A spongy nickel-organic CO₂ reduction photocatalyst for nearly 100% selective CO production

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Solar-driven photocatalytic conversion of CO₂ into fuels has attracted a lot of interest; however, developing active catalysts that can selectively convert CO₂ to fuels with desirable reaction products remains a grand challenge. For instance, complete suppression of the competing H₂ evolution during photocatalytic CO₂-to-CO conversion has not been achieved before. We design and synthesize a spongy nickel-organic heterogeneous photocatalyst via a photochemical route. The catalyst has a crystalline network architecture with a high concentration of defects. It is highly active in converting CO₂ to CO, with a production rate of ~1.6 × 10⁴ μmol hour⁻¹ g⁻¹. No measurable H₂ is generated during the reaction, leading to nearly 100% selective CO production over H₂ evolution. When the spongy Ni-organic catalyst is enriched with Rh or Ag nanocrystals, the controlled photocatalytic CO₂ reduction reactions generate formic acid and acetic acid. Achieving such a spongy nickel-organic photocatalyst is a critical step toward practical production of high-value multicarbon fuels using solar energy.

INTRODUCTION

Rapid fossil fuel consumption induces environmental burden and energy crisis (1–3). Excessive anthropogenic CO₂ emission is a significant concern because of its hastening impact on climate change (4–6), acidification of ocean (7), crop yield reduction (8), extinction of animal species (9), and damage to human health (10, 11). Removal of excessive CO₂ from the atmosphere (12), particularly converting CO₂ to fuels using solar energy, is currently a global research endeavor (13–15). Discovering novel catalysts that can reduce the stable CO₂ molecules and convert them to liquids with high activity and selectivity is essential (13, 14). To date, despite the progress that has been made in investigating the photocatalytic reduction of CO₂ (15–19), controlling the reaction to yield a specific product among many possible reaction species, including CO, H₂, CH₄, and formic acid, remains a great challenge (16, 20, 21). Finding photocatalysts that can efficiently convert CO₂ to CO and largely suppress other competing photocatalytic reactions, such as H₂ evolution, would be a critical step forward toward practical solar-to-fuels conversion for the production of high-value multicarbon fuels (15, 17, 22).

We recently developed a laser-chemical method and synthesized active transition metal hydroxide catalysts with a high concentration of defects for water oxidation (23). Specifically, we used an unfocused infrared laser to initiate the reactions between transition metal ions and triethylene glycol (TEG) and obtained a series of metal hydroxide–TEG composites with a distorted layered structure (23). This disordered structure enhances the accessibility of water molecules to the active sites and enables efficient electrocatalysis of alkaline water oxidation (23). Such a laser-chemical strategy may be applied to the discovery of many other catalysts, for instance, novel nanostructured metal-organic heterogeneous catalysts for CO₂ reduction reaction.

When designing catalysts for CO₂ reduction, the material’s ability to capture the CO₂ molecules is another significant consideration (24). Metal-organic frameworks (MOFs) with high surface area and tunable pores have been used for gas capture and heterogeneous catalysis (25, 26). Typically, MOFs have highly ordered crystalline structures constructed by coordinating metal ions or clusters with rigid organic linkers, most often the aromatic carboxylic acid molecules (27), such as terephthalic acid (TPA). In light of MOF structure design, we replace part of the rigid linkers (for example, TPA) in traditional MOFs with soft molecules (for example, TEG) by laser, considering the comparable molecular length of TEG to TPA (fig. S1). When the TEG molecules, which lack essential carboxylic groups for the perfect framework construction, are weaved into the metal–TPA framework, their substitution of TPA linkers may frustrate the growth of highly ordered MOF crystals, resulting in disordered and defective metal-organic hybrids for effective CO₂ fixation.

Here, we design a model metal-organic CO₂ reduction catalyst, with Ni₂⁺ ions as active metal centers, TPA as a rigid linker, TEG as a soft linker, and dimethylformamide (DMF) as a solvent, via laser-induced solution reactions. The as-synthesized catalyst, labeled as Ni(TPA/TEG), has a crystalline network architecture with considerable defects and performs nearly 100% selective gas production (CO over H₂ evolution) with a high CO production rate of ~1.6 × 10⁴ μmol hour⁻¹ g⁻¹. Further metal decorations (that is, Rh and Ag) of the Ni(TPA/TEG) catalyst lead to controlled photocatalytic CO₂ reduction reactions that generate formic acid and acetic acid.

RESULTS AND DISCUSSION

Structure determination of the Ni(TPA/TEG) catalyst

As shown in Fig. 1A, the Ni(TPA/TEG) composite forms a disordered spongy network structure, in which Ni, O, and C are uniformly distributed (fig. S2). In comparison, the solution without TEG, Ni(TPA) only,
forms large particles (Fig. 1B). A three-dimensional electron tomographic reconstruction of the spongy Ni(TPA/TEG) architecture reveals various mesopores in the structure (Fig. 1C and movies S1), which closely resembles the pore features identified from the N2 physisorption measurements (Fig. S3). Figure 1D shows a typical transmission electron microscopy (TEM) image of the spongy Ni(TPA/TEG) composite, where defective lattices with a \( d \)-spacing of 1.02 nm are captured. To further interpret the structure of Ni(TPA/TEG) composite, we acquire a scanning nanobeam diffraction data set using an electron beam with a size of ~3 nm, a total beam current of ~5 pA, and an exposure time of 0.5 s, where the electron beam damage to the metal-organic material has been evidently minimized (Fig. 1E). Single-crystalline diffraction patterns along the [100] and [111] orientations of the orthorhombic Ni(TPA/TEG) composite (movies S2 and S3) are captured from two different regions of the spongy network (Fig. 1F and movies S2 and S3), showing an orthorhombic structure similar to that of the Ni(TPA) particles (fig. S4). Changes of the diffraction patterns are observed from movies S2 and S3, indicating defects (that is, grain boundaries) in the spongy Ni(TPA/TEG) catalyst (fig. S5).

To verify that the soft TEG molecules have been incorporated into the Ni(TPA) framework (Fig. 2A), we compare the structure of laser-synthesized Ni(TPA) and Ni(TPA/TEG) composites in detail. The x-ray diffraction (XRD) pattern (Fig. 2B) shows that the Ni(TPA) composite has an orthorhombic structure similar to that of Ni(TPA), but slight differences exist in the peak positions and widths of the x-ray lines, which may result from the exchange of linkers and solvent molecules in the structure (28). In the Fourier transform infrared (FTIR) spectra (Fig. 2C), we can see that both samples have clear bands of \( \nu (\text{COO}^-) \) (1375 and 1575 cm\(^{-1}\)) and ring breathing (815 cm\(^{-1}\)) from the TPA linkers. Characteristic bands of \( \delta \text{NCO} \) (690 cm\(^{-1}\)) and \( \nu (\text{CO}) \) (1685 cm\(^{-1}\)) from DMF molecules are found in Ni(TPA), indicating that DMF molecules may reside in the MOF cavities and coordinate with Ni\(^{2+}\) through carbonyl groups (28). Meanwhile, distinct bands of \( \nu (\text{OH}) \) (1065 and 3374 cm\(^{-1}\)) related to TEG linkers are found exclusively in Ni(TPA/TEG), and no DMF bands are detected, indicating that TEG molecules exist in Ni(TPA/TEG); DMF molecules that originally occupied the Ni(TPA) framework cavities prefer to leave. The EDX spectrum (Fig. 2D) also shows that no nitrogen (from DMF) can be detected in Ni(TPA/TEG). Thermo-gravimetric analysis (TGA) curves in Fig. 2E, both displaying three stages of mass losses, indicate differences between the two samples. For Ni(TPA), the mass loss measurements of 6, 18, and 50% are from H\(_2\)O, DMF, and TPA, respectively (28); for Ni(TPA/TEG), the mass losses of 16, 26, and 26% are from H\(_2\)O, TEG, and TPA, respectively. The large differences in the mass loss of TPA in the two samples (50% versus 26%) indicate that soft TEG molecules have replaced part of the rigid TPA linkers, causing the formation of a spongy Ni(TPA/TEG) composite.
network. Because of the varying chemical environment of Ni\(^{2+}\), the Ni\(^{2p}\) peaks in the x-ray photoelectron spectroscopy (XPS) spectra (Fig. 2F) shifted to the right in Ni(TPA/TEG) compared to Ni(TPA). We also find that it is more effective to use lasers to cross-link the soft TEG and rigid TPA molecules together with the Ni\(^{2+}\) centers than to use a traditional heating process (see the Supplementary Materials). Laser irradiation appears to produce Ni-TEG building units that are indispensable for the spongy Ni(TPA/TEG) network construction (Fig. 2A). Because of the soft character of TEG molecules, various inhomogeneous configurations of Ni-TEG units can be generated. Mismatches between the soft Ni-TEG units and the rigid Ni-TPA units may introduce considerable defects, leading to the formation of disordered spongy Ni(TPA/TEG).

**Evaluation of the photocatalytic activity for CO production**

We apply the as-synthesized Ni-organic composites for visible light–driven photocatalytic CO\(_2\) reduction in a solvent mixture of acetonitrile/water [considering the high solubility of CO\(_2\) in acetonitrile (29)] under mild reaction conditions (20°C and 400 torr of CO\(_2\)), with triethanolamine (TEOA) as a sacrificial reducing agent and Ru(bpy)\(_3\)Cl\(_2\) as a photosensitizer (18, 30). Five samples (figs. S6 to S8), that is, Ni(TPA) (L), Ni(TPA) (H), Ni(TPA/TEG) (L), Ni(TPA/TEG) (H), and Ni(TEG) (L), synthesized by both laser irradiation (L) and traditional heating (H), are examined. Figure 3A shows the CO evolutions from these five Ni-organic catalysts in a 6-hour photocatalytic reaction. The spongy Ni(TPA/TEG) (L) composite shows the highest activity, and the amount of CO is 95.2 μmol after a 2-hour reaction, giving a CO production rate of 15,866 μmol hour\(^{-1}\) g\(^{-1}\), which is several times higher than that from other samples. The total amount of CO produced on the spongy Ni(TPA/TEG) catalyst in 6 hours reaches 136.9 μmol (Fig. 3A), giving a turnover number of 11.5 for the 6-hour reaction (table S1). The CO production rate is also superior compared with many other reported heterogeneous CO evolution photocatalysts to the best of our knowledge (17), such as the Co\(_3\)O\(_4\) platelets with [Ru(bpy)\(_3\)]Cl\(_2\) as a photosensitizer (3523 μmol hour\(^{-1}\) g\(^{-1}\)) (18), the sensitized TiO\(_2\) particles with enzyme as a cocatalyst (300 μmol hour\(^{-1}\) g\(^{-1}\)) (31), and the sensitized BaLa\(_4\)Ti\(_4\)O\(_{15}\) particles with Ag as a cocatalyst (22 μmol hour\(^{-1}\) g\(^{-1}\)) (16). Note that the soluble homogeneous metal complex catalysts, which have also been investigated for controlled CO\(_2\) reduction (32–34), are not categorized here for comparison.

By testing the 2-hour yield of CO on different amounts of the Ni(TPA/TEG) catalyst, we obtain a roughly linear relationship between the amount of evolved CO and the amount of the catalyst (Fig. 3B). However, kinetically, we found that the CO production rate actually decreases with the increase in the amount of the catalyst.
where 1.0 mg of the Ni(TPA/TEG) catalyst gives a CO production rate of ~26,620 m mol hour$^{-1}$ g$^{-1}$ in the same solution, indicating that more electrons generated from the photosensitizer molecules could have been transferred to the catalytic active sites. We have also tested the reusability of the spongy Ni(TPA/TEG) catalyst upon each 2-hour photocatalysis, where the catalyst has kept its activity and selectivity after recycling (Fig. 3C). It also exhibits excellent structural stability, and no obvious structural change is found after 24 hours of photocatalysis (fig. S10). To confirm the origin of the as-produced CO, we use isotopic $^{13}$CO$_2$ as feedstock gas for the photocatalytic reduction and examine the products by gas chromatography–mass spectrometry (GC-MS). A major signal at a mass/charge ratio of 29 on the mass spectrum corresponding to $^{13}$CO (Fig. 3D, red lines) appears, which confirms that the as-detected CO originates from the CO$_2$ gas source (fig. S11).

Transition metal ions with switchable electronic states have long been considered promising active sites for diverse photocatalytic or electrocatalytic reactions, such as water splitting (35–37), CO oxidation (38), and CO$_2$ reduction (39). Consequently, we use the laser-chemical method to synthesize four additional samples and compare different...
metal ions, that is, Ni$^{2+}$, Co$^{2+}$, and Cu$^{2+}$ (fig. S12), as active sites for the photocatalytic CO$_2$ reduction. Results show that the spongy Ni(TPA/TEG) catalyst is still the most active catalyst for CO evolution, and the activity worsens with the incorporation of Co and Cu ions (Fig. 3E). The amount of CO generated on the Ni(TPA/TEG) catalyst is almost two times that of the CO generated from Co(TPA/TEG) in a 6-hour photocatalytic reaction. The Cu(TPA/TEG) catalyst barely generates any CO, which is distinctly different from the metal Cu catalyst that has superior activity for CO evolution from the electrocatalytic reduction of CO$_2$ (40).

**Evaluation of the CO production selectivity**

We find that CO (CO$_2$ + H$_2$O + 2e$^-$$\rightarrow$ CO + 2OH$^-$) is the only detectable gas product from the photocatalytic CO$_2$ reduction on the TPA-containing Ni-organic catalysts. H$_2$ evolution (2H$_2$O + 2e$^-$$\rightarrow$ H$_2$ + 2OH$^-$), usually acting as a major competing reaction in the CO$_2$ reduction system for many transition metal–based catalysts (18, 20), has been completely suppressed (table S1). Thus, a near 100% selectivity of CO production (over H$_2$ evolution) is achieved. Note that no other potential competing gas products, such as CH$_4$ and C$_2$H$_4$ (17), have been detected in our experiments either.

For comparison, we have detected considerable amounts of H$_2$ from other laser-synthesized M(TPA/TEG) (M = Ni, Co, Cu) catalysts (Fig. 3, E and F, and fig. S13), and CO selectivity values of 96, 70, 78.2, and 4.8% were measured for NiCo(TPA/TEG), Co(TPA/TEG), NiCoCu(TPA/TEG), and Cu(TPA/TEG), respectively. We have also detected a fair amount of H$_2$ (25.7 µmol) from the hydroxylated TPA-free Ni(TEG) catalyst in addition to the CO evolution (96.5 µmol) after a 6-hour reaction (fig. S14), which is analogous to the Ni-based hydroxides for practical H$_2$ evolution from the electrocatalysis of water (36).

**Tuning the selectivity for liquid fuels production from CO$_2$**

Furthermore, considering the potential of noble metal electrodes for CO$_2$ reduction (41, 42), we decorate the spongy Ni(TPA/TEG) with noble metal nanocrystals, that is, Rh and Ag, in pursuit of tuning the selectivity of liquid fuels production from the photocatalytic CO$_2$ reduction (15, 21). Figure 4 (A to C) shows the Ag-decorated Ni(TPA/TEG) catalyst, where Ag nanocrystals with an average diameter of ~6 nm are well dispersed on the Ni(TPA/TEG). We examine the liquid products after a 6-hour reaction on three catalysts, that is, undecorated Ni(TPA/TEG), Rh-decorated Ni(TPA/TEG), and Ag-decorated Ni(TPA/TEG) (Fig. 4D). For the Ni(TPA/TEG) catalyst without any decoration, we measure formic acid (HCOOH) with a concentration of 29.2 µM and acetic acid (CH$_3$COOH) with a concentration of 72.5 µM in addition to CO. With the decoration of Rh and Ag nanocrystals, the amounts of CO decrease drastically, whereas the amounts of liquid products significantly increase. Formic acid (HCOOH) with a concentration of 313.5 µM is mainly obtained on the Rh-decorated Ni(TPA/TEG) catalyst, and CH$_3$COOH with a concentration of 195.6 µM is the major product for the Ag-decorated Ni(TPA/TEG) catalyst, where the origin of CH$_3$COOH has been confirmed from the flowing CO$_2$ source by MS (fig. S15). Note that CO, ...
formic acid, and acetic acid reflect the overall product distribution of the photocatalytic CO₂ reduction reaction; no other liquid products, such as methanol, ethanol, or propanol, are detected in this experiment.

### Enhanced production of liquid fuels from photocatalytic reduction of CO

By assuming that the evolved CO may be further consumed for the production of acids, we also conduct control experiments using CO instead of CO₂, as the gas feedstock for the photocatalytic reduction reaction. As a result, largely enhanced yields of acids are obtained. For instance, the amount of HCOOH evolved from the CO reduction is 24 times higher than that from the CO₂ reduction on the Ni(NPA/TEG) catalyst (Fig. 5A), and the amount of CH₃COOH produced from the CO reduction reaction for the Ag/Rh-decorated Ni(NPA/TEG) catalyst exhibits a sixfold increase over the results from the CO₂ reduction reaction (Fig. 5B and fig. S16). Besides the increase in acid production, another important C₂ product, that is, ethanol [with a concentration of 270.6 μM for Ni(NPA/TEG)-Rh and 262.2 μM for Ni(NPA/TEG)-Ag], has also emerged from the 6-hour photocatalytic CO reduction reaction.

### Possible mechanisms for the photocatalytic reduction reactions of CO₂ or CO

On the basis of the abovementioned results, we propose the following mechanism for the photocatalytic CO₂ reduction reactions on the spongy Ni(TPA/TEG) catalyst (Fig. 6). Upon visible light irradiation, the photosensitizer [Ru(bpy)₃]²⁺ is excited and then reductively quenched by the TEOA sacrificial electron donor (18, 30, 43), which gives rise to the reduced species of [Ru(bpy)₃]³⁻ (Fig. 6A). Subsequently, the reduced species of [Ru(bpy)₃]³⁻ could transfer an electron to the spongy Ni(TPA/TEG) catalyst, which then participates in reducing the CO₂ molecules fixed on the catalyst (Fig. 6B). In the 2-hour yield tests of CO production in the solution with different amounts of Ni(TPA/TEG), we have found that the CO production rate decreases with the increase in the amount of catalyst (fig. S9), indicating that the electron transfer from [Ru(bpy)₃]³⁻ to the catalyst could be a rate-determining step for the CO₂ reduction reaction, where a diffusion-limited event may have occurred in this heterogeneous catalytic system (44).

For the selective generation of CO from CO₂, we propose the formation of a basic intermediate (45) [that is, CO₂ radical anion (CO₂⁻)] in the initial reaction step (CO₂ + e⁻→CO₂⁻), which acts as a Brønsted base and reacts easily with H₂O to form CO (CO₂⁻ + H₂O + e⁻→CO + 2OH⁻). On the other hand, in our proton-deficient photocatalytic reaction medium (pH 8), water (the proton donor) could provide another nonbasic intermediate, H⁺(H₂O + e⁻→H⁺ + OH⁻), before the H₂ evolution (H₂O + H⁺ + e⁻→H₂ + OH⁻) (46). For the spongy Ni(TPA/TEG) catalyst, it is likely that the Ni-TPA coordination units are unfavorable for the binding of H⁺ on the active sites, limiting the proton transfer and the formation of H₂. In addition, the flexible Ni-TEG units, with superior structural resistance to the aqueous environment (23), enable the disordered spongy network construction, where the open and defective structure provides more accessible Ni²⁺ active sites to capture and stabilize the CO₂⁻ intermediates, leading to the efficient production of CO.

Subsequently, the evolved CO can be further reduced to liquid fuels through proton-coupled multielectron reaction processes (Fig. 6C and fig. S17). Figure 6D shows the proposed conversion pathways leading to the formation of HCOOH, CH₃COOH, and CH₃CH₂OH via proton-coupled one-, four-, and eight-electron steps, respectively, in the electrolyte with a pH value of ~8. For the formation of HCOOH, we propose the following reactions:

\[
H_2O + e^- \rightarrow H^+ + OH^- \tag{1}
\]

\[
*CO + H^+ \rightarrow *CHO \tag{2}
\]

\[
*CHO + OH^- \rightarrow HCOOH + e^- \tag{3}
\]

where *CO is quickly protonated to *CHO and then hydroxylated to HCOOH. In the pathway of CH₂COOH formation, *CO is continuously hydrated to *CHO→*CHOH→*CH₂OH→*CH₂OH, which bonds with the adsorbed *CO to form CH₂COOH. As to the formation of CH₃CH₂OH, the dehydroxylation of the as-formed *CHO could be a critical rate-limiting step to produce *C that can be further protonated to *CH→*CH₂→*CH₃ (47), and the C-C coupling between *CH₃ and multiprotonated *CO (that is, CH₂OH) could lead to the formation of CH₃CH₂OH (48).

To further tune the proposed reactions, we performed the CO reduction reaction on the Ni(TPA/TEG)-Ag in the electrolyte with a pH value of ~13 (fig. S18). We found that CH₃OH (184.09 μM), CH₃CH₂OH (149.39 μM), HCOOH (438.98 μM), and CH₃COOH (276.99 μM) are produced at pH 13 in 6 hours, which is distinct from the liquid products generated at pH 8, that is, CH₃OH (0 μM), CH₃CH₂OH...
(262.19 μM), HCOOH (263.58 μM), and CH₃COOH (1178.04 μM). At pH 13, the hydroxyl ions (OH⁻) are commonly available for the hydroxylation of °CHO, which favors the formation of HCOOH (fig. S18). On the contrary, the enhanced hydroxylation of °CHO may suppress the kinetics of the multiprotonation of °CO and the dehydroxylation of °CHO (fig. 6D), resulting in a lower amount of CH₃COOH and CH₃CH₂OH at pH 13. The appearance of CH₃OH may suggest a weak C–C coupling between °CH₃OH and °CO at pH 13 (leading to CH₃COOH at pH 8), which should be considered for the future CO₂/CO reduction catalyst design (49).

In summary, we have demonstrated a photochemical strategy for the design of novel nanostructured metal-organic materials, where the rigid TPA and soft TEG molecules are successfully cross-linked together with the Ni²⁺ centers. A spongy Ni(TPA/TEG) hybrid structure with a considerably high concentration of defects has been achieved. We found that the Ni(TPA/TEG) catalyst is remarkably active for CO production (with a production rate of 15,866 μmol hour⁻¹ g⁻¹) from the heterogeneous photocatalytic CO₂ reduction reaction, during which no other measurable competing gases such as H₂ or CH₄ are generated, thus giving a near 100% CO selectivity over other gases. When the spongy Ni-organic catalyst is enriched with Rh or Ag nanocrystals, formic acid and acetic acid can be produced selectively from the photocatalytic CO₂ reduction reactions. We propose to use the spongy Ni(TPA/TEG) catalyst in a “tandem catalyst” system to convert CO₂ into high-value liquid fuels using visible light, where the selectively CO₂-turned CO can be directly used as an intermediate gas reactant for the generation of liquid fuels, such as ethanol and acetic acid (fig. S19). More advanced metal-organic heterogeneous photocatalysts with improved CO₂ fixation and light-harvesting capabilities are expected to be fabricated using the photochemical strategy for efficient solar-to-fuels conversion.

**MATERIALS AND METHODS**

**Laser-chemical synthesis of metal-organic photocatalysts**

All chemicals including nickel nitrate (99.99%), cobalt nitrate (99%), copper nitrate trihydrate (99%), ethanol (99.5%), DMF, TPA, and TEG were purchased from Sigma-Aldrich and used as received. The deionized water was produced by the Milli-Q Integral water purification system. A Continuum Surelite III nanosecond pulsed laser was used as a power source, and the following were the typical parameters of operation: wavelength, 1064 nm; frequency, 10 Hz; pulse width, 7 to 8 ns; beam diameter, 0.9 cm; and 700 mJ per pulse. TEG solutions (1 ml) of 1.5 M transition metal nitrates were added into 5 ml of DMF solution of 0.5 M TPA; the mixed solutions were stirred for 30 min before laser irradiation. For the syntheses of NiCo-, NiCoCu-organic frameworks, molar ratios of Ni/Co (1:1) and Ni/Co/Cu (1:1:1) were used, respectively. Typically, 3-hour laser irradiation was required for a 6-ml mixed precursor solution to complete the reaction. For the
syntheses by heating method, the same precursor solutions in glass vials were heated at 110°C in an oven for 48 hours. Precipitates produced after laser irradiations or heating were rinsed with acetone/ethanol, centrifuged at 9000 rpm for three times, and then dried in air at 60°C to obtain powders.

**Photocatalysis measurement of CO₂ reduction reactions**

The visible light–driven photocatalytic CO₂ reduction was conducted in a closed gas circulation and evacuation system fitted with a top window Pyrex cell. A circulating cooling water system was used to maintain the reactor at around 20°C. In a typical reaction, 3 mg of catalyst (for each test), 2.5 mmol of Ru(bpy)₃Cl₂·6H₂O, and 2 ml of TEOA were added to 10 ml of acetonitrile/H₂O solvent mixture (CH₃CN/H₂O = 8:2). The pH value of the solution was ~8, which was tuned to 13 by 1 M NaOH aqueous solution for control experiment. The light source was a 300-W Xe lamp with a long-pass cutoff filter (λ > 420 nm). Before light irradiation, the reaction system was evacuated and refilled by high-purity CO₂ (99.995%; SOXAL) several times to remove air inside and finally filled with CO₂ gas to reach a pressure of 400 torr. The evolved gas was detected by online GC (Agilent 7890A) equipped with a thermal conductivity detector and a flame ionization detector (FID) at different times of the photoreaction. To evaluate catalyst reusability, 15 mg of the catalyst was applied for photocatalysis and recycled by centrifugation after a 2-hour reaction, and then mixed with 8 ml of CH₂CN, 2 ml of H₂O, 2 ml of TEOA, and 18 mg of Ru(bpy)₃Cl₂·6H₂O for the next run. The solution products in the liquid phase were analyzed separately at the end of the photoreaction. Alcohols in the liquid phase were analyzed by Agilent 7890A GC with an FID, a DB-WAX column, and helium as the carrier gas. Carboxylic acids in the liquid phase were analyzed using an Agilent 1260 high-performance liquid chromatograph (HPLC) with a PL Hi-Plex H column and variable wavelength detector (VWD) (210 nm). [Ru(bpy)₃]Cl₂·6H₂O (99.95%) and TEOA (≥99.0%) were obtained from Sigma-Aldrich. The acetonitrile (HPLC-grade) was obtained from Merck. The ¹³C CO₂ isotope tracer experiment was performed under similar photocatalytic reaction conditions. The reactor containing 3 mg of the catalyst, 2.5 mmol of Ru(bpy)₃Cl₂·6H₂O, 2 ml of TEOA, and 10 ml of the CH₂CN-H₂O mixture solution (CH₂CN/H₂O = 8:2) was first evacuated to ensure air removal and then refilled with ¹³C CO₂ gas (99 atomic % ¹³C; Aldrich) to reach a pressure of 400 torr. After 4 hours of irradiation, 200 µl of gas products was withdrawn using a gas-tight syringe and then injected into a GC-MS (Agilent, GC Model 6890N/MS Model 5973) with a molecular sieve 5 Å column for analysis. The carboxylic acid analysis was carried out separately by using a Thermo Finnigan LCQ MS system.

**SUPPLEMENTARY MATERIALS**

Supplementary material for this article is available at http://advances.sciencemag.org/cgi/content/full/3/7/e1700921/DC1

Materials characterizations

Three-dimensional tomographic reconstruction of the spongy Ni(TPA/TEG) network Comparison of the samples synthesized by laser-chemical route and traditional heating method Control experiments to confirm the origin of the evolved CO

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**REFERENCES AND NOTES**

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