

# Visible and NIR Light Assistance of the N<sub>2</sub> Reduction to NH<sub>3</sub> Catalyzed by Cs-promoted Ru Nanoparticles Supported on Strontium Titanate

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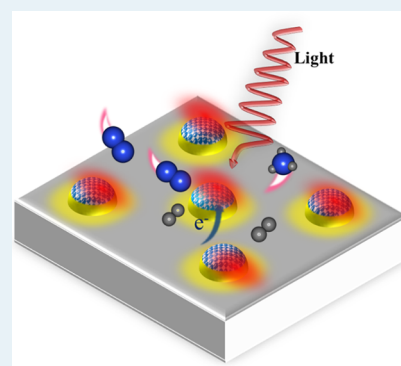


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Supporting Information

**ABSTRACT:** NH<sub>3</sub> production accounts for more than 1% of the total CO<sub>2</sub> emissions and is considered one of the most energy-intensive industrial processes currently ( $T > 400$  °C and  $P > 80$  bars). The development of atmospheric-pressure N<sub>2</sub> fixation to NH<sub>3</sub> under mild conditions is attracting much attention, especially using additional renewable energy sources. Herein, efficient photothermal NH<sub>3</sub> evolution in continuous flow upon visible and NIR light irradiation at near 1 Sun power using Cs-decorated strontium titanate-supported Ru nanoparticles is reported. Notably, for the optimal photocatalytic composition, a constant NH<sub>3</sub> rate near  $3500 \mu\text{mol}_{\text{NH}_3} \text{g}_{\text{catalyst}}^{-1} \text{h}^{-1}$  was achieved for 120 h reactions, being among the highest values reported at atmospheric pressure under 1 Sun irradiation.



**KEYWORDS:** photothermal, ammonia synthesis, Cs promoter, titanium perovskite, Ru nanoparticles

## 1. INTRODUCTION

The most relevant chemical industrial processes (*i.e.*, Bessemer, Gattermann, Haber–Bosch, Fischer–Tropsch, Solvay, *etc.*) are characterized by the enormous energy input required to achieve cost-effective and competitive product formation rates. This is consequence of the unfavorable thermodynamics and sluggish kinetics characteristic of these chemical reactions, making necessary the use of very large pressures and temperatures to achieve adequate production rates. In the context of finding alternative processes to the conventional thermocatalytic reaction, the conversion of inexhaustible and clean sunlight into chemicals and fuels by means of a (photo)catalyst, in the so-called photothermal process, is being increasingly considered an appealing approach to alleviate the high required energy input. The use of solar light should contribute to mitigate the environmental impact caused by the massive consumption of fossil fuels to obtain these enormous amounts of energy.

It is increasingly recognized that the yield of a large variety of chemical reactions of special interest for industry, such as hydrogenations,<sup>1–3</sup> oxidations,<sup>4,5</sup> couplings,<sup>6,7</sup> rearrangements,<sup>8</sup> among others,<sup>9,10</sup> can be notably enhanced by the assistance of light. In the conventional photothermal mechanism, light absorption at catalytically active metal nanoparticles (NPs) causes an increase of the local temperature at the nanoscale that is not measurable by conventional macroscopic methods.<sup>11</sup> This localized temperature increase can result in a much enhanced catalytic activity of the metal

NP.<sup>12,13</sup> This photothermal mechanism causes the selective heating of the absorber, particularly when the thermal conductivity of the support is low, because heat dissipation becomes slower.<sup>14</sup> When the absorber is a catalytically active site, then, remarkable increases in activity, much higher than those expected according to the macroscopic temperature, can be observed, simply because of the selective conversion of the photon into heat at the proper site. Moreover, in certain cases, it has been proved that chemical reactions can be driven simultaneously by photothermal (selective heating at the NP, not detectable by macroscopic thermocouples) and photocatalytic (photoinduced electron/hole separation) mechanisms.<sup>15,16</sup>

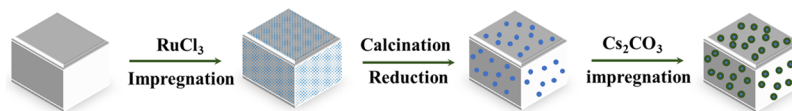
One clear example of photoassisted hydrogenation of an inert molecule is the Sabatier hydrogenation of CO<sub>2</sub> into CH<sub>4</sub>.<sup>17,18</sup> The Sabatier reaction is well known to take place at high temperatures above 500 °C in order to achieve production rates convenient for the industry. For this reaction, it has been demonstrated that illumination, even with solar light, increases the CH<sub>4</sub> formation rate at values required at

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Scheme 1. Illustration of the Synthesis of the  $\text{Cs}_y\text{Ru}_x@ST$  Catalyst by the Incipient Wetness Impregnation Method

much higher temperatures and pressures for the pure thermocatalytic process.

Going a step beyond the relatively large number of photothermal and photocatalytic  $\text{CO}_2$  hydrogenation reports,<sup>17–19</sup> herein, we present an optimized photocatalyst for  $\text{N}_2$  reduction to  $\text{NH}_3$  operating in continuous flow under about 1 Sun illumination. This light-assisted  $\text{N}_2$  fixation has been barely studied so far. Previous studies of photocatalytic  $\text{N}_2$  reduction have focused on semiconductor metal oxides, especially  $\text{TiO}_2$ .<sup>20</sup> However,  $\text{TiO}_2$  presents a 3.2 eV band gap, limiting light harvesting to the UV range, which accounts for less than 5% of the solar light spectrum. Thus,  $\text{TiO}_2$  photocatalysts are not suitable for solar light assistance. Besides metal oxides, Cd-containing dichalcogenides, basically in the form of sulfides, have attracted considerable attention due to their visible light photoresponse. However, the high metal toxicity and poor stability makes Cd and metal chalcogenides far from any practical use. In other study, illumination with up to 47 Suns power, without external heating, of  $\text{Cs}^+$ -promoted Ru NPs supported in MgO resulted in a temperature gradient in a thick catalyst bed with a hotter external surface and cooler interior that was found to be a thermodynamic pump favoring  $\text{NH}_3$  formation.<sup>21</sup>

In the present article, we demonstrate that visible and NIR light can assist the  $\text{N}_2$  hydrogenation at atmospheric pressure and temperatures below 400 °C under near 1 Sun ( $1080 \text{ W m}^{-2}$ ) light intensity using as photocatalyst Cs-promoted ruthenium NPs supported on strontium titanate ( $\text{Cs}_y\text{Ru}_x@ST$ ,  $x$  referring to Ru loading and  $y$  to the Cs/Ru atomic ratio of these elements and ST corresponding to strontium titanate). This photoassisted process will have the advantage of atmospheric pressure and lower macroscopic temperatures than the current industrial process, while using *ca.* 95% of the solar light spectrum. Upon visible and NIR light harvesting, the photocatalyst with optimal composition promotes  $\text{NH}_3$  formation under continuous flow at rates about 68% higher than those obtained in the thermal process under dark conditions and reaching values that are among the highest reported so far for any photocatalytic  $\text{N}_2$  fixation. Moreover, we have studied in detail, using *in situ* spectroscopic techniques, the  $\text{NH}_3$  formation mechanism as well as the role of Cs as the reaction promoter. The Cs-promoted Ru NPs supported on the ST photocatalyst has also demonstrated to be very stable under these reaction conditions for 120 h continuous operation. We believe that this work sets a precedent for the photothermal  $\text{N}_2$  hydrogenation reaction in continuous operation, which is considered a prerequisite for any industrial purpose.

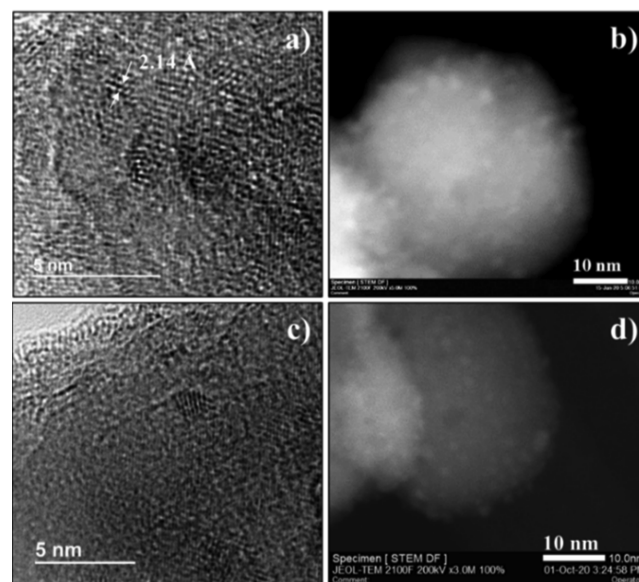
## 2. RESULTS AND DISCUSSION

### 2.1. Catalyst Preparation and Characterization.

Ru NPs supported on strontium titanate ( $\text{Ru}_x@ST$ ) containing different Ru loadings (1, 2.5, and 5 wt %) were first prepared to screen out the optimal amount of Ru loading for the  $\text{N}_2$  hydrogenation reaction, and the synthetic procedure is described in detail in the Experimental Section in Supporting

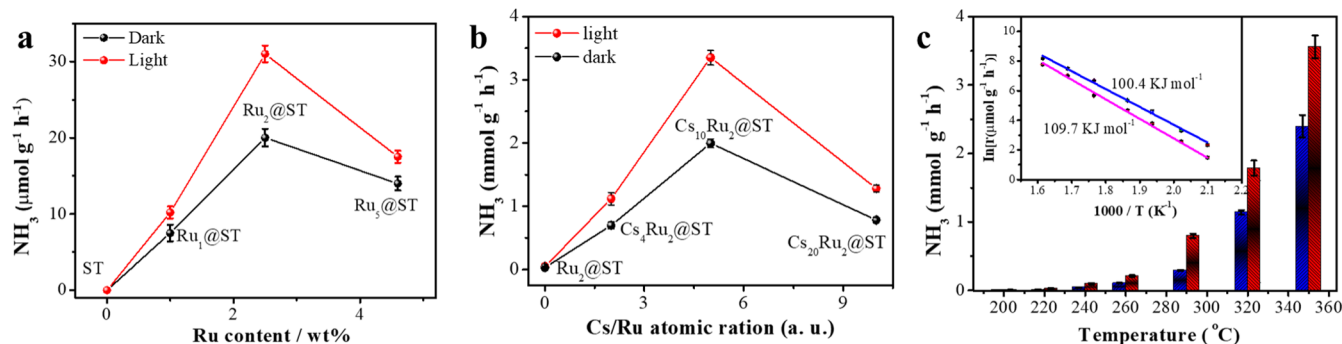
Information and illustrated in Scheme 1. Commercial ST particles were impregnated with  $\text{RuCl}_3$  solutions at different concentrations. After drying, the  $\text{Ru}^{3+}/ST$  solids were submitted to calcination at 250 °C and, subsequently, reduced at 350 °C in a  $\text{H}_2$  atmosphere, obtaining the  $\text{Ru}_x@ST$  catalysts (being  $x$  the  $\text{Ru}^{3+}$  wt %). The amount of Ru loaded in the final  $\text{Ru}_x@ST$  catalysts was measured by X-ray fluorescence spectroscopy (XRF) and the Ru content of  $\text{Ru}_x@ST$  catalysts ( $x = 1, 2, \text{ and } 5$ ) was determined to be 0.94, 2.48, and 4.6 wt %, respectively, based on a previous calibration by XRF using  $\text{RuO}_2/ST$  mixtures with different Ru known concentrations.

The powder X-ray diffraction (PXRD) patterns obtained from  $\text{Ru}_x@ST$  samples correspond solely to the STO support (see Figure S1 in Supporting Information), and the absence of diffraction peaks from Ru species (neither metallic Ru nor  $\text{RuO}_2$ ) could be justified by its small particle size and homogeneous distribution on the ST surface. The high dispersity of Ru NPs on the ST surface was further confirmed by high-resolution transmission electron microscopy (HR-TEM) and annular dark-field scanning transmission electron microscopy (ADF-STEM). Figure 1a shows the HR-TEM



**Figure 1.** (a) Representative HR-TEM images and (b) ADF-STEM images obtained from the  $\text{Ru}_2@ST$  catalyst. (c) Representative HR-TEM images and (d) ADF-STEM images of the  $\text{Cs}_{10}\text{Ru}_2@ST$  catalyst.

image obtained from  $\text{Ru}_2@ST$ , where highly crystalline NPs, homogeneously deposited on the support surface, can be discerned. A planar distance of 2.14 Å in the crystalline NPs was determined, which matches well with the (002) lattice plane of the metallic Ru phase (PDF #88-1734). ADF-STEM images in Figures 1b and S2 (see Supporting Information) allowed us to measure the dimensions of the Ru NPs, and the average size of  $\text{Ru}_1@ST$ ,  $\text{Ru}_2@ST$ , and  $\text{Ru}_5@ST$  was  $0.9 \pm 0.3$ ,  $1.4 \pm 0.4$ , and  $1.8 \pm 0.5$  nm, respectively.



**Figure 2.** (a) Dependence of the NH<sub>3</sub> production rate on the amount of Ru loading both under light irradiation (red spheres) and dark conditions (black spheres), (b) NH<sub>3</sub> production rate obtained from Ru<sub>2.5</sub>@ST with different Cs loadings in both light (red spheres) and dark (black spheres) reaction conditions, (c) NH<sub>3</sub> production rate obtained from Cs<sub>10</sub>Ru<sub>2</sub>@ST at different reaction temperatures, in both light (red) and dark (blue) conditions. Inset: Arrhenius plot based on the ammonia production rate at different temperatures and the linear fitting in both light (blue) and dark (pink) conditions. The corresponding activation energies are also indicated in the figure. Reaction conditions: 50 mg catalyst, 10 mL min<sup>-1</sup> N<sub>2</sub> + 30 mL min<sup>-1</sup> H<sub>2</sub>, 350 °C, and 0.1 MPa (unless otherwise specified).

## 2.2. Photothermal Catalytic Ammonia Production

**Tests.** Photothermal N<sub>2</sub> fixation reaction was conducted using the obtained Ru<sub>x</sub>@ST samples, targeting to screen out the optimal Ru loading. The reaction was performed using a customized glass flow reactor at 350 °C upon external heating, and light enhancement was evaluated by shifting from dark to light condition (1080 W/m<sup>2</sup>) (see Experimental Section and Scheme S1 for further details of the reaction setup). It is important to note that the thickness of the photocatalytic film (about 1 mm) and the collimated light irradiation should avoid penetration gradients in the catalyst bed (circular 1 cm diameter). In addition, the macroscopic temperature was measured with a thermocouple in contact with the photocatalyst. As shown in Figure 2a, bare ST support exhibited no activity, while 7.02 μmol g<sup>-1</sup> h<sup>-1</sup> NH<sub>3</sub> production was achieved using the Ru<sub>1</sub>@ST catalyst in dark conditions. The production rate further increased to 20.14 μmol g<sup>-1</sup> h<sup>-1</sup> with 2.5 wt % Ru loading (Ru<sub>2</sub>@STO) and then decreased to 14.21 μmol g<sup>-1</sup> h<sup>-1</sup> when Ru loading reaches 4.6 wt % (Ru<sub>5</sub>@ST). Upon light irradiation, the NH<sub>3</sub> production rates followed the same trend as that of the dark condition; however, 10.19, 30.90, and 20.35 μmol g<sup>-1</sup> h<sup>-1</sup> for Ru<sub>1</sub>@ST, Ru<sub>2</sub>@ST, and Ru<sub>5</sub>@ST, respectively, were obtained. The optimal Ru loading can be explained as a result of two contradictory factors: (a) the increase of Ru NPs as the active sites must promote the N<sub>2</sub> hydrogenation,<sup>22</sup> while (b) further increase in the Ru loading results in larger average particle size, and as a consequence, a decrease in active site surface per mass unit. It must be noticed that the light-enhanced activities are over 50%, which, to the best of our knowledge, is the first reported case of photoassisted N<sub>2</sub> hydrogenation to NH<sub>3</sub>.

To confirm N<sub>2</sub> as the main source of NH<sub>3</sub>, control experiments using <sup>15</sup>N<sub>2</sub>-labeled gas were carried out (see Experimental Section in Supporting Information for further details), and the resultant ammonia products were determined by <sup>1</sup>H NMR. For comparison purposes, a reaction using <sup>14</sup>N<sub>2</sub> was also carried out and the obtained products were analyzed by <sup>1</sup>H NMR. As can be seen from Figure S3, the <sup>1</sup>H NMR spectrum conclusively confirms N<sub>2</sub> as the source of NH<sub>3</sub>.<sup>23</sup>

It is worth noticing that the catalytic activity of Ru<sub>2</sub>@ST is below the typical values in the state of the art under similar reaction conditions (see Table S1 in Supporting Information), albeit a 50% of enhancement was achieved. For that reason, further modification of Ru<sub>2</sub>@ST has been carried out. It has

been previously reported that the incorporation of alkali or alkaline-earth metal as promoters in some catalysts can remarkably increase the catalytic activity or selectivity in some reactions such as CO<sub>2</sub> hydrogenation, alkanes dehydrogenation, NO reduction, and N<sub>2</sub> hydrogenation, among others.<sup>24–27</sup> Hence, Cs<sub>2</sub>CO<sub>3</sub> was introduced onto the Ru<sub>2</sub>@ST surface by wet impregnation (see Scheme 1 and Experimental Section in Supporting Information for detailed information). Three Cs<sub>x</sub>Ru<sub>2</sub>@ST catalysts with Cs/Ru equal to 2, 5, and 10 were prepared and labeled as Cs<sub>2</sub>Ru<sub>2</sub>@ST, Cs<sub>5</sub>Ru<sub>2</sub>@ST, and Cs<sub>10</sub>Ru<sub>2</sub>@ST, respectively. Figure 1c,d shows the representative HR-TEM and ADF-STEM images of the resultant Cs<sub>10</sub>Ru<sub>2</sub>@ST, which exhibit no particle size or morphology change compared with the Ru<sub>2</sub>@ST sample. PXRD of Cs<sub>x</sub>Ru<sub>2</sub>@ST catalysts exhibited only diffraction patterns from ST (Figure S2), which could be explained by the high dispersity of Cs species that can migrate to the vicinity of Ru NPs under a H<sub>2</sub> atmosphere at the reaction temperature, as reported before.<sup>28</sup> The migration of Cs species to Ru NPs was lately confirmed by EDX analysis, which revealed the co-presence of Cs and Ru in the high contrast region of the spectrum 1 (Figure S4b in Supporting Information), while no Cs was detected far from the Ru NPs in the low contrast region of spectrum 2. In comparison, no obvious image contrast differences can be observed in samples before the activation (Figure S4a). These differences in contrast are compatible with the coverage of Cs on the whole surface of the support and hence further prove the migration of Cs during the activation treatment. Further evidence of Cs migration to the Ru surroundings has been obtained from EDS analysis of randomly selected regions and elemental mapping of representative STEM images of the Cs@ST sample lacking Ru for which a homogeneous Cs distribution was observed (Figure S4c,d–h, respectively). Preferential Cs deposition near Ru NPs has been previously justified as derived from the higher adsorption energy of Cs for Ru rather than basic support and the same can apply in the present case.<sup>31</sup>

The results of photothermal N<sub>2</sub> hydrogenation demonstrated a dramatic increase in activity for NH<sub>3</sub> production in the presence of the Cs promoter, being 3345 μmol g<sup>-1</sup> h<sup>-1</sup> under 1080 W/m<sup>2</sup> light irradiation with the optimal Cs loading (Cs<sub>10</sub>Ru<sub>2</sub>@ST), 100 times higher compared to Ru<sub>2</sub>@ST (Figure 2b). In addition, compared to dark conditions (1989 μmol g<sup>-1</sup> h<sup>-1</sup>), light irradiation enhanced the NH<sub>3</sub> production

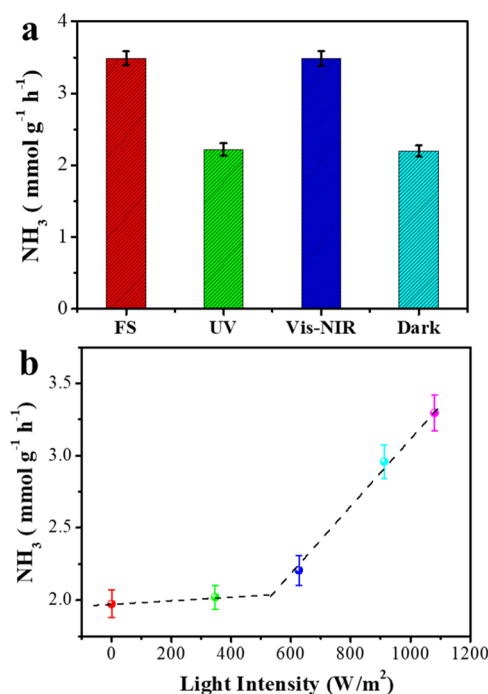
rate by over 68%. To the best of our knowledge, this activity is among the highest values reported so far (see Table S1 in Supporting Information). The dependence of catalytic activity with the Cs loaded follows a similar trend in both light and dark conditions and could be explained also by the simultaneous concurrence of two opposite effects: (1) the presence of Cs species as the promoter can facilitate the nitrogen activation and dissociation (*via infra*), therefore resulting in activity enhancement, while (2) excessive Cs amounts lying next to the Ru NPs can cover partially or completely the Ru NPs, thus decreasing the amount of exposed active sites.<sup>29</sup> Therefore, the optimal Cs loading must be a compromise between these two factors.

The partial coverage of Cs on the Ru NP surface was investigated by CO chemisorption measurement, and the gas sorption volume versus pressure profiles of Cs<sub>10</sub>Ru<sub>2</sub>@ST and Ru<sub>2</sub>@ST samples are presented in Figure S5. The average crystal size of Ru<sub>2</sub>@ST was 1.464 nm, which is in good agreement with the value obtained from the ADF-STEM image. The calculated Ru surface area was 6.663 m<sup>2</sup>/g, with the metal dispersion of 91.13%. However, the average crystal size obtained in the Cs<sub>10</sub>Ru<sub>2</sub>@ST sample was 3.93 nm, and the surface area has decreased to 2.843 m<sup>2</sup>/g. As confirmed by HR-TEM and AD-STEM images (Figure 1), the Ru particle size remains unchanged after the incorporation of Cs, and thus, the only explanation for the active surface area decreases and the calculated particle size increase is the partial coverage of Cs species on Ru NPs, which is also in agreement with the results obtained from the EDX analysis (Figure S4), while a homogeneous Cs distribution was observed in the absence of Ru after activation (Figure S4c). The partial coverage of Cs on Ru sites can be justified by the fact that the partial reduction of Cs<sup>+</sup> (*vide infra*) occurred on the Ru sites due to the high concentration of adsorbed H<sub>2</sub> on its surface.<sup>30–32</sup> In addition, the Sr component contributes to the high basicity of ST support and thus also favors the migration of Cs species to Ru NPs at high temperature.<sup>33</sup>

**2.3. Photothermal Effects.** To investigate the underlying mechanism of the light enhancement, the reaction was conducted using Cs<sub>10</sub>Ru<sub>2</sub>@ST at temperatures in the range of 200 to 350 °C, either in dark or in light irradiation conditions. As shown in Figure 2c, the NH<sub>3</sub> production activity exhibited an exponential increase with elevating reaction temperature, both in dark and in light conditions, and over 50% of light enhancement was achieved in all the tested temperature ranges. The activation energies under light and dark conditions were 100.4 and 109.7 kJ/mol, respectively, calculated based on Arrhenius plots (inset in Figure 2c) and equation (eq 1). A similar activation energy value in light and dark conditions indicates the same rate determine step in both cases, suggesting that light-assisted and thermal reactions share the same limiting step in the reaction mechanism.

$$K(T) = Ae^{-E_a/RT} \quad (1)$$

The influence of the irradiation wavelength and the light intensity on the NH<sub>3</sub> production was also investigated. Cs<sub>10</sub>Ru<sub>2</sub>@ST was selectively irradiated with full spectrum (FS), visible, and near infrared (vis–NIR) or ultraviolet (UV) light. As can be observed in Figure 3a, light enhancement originates from the vis–NIR range, while the NH<sub>3</sub> production rate from UV irradiation can be considered negligible. Furthermore, 515, 610, 695, and 830 nm cut-off filters were applied to evaluate the spectral response. As shown in Figure



**Figure 3.** (a) NH<sub>3</sub> production rate with FS light, UV light, and vis–NIR light irradiation, as well as in dark condition (dark). Light source, 300 W Xenon lamp with FS 1080 W/m<sup>2</sup>. (b) Dependence of the NH<sub>3</sub> production rate and the irradiation light intensity. Light source, 300 W Xenon lamp with FS 1080 W/m<sup>2</sup>. 50 mg Cs<sub>10</sub>Ru<sub>2</sub>@ST, 10 mL min<sup>-1</sup> N<sub>2</sub> + 30 mL min<sup>-1</sup> H<sub>2</sub>, 350 °C and 0.1 MPa.

S6a, the NH<sub>3</sub> production rates using the 515 and 610 nm filters were almost identical to those of FS irradiation, indicating that light contribution of wavelengths below 610 nm is negligible. On the contrary, 70% of light enhancement has been observed using an 830 nm cut-off filter, suggesting that the light-induced effects mainly originate from the NIR.

Overall, the spectral response of this photothermal NH<sub>3</sub> system makes it promising for solar light irradiation. The influence of light intensity on the activity was also studied with a 300 W Xe lamp by passing the incident light through neutral density filters before reaching the catalyst. The temperature variations with light intensity in the catalyst surface have been measured (Figure S7), obtaining a maximum temperature increase of 12 °C at the maximum light intensity. Consequently, the external heating temperature controller has been corrected accordingly in order to have constant 350 °C at the different light intensities. Figure 3b shows that the activity increased linearly with the light intensity after reaching 500 W/m<sup>2</sup>, while in the range of 0–500 W/m<sup>2</sup>, negligible activity enhancement was achieved, which indicates that a minimum intensity was required to have a significant activity increase. A similar trend was also observed using the monochromatic 980 nm laser as the light source. In agreement with Figure 2c, this minimum light intensity to form NH<sub>3</sub> is probably related to the requirement of a minimum local temperature at the Ru NPs to promote N<sub>2</sub> activation. In addition, as can be seen in Figure S6b, remarkable activity enhancement occurred after the laser power reached 0.5 W, and noticeably, a plateau was reached beyond 1.5 W, which indicates a saturation of the photogenerated carriers. It must be noticed that this conversion is far below the theoretical equilibrium, and even at 450 °C, the decomposition of the

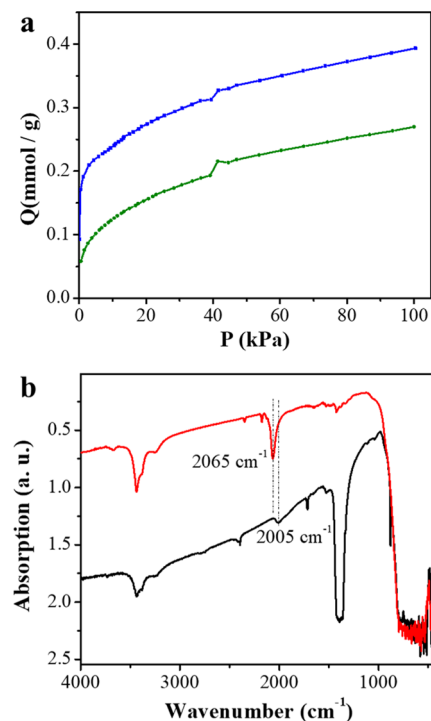
ammonia products can be neglected (Figure S8). It is well-known that for pure thermal catalysis, the conversion rates increase with temperature, which can be modulated by the incident light intensity.<sup>34</sup> In the present case, pure thermal effects induced by simple light-induced heating can be ruled out, as derived from the saturation effect observed in this experiment, and thus, synergistic thermal and photo-induced hot carrier mechanisms are proposed as responsible for the NH<sub>3</sub> enhancement upon light irradiation, as reported elsewhere.<sup>35</sup>

Diffuse reflectance spectra from Ru<sub>2</sub>@ST and Cs<sub>10</sub>Ru<sub>2</sub>@ST catalysts are presented in Figure S9a in the Supporting Information, showing very similar features. In this, a predominant band in the UV light range ( $\lambda < 350$  nm), originated from light absorption of ST support, can be observed. Additionally, other absorption bands, starting from 550 nm (inset in Figure S9a), and extended to the NIR range (Figure S9b), can be attributed to light absorption by Ru metal.<sup>18</sup> Noticeably, the vis–NIR absorption results are coincident with the rate–wavelength dependence observed in Figure S6a, which strongly confirms the light-enhanced NH<sub>3</sub> production rate originated from the interaction of supported Ru NPs and incident light.

Because 70% of the light enhancement is derived from the NIR range, while the UV light, where the ST has strong response, does not contribute to the NH<sub>3</sub> production, it is then of interest to study if ST only acts as an inert support for the Ru NPs dispersity. Hence, different supports (TiO<sub>2</sub>, P25, aluminate, and aluminosilicate) loaded with the same amount of Ru and Cs were used as photocatalysts in the NH<sub>3</sub> production. As observed in Figure S10, the activity of Cs<sub>10</sub>Ru<sub>2</sub>@ST is over two orders of magnitude compared with those using P25, aluminate, and aluminosilicate as the support, despite the lowest surface area it processes (surface area: 46, 71, 275, and 900 m<sup>2</sup> g<sup>-1</sup> for ST, P25, aluminate, and aluminosilicate, respectively). In addition, the light enhancement in the case of ST is the highest, being 65, 34, 20, and 31% for ST, P25, aluminate, and aluminosilicate, respectively. Therefore, it is clear that ST is not just an inert support, but can synergistically facilitate the N<sub>2</sub> reduction reaction. It has been reported that support with higher basicity favors the migration of alkali metals (promoter) to the vicinity of the metal active site during the activation process, and as a consequence, the strongly interaction of the promoter with the active site induces a high catalytic activity.<sup>28</sup> In the present case, the presence of Sr in the structure enables the ST process to have the highest basicity among those supports; thus it exhibited the highest activity even in dark conditions. Regarding the photothermal enhancement, it has been reported in previous works that supports that possess low thermal conductivity could avoid the heat dissipation from high temperature zones localized in the NPs to the surrounding through the support, and hence a considerable high photothermal effects can be achieved.<sup>14</sup> This mechanism can explain the high photoenhancement of the NH<sub>3</sub> rate in the case of ST, as its thermal conductivity is as low as 4.5 W m<sup>-1</sup> K<sup>-1</sup> at 350 °C; therefore, ST is an ideal support for the photothermal N<sub>2</sub> hydrogenation reaction.

**2.4. Promoter Effects.** Alkali and alkaline earth transition metals have been used as promoters in reactions, and the promoting effects have been attributed to basicity/acidity changes, structure regulation, and electronic modulation.<sup>29,36–42</sup> High basicity could promote NH<sub>3</sub> desorption, as

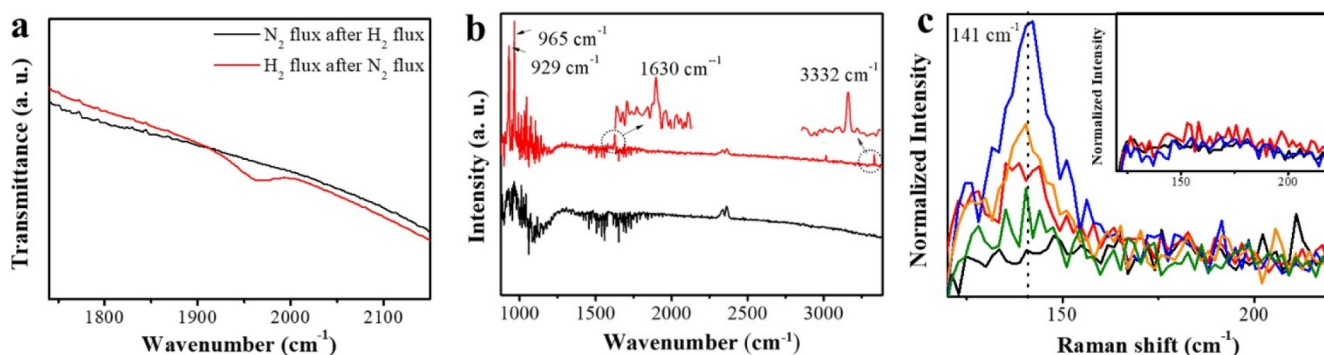
well as electron density migration from the Cs promoter to the Ru active centres. Hence, the basicity in Ru<sub>2</sub>@ST and Cs<sub>10</sub>Ru<sub>2</sub>@ST was evaluated by the measurement of isothermal CO<sub>2</sub> adsorption. As can be observed in Figure 4a, the CO<sub>2</sub>



**Figure 4.** (a) Isothermal CO<sub>2</sub> adsorption profiles of Cs<sub>10</sub>Ru<sub>2</sub>@ST (blue line) and Ru<sub>2</sub>@ST (green line) obtained at 273 K. (b) FT-IR spectrum of CO chemisorption obtained from Cs<sub>10</sub>Ru<sub>2</sub>@ST (black line) and Ru<sub>2</sub>@ST (red line).

adsorption capacity was twofold increased after the incorporation of the Cs promoter. Because no morphology or structure changes have been observed by XRD and ADF-SEM characterization (see Figures 1 and S2), the only explanation can be the increase in the basicity sites. Electron-donating effects from the promoter to the Ru active site were first observed by monitoring CO chemisorption on the catalysts by Fourier transformed infrared spectroscopy (FT-IR). As can be seen in Figure 4b, FT-IR spectra from the Ru<sub>2</sub>@ST catalyst exhibits a peak at 2065 cm<sup>-1</sup>, attributed to C=O vibration from linear adsorbed CO on the Ru surface. This vibration peak shows 55 cm<sup>-1</sup> shift toward lower wavenumbers in Cs<sub>10</sub>Ru<sub>2</sub>@ST, which is, as reported previously, due to more electron back-donating to the adsorbed CO molecule, and in other words, indicate higher electron density on the Ru NP surface.<sup>43,44</sup>

The electron-donating effect was confirmed by monitoring UV–vis absorption of tetracyanoethylene (TCNE) molecules in the presence of Ru<sub>2</sub>@ST and Cs<sub>10</sub>Ru<sub>2</sub>@ST catalyst. It is well-known that TCNE can accept electrons from the vicinity of electron-rich surfaces, forming e<sup>-</sup>/TCNE complexes that exhibit a characteristic UV–vis absorption peak at ~300 nm.<sup>45</sup> As shown in Figure S11a, the intensity of the absorption band centered at ~300 nm gradually increased with time upon the addition of the Ru<sub>2</sub>@ST catalyst. This result indicates that the TCNE molecule can accept electrons from the adjacent Ru active sites and form e<sup>-</sup>/TCNE complexes. Impressively, the intensity of the peak corresponding to the e<sup>-</sup>/TCNE complex



**Figure 5.** (a) *In situ* FT-IR spectrum of  $\text{Cs}_{10}\text{Ru}_2@\text{ST}$  after activation with  $\text{H}_2$  and purging with argon. Subsequently flush with  $\text{N}_2$  followed by  $\text{H}_2$  flux (red line) or  $\text{H}_2$  followed by  $\text{N}_2$  flux (black line). (b) *In situ* FT-IR spectrum in gas after each step in figure (a), that is, the gas phase spectrum of  $\text{N}_2$  after  $\text{H}_2$  flux (red line) and  $\text{H}_2$  after the  $\text{N}_2$  spectrum (black line). (c) *In situ* Raman spectrum of  $\text{Cs}_{10}\text{Ru}_2@\text{ST}$  after activation with  $\text{H}_2$  and purging with Ar (black line), followed by  $\text{N}_2$  flux after 10 min (red line) and 20 min (blue line). Afterward, it shifted to  $\text{H}_2$  flow for 10 min (orange line) and 20 min (green line). [Inset: *in situ* Raman spectrum of  $\text{Ru}_2@\text{ST}$  with the same test steps as that in figure (c); however no signal detected].

grew 3 times faster in the presence of the  $\text{Cs}_{10}\text{Ru}_2@\text{ST}$  catalyst compared to that with  $\text{Ru}_2@\text{ST}$ , reaching 3 at 11 min (Figure S11b), suggesting the higher electron density on the Cs-decorated catalysts.

X-ray photoemission spectroscopy (XPS) measurements were performed to further confirm the electron density transfer from the Cs species to the Ru active sites. Figure S12 presents the XPS spectrum acquired from  $\text{Ru}_2@\text{ST}$  and  $\text{Cs}_{10}\text{Ru}_2@\text{ST}$  catalysts. As can be seen in Figure S10a, the binding energy of Ru 3d is located in the same range with C 1s and Sr 3p. After deconvolution, clearly distinguished peaks can be seen at 280.53 and 284.73 eV corresponding to metallic Ru. In addition, the deconvolution from Ru 3p + Ti 2p spectra also confirms the reduced Ru species with binding energies of 461.36 and 483.56 eV, together with peaks at 458.46 and 464.16 eV corresponding to  $\text{Ti}^{4+}$  (Figure S12b). Ru 3d + C 1s + Sr 3p spectra of  $\text{Cs}_{10}\text{Ru}_2@\text{ST}$  clearly exhibit the presence of  $\text{CO}_3^{2-}$  species; besides, Ru 3d is 0.33 eV shifted to lower binding energy, with components at 280.20 and 284.40 eV (Figure S12c), which can be ascribed to higher electron density of Ru.<sup>28</sup> This lower binding energy shift was also observed in the Ru 3p spectrum, which shows a binding energy of 461.10 eV for the  $3p_{3/2}$  orbital. Therefore, the XPS results demonstrated a higher electron density of Ru active sites in the presence of Cs as a result of the electron back-donating effects from the Cs promoter. To further demonstrate the underlying reason why the Cs can donate electrons to Ru, XPS analysis of Cs 3d before (Figure S12g) and after (Figure S12e) the  $\text{H}_2$  activation was carried out. As can be observed, the Cs 3d of the sample shows a binding energy of 724.92 eV before activation, while this binding energy is shifted to 725.58 eV after the  $\text{H}_2$  activation. It must be noted that the Cs exhibited inverse chemical shifts to higher binding energy when the oxidation state decreases.<sup>28</sup> It is well-known that the binding energy of Cs metal is  $\sim 726.1$  eV, and hence, it can be concluded that the valence state of Cs species after the activation is between 0 and 1, namely,  $\text{Cs}^{\delta+}$  ( $0 < \delta < 1$ ), which is also in good agreement with the previous report.<sup>28</sup>

Overall, it has been demonstrated that the partially reduced Cs species can have a strong electron-donating ability, transferring electrons to the adjacent Ru sites. Accompanying a partial reduction,  $\text{Cs}_2\text{CO}_3$  species also underwent partial decomposition, as observed from O1s spectra from the  $\text{Cs}_{10}\text{Ru}_2@\text{ST}$  sample before and after the activation (Figure

S12f,h). As can be seen there, before the activation, the O 1s spectrum is composed of two peaks at 530 and 531.8 eV, which can be attributed to Ti–O and  $\text{CO}_3^{2-}$ .<sup>46,47</sup> After the activation, a new component at 532.3 eV (30%) related to –OH species can be observed.<sup>48</sup> The appearance of this new component after the activation indicates  $\text{Cs}_2\text{CO}_3$  partial decomposition to CsOH.

**2.5. Reaction Mechanism.**  $\text{N}_2$  hydrogenation to  $\text{NH}_3$  involves  $\text{N}_2$  activation and dissociation, as well as the  $\text{H}_2$  splitting to hydride. Therefore, the following control experiments were implemented to make out the key intermediates for the  $\text{NH}_3$  production. Specifically,  $\text{Cs}_{10}\text{Ru}_2@\text{ST}$  was Ar-purged in the reaction system for 15 min to remove all the  $\text{H}_2$  gas from the activation process. Then,  $\text{N}_2$  flux was introduced in the reaction chamber at 350 °C for 15 min to form any possible activated N species. Afterward, the reactor was Ar-purged for 15 min to remove the free  $\text{N}_2$  molecules, followed by shifting the gas flow to  $\text{H}_2$ . In this way, it is expected that  $\text{H}_2$  will react with the adsorbed N species, forming  $\text{NH}_3$ . In the end of this experiment, 7.1  $\mu\text{g}$  of  $\text{NH}_3$  was collected. However, by changing the gas sequence to react the adsorbed hydride with  $\text{N}_2$ , only 0.5  $\mu\text{g}$  of  $\text{NH}_3$  was obtained. These results indicate that the key step for the hydrogenation is the  $\text{N}_2$  adsorption and activation, instead of  $\text{H}_2$  adsorption or splitting.

Similar results were observed from *in situ* FT-IR characterization. For this study,  $\text{Cs}_{10}\text{Ru}_2@\text{ST}$  was first activated at 350 °C for 2 h *in situ* in the chamber, followed by introducing  $\text{H}_2$  and  $\text{N}_2$  (or  $\text{N}_2$  and  $\text{H}_2$ ) in sequence, and then the FT-IR spectrum was collected. It must be noted that Ar flow was applied for 15 min before introducing  $\text{N}_2$  or  $\text{H}_2$  gas to ensure that the gas phase molecules previously introduced had been totally removed. Figure 5a shows the representative spectrum obtained in this study, where a new peak was observed at 1962  $\text{cm}^{-1}$  in the red curve corresponding to the spectrum collected in  $\text{H}_2$  atmosphere after  $\text{N}_2$  flux, whilst no new signal appeared in the reverse gas feeding sequence. The peak at 1962  $\text{cm}^{-1}$  cannot be attributed to Ru–H or Ru– $\text{H}_2$  species as previously reported<sup>43,49,50</sup> because no signal appeared under the  $\text{H}_2$  atmosphere, even after 8 h holding in such conditions, but appeared only when  $\text{N}_2$  was introduced in the previous step. Therefore, this peak at 1962  $\text{cm}^{-1}$  has been assigned instead to adsorbed azide ( $\text{N}=\text{N}=\text{N}$ ) or oxy-azide ( $\text{N}=\text{N}=\text{O}$ ) species, as previously reported.<sup>51</sup> Accordingly,  $\text{NH}_3$  was detected in the FT-IR chamber gas phase only in the  $\text{H}_2$

atmosphere after  $N_2$  flux, with featured peaks 3332, 1630, 965, and  $929\text{ cm}^{-1}$ , as marked in Figure 5b. It must be highlighted that this result is coincident with that previously performed in the reactor (*vide ante*). Combining the results in Figure 5a,b, it can be concluded that the produced  $NH_3$  derives from the nitrogen species detected at  $1962\text{ cm}^{-1}$ , as shown in Figure 5a, in the *in situ* FTIR experiments and is attributed to azide or oxo-azide species.

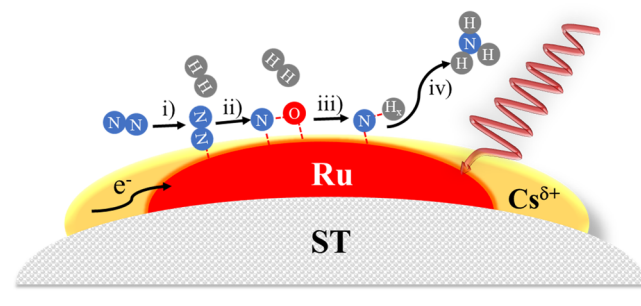
The formation of adsorbed  $N_2$  species was further studied by *in situ* Raman spectroscopy using  $Cs_{10}Ru_2@ST$  and  $Ru_2@ST$  catalysts, and the results are shown in Figure 5c. As can be observed in Figure 5c,  $N_2$  flux on  $Cs_{10}Ru_2@ST$  after the *in situ* activation results in a new peak at  $141\text{ cm}^{-1}$ , which has been attributed to  $\delta NRuO$  species formed on the  $Cs_{10}Ru_2@ST$  surface,<sup>52</sup> and the intensity increases with time in contact with  $N_2$ . Then, the intensity gradually decreased with time after shifting the gas to  $H_2$ , which can be ascribed to the consumption of  $\delta NRuO$  species by  $H_2$ , forming  $NH_3$  products. On the contrary, no signal was observed in the case of  $Ru_2@ST$  (Inset in Figure 5c), indicating that the  $N_2$  could not be easily bonded to the Ru surface or activated, while the presence of the Cs promoter can increase the interaction ability of Ru with  $N_2$  and therefore facilitate the  $N_2$  activation.

The strong interaction of Ru with  $N_2$  in the case of  $Cs_{10}Ru_2@ST$  was also confirmed by thermo-programmed desorption (TPD) experiments. In a first test, the  $Cs_{10}Ru_2@ST$  sample was subjected to  $H_2$  at  $350\text{ }^\circ\text{C}$  for 2 h and then cooled to room temperature with the purge of Ar before  $H_2$  TPD measurement. Alternatively,  $Cs_{10}Ru_2@ST$  was in contact with a mixture of  $H_2$  and  $N_2$  flow (3:1 in volume ratio) at  $350\text{ }^\circ\text{C}$  for 2 h after the activation, and then  $H_2$  TPD measurement was performed after the system cooled to room temperature under Ar flow. As can be observed in Figure S13 in Supporting Information, the  $Cs_{10}Ru_2@ST$  exhibited an intense  $H_2$  desorption peak from 400 to  $600\text{ }^\circ\text{C}$  when the measurement followed after the activation step, which demonstrates that the  $H_2$  can be adsorbed efficiently on the surface of Ru or even penetrate deep in its subsurface. However, the desorption peak was drastically decreased when treating  $Cs_{10}Ru_2@ST$  by the  $H_2/N_2$  mixture, indicating that the  $N_2$  molecule could strongly interact with Ru active sites, which should compete to  $H_2$  adsorption, and therefore, results in low  $H_2$  desorption intensity.

To sum up, results from the abovementioned studies revealed that the partially reduced Cs promoter can be responsible not only for the electron density transfer to Ru sites, but also for the formation of oxy-azide-related species, which can be then sequentially hydrogenated to the final  $NH_3$  product. Moreover, upon light irradiation, Ru NPs absorb visible-NIR light, generating hot electrons and elevating the local temperature at the NP, thus, greatly assisting in the activation of dinitrogen molecules. Subsequently, the activated nitrogen is sequentially hydrogenated and, eventually, ammonia is produced and released. A scheme of the proposed mechanism is presented as Scheme 2.

**2.6. Catalyst Stability.** The stability of the  $Cs_{10}Ru_2@ST$  catalyst was evaluated by conducting the  $N_2$  fixation reaction at  $350\text{ }^\circ\text{C}$  continuously for 120 h under 300 W Xenon lamp irradiation. The temporal profile of the ammonia production rate is shown in Figure S14. As shown in the graph, a slight decay of the activity occurred during a continuous 120 h reaction, demonstrating the high stability of the  $Cs_{10}Ru_2@ST$  catalyst. More specifically, it can be found that only 10% of

**Scheme 2. Illustration of the Proposed Mechanism: (i)  $N_2$  Adsorption and Activation, (ii)  $N_2$  Dissociation and Formation of Intermediate Species, (iii) Hydrogenation of Activated Nitrogen Species, and (iv)  $NH_3$  Formation and Desorption**



activity is lost in 60 h compared to the fresh sample, and additional 10% activity loss at 120 h. In addition, it can be observed from HR-TEM images (Figure S15) that the Ru particle size and dispersity of  $Cs_{10}Ru_2@ST$  remained unchanged before and after 120 h reaction, further confirming the high stability of the  $Cs_{10}Ru_2@ST$  catalyst.

### 3. CONCLUSIONS

This study reports the visible and NIR light-driven improvement in the  $N_2$  hydrogenation reaction to  $NH_3$ , when optimized Cs-decorated strontium titanate-supported Ru NPs is used as a catalyst. Isotopic  $^{15}N$  labeling experiments firmly confirm  $N_2$  as the origin of  $NH_3$ . The  $NH_3$  production is 68% increased upon  $1080\text{ W m}^{-2}$  irradiation (corresponding to about  $600\text{ W m}^{-2}$  of effective light power) and depends on the wavelength, the NIR radiations being more efficient than wavelengths in the visible range, while UV light does not contribute to the  $NH_3$  production rate. This makes the photocatalyst very appropriate for the use of solar light. Mechanistic studies indicate that the reaction occurs upon light absorption by Ru active sites causing a localized temperature increase (photothermal mechanism) and that ST plays a key role due to its basicity and low thermal conductivity. *In situ* FTIR experiments have been carried out to determine the reaction mechanism. Considering that upon illumination the  $Cs_{10}Ru_2@ST$  photocatalyst reported here is among the most efficient catalysts reported so far for  $NH_3$  formation, the present finding opens the door for further exploration of light assistance to ammonia synthesis and helps in the development of more advanced photo-responsive catalysts.

### ■ ASSOCIATED CONTENT

#### Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/acscatal.2c00509>.

Catalyst preparation and characterization, reaction setup and test methodology, catalyst performance and stability data, XRD,  $^1H$  NMR, EDS and mapping, HR-TEM, CO chemisorption, UV-vis absorption, XPS, TPD data, and a table list of the state-of-the-art of the catalysts for nitrogen fixation (PDF)

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## Author Contributions

Y.P. prepared the catalysts and performed the photocatalytic reactions. A.F. characterized the samples. P.C. performed the CO-FTIR measurements and *in situ* FT-IR and Raman study. The work was supervised by J.A. and H.G. who wrote the draft. The manuscript was revised by all the authors

## Notes

The authors declare no competing financial interest.

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## REFERENCES

- (1) Mateo, D.; De Masi, D.; Albero, J.; Lacroix, L. M.; Fazzini, P. F.; Chaudret, B.; García, H. Synergism of Au and Ru Nanoparticles in Low-Temperature Photoassisted CO<sub>2</sub> Methanation. *Chem.—Eur. J.* **2018**, *24*, 18436–18443.
- (2) Zhou, S.; Shang, L.; Zhao, Y.; Shi, R.; Waterhouse, G. I. N.; Huang, Y. C.; Zheng, L.; Zhang, T. Pd Single-Atom Catalysts on Nitrogen-Doped Graphene for the Highly Selective Photothermal Hydrogenation of Acetylene to Ethylene. *Adv. Mater.* **2019**, *31*, 1900509.
- (3) Ling, L.-L.; Yang, W.; Yan, P.; Wang, M.; Jiang, H.-L. Light-Assisted CO<sub>2</sub> Hydrogenation over Pd<sub>3</sub>Cu@UiO-66 Promoted by Active Sites in Close Proximity. *Angew. Chem., Int. Ed.* **2022**, *61*, 202116396.
- (4) Espinosa, J. C.; Catalá, C.; Navalón, S.; Ferrer, B.; Álvaro, M.; García, H. Iron oxide nanoparticles supported on diamond nano-

particles as efficient and stable catalyst for the visible light assisted Fenton reaction. *Appl. Catal., B* **2018**, *226*, 242–251.

(5) Sun, Z.; Fang, S.; Lin, Y.; Hu, Y. H. Photo-assisted methanol steam reforming on solid solution of Cu-Zn-Ti oxide. *Chem. Eng. J.* **2019**, *375*, 121909.

(6) Prajapati, P. K.; Saini, S.; Jain, S. L. Nickel mediated palladium free photocatalytic Suzuki-coupling reaction under visible light irradiation. *J. Mater. Chem. A* **2020**, *8*, 5246–5254.

(7) Mori, K.; Kawashima, M.; Yamashita, H. Visible-light-enhanced Suzuki–Miyaura coupling reaction by cooperative photocatalysis with an Ru–Pd bimetallic complex. *Chem. Commun.* **2014**, *50*, 14501–14503.

(8) Wrighton, M.; Hammond, G. S.; Gray, H. B. Group VI metal carbonyl photoassisted isomerization of olefins. *J. Organomet. Chem.* **1974**, *70*, 283–301.

(9) Kim, C.; Lee, H. Light-assisted surface reactions on metal nanoparticles. *Catal. Sci. Technol.* **2018**, *8*, 3718–3727.

(10) Cheng, G.; Tan, X.; Song, X.; Chen, X.; Dai, W.; Yuan, R.; Fu, X. Visible light assisted thermocatalytic reaction of CO+ NO over Pd/LaFeO<sub>3</sub>. *Appl. Catal., B* **2019**, *251*, 130–142.

(11) Hogan, N. J.; Urban, A. S.; Ayala-Orozco, C.; Pimpinelli, A.; Nordlander, P.; Halas, N. J. Nanoparticles heat through light localization. *Nano Lett.* **2014**, *14*, 4640–4645.

(12) Fuku, K.; Hayashi, R.; Takakura, S.; Kamegawa, T.; Mori, K.; Yamashita, H. The synthesis of size- and color-controlled silver nanoparticles by using microwave heating and their enhanced catalytic activity by localized surface plasmon resonance. *Angew. Chem.* **2013**, *125*, 7594–7598.

(13) Zhang, X.; Li, X.; Reish, M. E.; Zhang, D.; Su, N. Q.; Gutiérrez, Y.; Moreno, F.; Yang, W.; Everitt, H. O.; Liu, J. Plasmon-enhanced catalysis: Distinguishing thermal and nonthermal effects. *Nano Lett.* **2018**, *18*, 1714–1723.

(14) Ghossoub, M.; Xia, M.; Duchesne, P. N.; Segal, D.; Ozin, G. Principles of photothermal gas-phase heterogeneous CO<sub>2</sub> catalysis. *Energy Environ. Sci.* **2019**, *12*, 1122–1142.

(15) Gan, Z.; Wu, X.; Meng, M.; Zhu, X.; Yang, L.; Chu, P. K. Photothermal contribution to enhanced photocatalytic performance of graphene-based nanocomposites. *ACS Nano* **2014**, *8*, 9304–9310.

(16) Lu, S.; Liu, F.; Qiu, P.; Qiao, M.; Li, Y.; Cheng, Z.; Xue, N.; Hou, X.; Xu, C.; Xiang, Y.; Peng, F.; Guo, Z. Photothermal-assisted photocatalytic degradation with ultrahigh solar utilization: Towards practical application. *Chem. Eng. J.* **2020**, *379*, 122382.

(17) Mateo, D.; Albero, J.; García, H. Photoassisted methanation using Cu<sub>2</sub>O nanoparticles supported on graphene as a photocatalyst. *Energy Environ. Sci.* **2017**, *10*, 2392–2400.

(18) Mateo, D.; Albero, J.; García, H. Titanium-Perovskite-Supported RuO<sub>2</sub> Nanoparticles for Photocatalytic CO<sub>2</sub> Methanation. *Joule* **2019**, *3*, 1949–1962.

(19) Barrio, J.; Mateo, D.; Albero, J.; García, H.; Shalom, M. A Heterogeneous Carbon Nitride–Nickel Photocatalyst for Efficient Low-Temperature CO<sub>2</sub> Methanation. *Adv. Energy Mater.* **2019**, *9*, 1902738.

(20) Chen, X.; Li, N.; Kong, Z.; Ong, W.-J.; Zhao, X. Photocatalytic fixation of nitrogen to ammonia: state-of-the-art advancements and future prospects. *Mater. Horiz.* **2018**, *5*, 9–27.

(21) Li, X.; Zhang, X.; Everitt, H. O.; Liu, J. Light-Induced Thermal Gradients in Ruthenium Catalysts Significantly Enhance Ammonia Production. *Nano Lett.* **2019**, *19*, 1706–1711.

(22) Li, J.; Wu, J.; Wang, H.; Lu, Y.; Ye, T.; Sasase, M.; Wu, X.; Kitano, M.; Inoshita, T.; Hosono, H. Acid-durable electrode with layered ruthenium for ammonia synthesis: boosting the activity via selective etching. *Chem. Sci.* **2019**, *10*, 5712–5718.

(23) Chu, K.; Liu, Y.-p.; Wang, J.; Zhang, H. NiO Nanodots on Graphene for Efficient Electrochemical N<sub>2</sub> Reduction to NH<sub>3</sub>. *ACS Appl. Energy Mater.* **2019**, *2*, 2288–2295.

(24) Liu, L.; Puga, A. V.; Coreo, J.; Concepción, P.; Pérez-Dieste, V.; García, H.; Corma, A. Sunlight-assisted hydrogenation of CO<sub>2</sub> into ethanol and C<sub>2+</sub> hydrocarbons by sodium-promoted Co@C nanocomposites. *Appl. Catal., B* **2018**, *235*, 186–196.



- (25) Pham, H. N.; Sattler, J. J. H. B.; Weckhuysen, B. M.; Datye, A. K. Role of Sn in the Regeneration of Pt/ $\gamma$ -Al<sub>2</sub>O<sub>3</sub> Light Alkane Dehydrogenation Catalysts. *ACS Catal.* **2016**, *6*, 2257–2264.
- (26) Burch, R.; Coleman, M. D. An Investigation of Promoter Effects in the Reduction of NO by H<sub>2</sub> under Lean-Burn Conditions. *J. Catal.* **2002**, *208*, 435–447.
- (27) Rossetti, L.; Pernicone, N.; Forni, L. Promoters effect in Ru/C ammonia synthesis catalyst. *Appl. Catal., A* **2001**, *208*, 271–278.
- (28) Larichev, Y. V.; Moroz, B. L.; Moroz, E. M.; Zaikovskii, V. I.; Yunusov, S. M.; Kalyuzhnaya, E. S.; Shur, V. B.; Bukhtiyarov, V. I. Effect of the Support on the Nature of Metal-Promoter Interactions in Ru-Cs+/MgO and Ru-Cs+-Al<sub>2</sub>O<sub>3</sub> Catalysts for Ammonia Synthesis. *Kinet. Catal.* **2005**, *46*, 891–899.
- (29) Aika, K.-i. Role of alkali promoter in ammonia synthesis over ruthenium catalysts—Effect on reaction mechanism. *Catal. Today* **2017**, *286*, 14–20.
- (30) Tang, Y.; Kobayashi, Y.; Masuda, N.; Uchida, Y.; Okamoto, H.; Kageyama, T.; Hosokawa, S.; Loyer, F.; Mitsuhara, K.; Yamanaka, K.; Tamenori, Y.; Tassel, C.; Yamamoto, T.; Tanaka, T.; Kageyama, H. Metal-Dependent Support Effects of Oxyhydride-Supported Ru, Fe, Co Catalysts for Ammonia Synthesis. *Adv. Energy Mater.* **2018**, *8*, 1801772.
- (31) Kitano, M.; Inoue, Y.; Ishikawa, H.; Yamagata, K.; Nakao, T.; Tada, T.; Matsuishi, S.; Yokoyama, T.; Hara, M.; Hosono, H. Essential role of hydride ion in ruthenium-based ammonia synthesis catalysts. *Chem. Sci.* **2016**, *7*, 4036–4043.
- (32) Siporin, S.; Davis, R. J. Use of kinetic models to explore the role of base promoters on Ru/MgO ammonia synthesis catalysts. *J. Catal.* **2004**, *225*, 359–368.
- (33) Aika, K.; Takano, T.; Murata, S. Preparation and characterization of chlorine-free ruthenium catalysts and the promoter effect in ammonia synthesis: 3. A magnesia-supported ruthenium catalyst. *J. Catal.* **1992**, *136*, 126–140.
- (34) Yaghoobi, P.; Vahdani Moghaddam, M.; Nojeh, A. Solar electron source and thermionic solar cell. *AIP Adv.* **2012**, *2*, 042139.
- (35) Luo, S.; Ren, X.; Lin, H.; Song, H.; Ye, J. Plasmonic photothermal catalysis for solar-to-fuel conversion: current status and prospects. *Chem. Sci.* **2021**, *12*, 5701–5719.
- (36) McClory, M.; Gonzalez, R. D. The role of alkali metals as promoters in the methanation and Fischer-Tropsch reaction: An in situ study. *J. Catal.* **1984**, *89*, 392–403.
- (37) Resasco, J.; Chen, L. D.; Clark, E.; Tsai, C.; Hahn, C.; Jaramillo, T. F.; Chan, K.; Bell, A. T. Promoter Effects of Alkali Metal Cations on the Electrochemical Reduction of Carbon Dioxide. *J. Am. Chem. Soc.* **2017**, *139*, 11277–11287.
- (38) Kuei, C.-K.; Lee, J.-F.; Lee, M.-D. The Promoter Effects of Alkali Metal to Supported Iron Catalyst in Carbon Monoxide Hydrogenation, The Influence of Alkali on Carbon. *Chem. Eng. Commun.* **1991**, *101*, 77–92.
- (39) Aika, K.-i.; Toshikazu, K.; Shuzo, M.; Takaharu, O. Promoter Effect of Alkali Metal Oxides and Alkali Earth Metal Oxides on Active Carbon-Supported Ruthenium Catalyst for Ammonia Synthesis. *Bull. Chem. Soc. Jpn.* **1990**, *63*, 1221–1225.
- (40) Sattler, J. J. H. B.; Ruiz-Martinez, J.; Santillan-Jimenez, E.; Weckhuysen, B. M. Catalytic Dehydrogenation of Light Alkanes on Metals and Metal Oxides. *Chem. Rev.* **2014**, *114*, 10613–10653.
- (41) Liu, Y.-J.; Deng, X.; Huang, W. Effect of non-metal promoters on higher alcohols synthesis from syngas over Cu-based catalyst. *J. Energy Inst.* **2018**, *91*, 1136–1142.
- (42) Huo, C.-F.; Wu, B.-S.; Gao, P.; Yang, Y.; Li, Y.-W.; Jiao, H. The mechanism of potassium promoter: enhancing the stability of active surfaces. *Angew. Chem.* **2011**, *123*, 7541–7544.
- (43) Kita, Y.; Kuwabara, M.; Yamadera, S.; Kamata, K.; Hara, M. Effects of ruthenium hydride species on primary amine synthesis by direct amination of alcohols over a heterogeneous Ru catalyst. *Chem. Sci.* **2020**, *11*, 9884–9890.
- (44) Yang, K.; Zhang, Y.; Li, Y.; Huang, P.; Chen, X.; Dai, W.; Fu, X. Insight into the function of alkaline earth metal oxides as electron

promoters for Au/TiO<sub>2</sub> catalysts used in CO oxidation. *Appl. Catal., B* **2016**, *183*, 206–215.

(45) Wang, F.; Ueda, W.; Xu, J. Detection and Measurement of Surface Electron Transfer on Reduced Molybdenum Oxides (MoOx) and Catalytic Activities of Au/MoOx. *Angew. Chem., Int. Ed.* **2012**, *51*, 3883–3887.

(46) Bharti, B.; Kumar, S.; Lee, H.-N.; Kumar, R. Formation of oxygen vacancies and Ti<sup>3+</sup> state in TiO<sub>2</sub> thin film and enhanced optical properties by air plasma treatment. *Sci. Rep.* **2016**, *6*, 32355.

(47) Ahmad, Z.; Najeeb, M. A.; Shakoor, R. A.; Alshraf, A.; Al-Muhtaseb, S. A.; Soliman, A.; Nazeeruddin, M. K. Instability in CH<sub>3</sub>NH<sub>3</sub>PbI<sub>3</sub> perovskite solar cells due to elemental migration and chemical composition changes. *Sci. Rep.* **2017**, *7*, 15406.

(48) Banger, K. K.; Yamashita, Y.; Mori, K.; Peterson, R. L.; Leedham, T.; Rickard, J.; Sirringhaus, H. Low-temperature, high-performance solution-processed metal oxide thin-film transistors formed by a ‘sol-gel on chip’ process. *Nat. Mater.* **2011**, *10*, 45–50.

(49) Choi, J.-H.; Schloerer, N. E.; Berger, J.; Precht, M. H. G. Synthesis and characterisation of ruthenium dihydrogen complexes and their reactivity towards B–H bonds. *Dalton Trans.* **2014**, *43*, 290–299.

(50) Ashok, M.; Prasad, A. V. S. S.; Ravinder, V. Synthesis, spectral studies and catalytic activity of ruthenium(II) complexes with organic amide ligands. *J. Braz. Chem. Soc.* **2007**, *18*, 1492–1499.

(51) Maji, S.; Sarkar, B.; Patra, M.; Das, A. K.; Mobin, S. M.; Kaim, W.; Lahiri, G. K. Formation, Reactivity, and Photorelease of Metal Bound Nitrosyl in [Ru(trpy)(L)(NO)]<sup>n+</sup> (trpy = 2,2′:6′,2″-Terpyridine, L = 2-Phenylimidazo[4,5-f]1,10-phenanthroline). *Inorg. Chem.* **2008**, *47*, 3218–3227.

(52) Villalba, M. E. C.; Echeverría, G.; Aymonino, P. J.; Varetto, E. L. Crystal Structure and Vibrational Spectra of [Ru(NH<sub>3</sub>)<sub>4</sub>(OH)(NO)]ZnCl<sub>4</sub>. *J. Chem. Crystallogr.* **2011**, *41*, 508–513.

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