The GC temperature program involved an initial temperature of $40 \text{ }^\circ\text{C}$ for 15 min, followed by a ramp of $+25 \text{ }^\circ\text{C min}^{-1}$ to $200 \text{ }^\circ\text{C}$, which was maintained for 20 min. Mass spectrometry parameters included a 70 eV ionization energy, an ion source temperature of $230 \text{ }^\circ\text{C}$ in electron ionization mode, an interface temperature of $250 \text{ }^\circ\text{C}$, a solvent cut time of 1 min, a detector gain of 0.8 kV, a threshold of 5, a start time of 2 min, an end time of 5.58 min, and a selected ion monitoring mode for a mass range of 0 to 100 m/z . The acquired data were analyzed using GC-MS Solution Software (version 4.30, Shimadzu).⁶⁹

Additionally, an UltiMate 3000 HPLC System (Thermo Fisher Scientific, USA) and an EC 100/2 Nucleoshell Hilic column, $2.7 \mu\text{m}$ (Macherey-Nagel, Germany) were used. The sample was filtered using $0.22 \mu\text{m}$ membrane filters before analysis. A solvent mixture of 80% acetonitrile and 20% methanol was used, with the column temperature set to $40 \text{ }^\circ\text{C}$ and the flow rate of $300 \mu\text{L min}^{-1}$. The analysis run time was 10 min. Q Exactive Plus Q-Orbitrap MS instruments (Thermo Fisher Scientific, USA) were used in this study. Electrospray-ionization MS was used for analyzing the standard samples and xylose, while high-energy collision dissociation (HCD) was employed for MS-MS analysis. The data acquisition mode targets fragment ions from HCD events with a normalized collision energy (NCE) of 35 eV on eight precursors (195.04967, 165.03900, 149.04402, 119.03340, 105.01778, 89.02254, 89.02287, and 75.00724), detected using an Orbitrap mass analyzer. A negative mode was applied with a 2.5 kV spray voltage, a sheath gas flow rate of 35, an auxiliary gas flow rate of 10, a capillary temperature of $275 \text{ }^\circ\text{C}$, and an S-lens level of 50. The resolutions were set at 70 000

and 17 500 for full MS and MS-MS, respectively. The automatic gain control target was 3×10^6 for full scan and 2×10^5 for MS-MS, with an injection time of 200 ms. Data analysis, including full MS spectra and fragment ions, was conducted using Xcalibur 4.1.31.9.

Author contributions

Van-Can Nguyen: conceptualization, data curation, formal analysis, methodology, writing – original draft, writing – review & editing. Meyta Sanoe: investigation, data curation, formal analysis, methodology, writing – original draft. Novy Pralisa Putri: data curation, formal analysis. Yuh-Lang Lee: data curation, validation. Hsisheng Teng: conceptualization, project administration, resources, supervision, validation, investigation, writing – original draft, writing – review & editing. All authors approved and contributed to the manuscript.

Conflicts of interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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