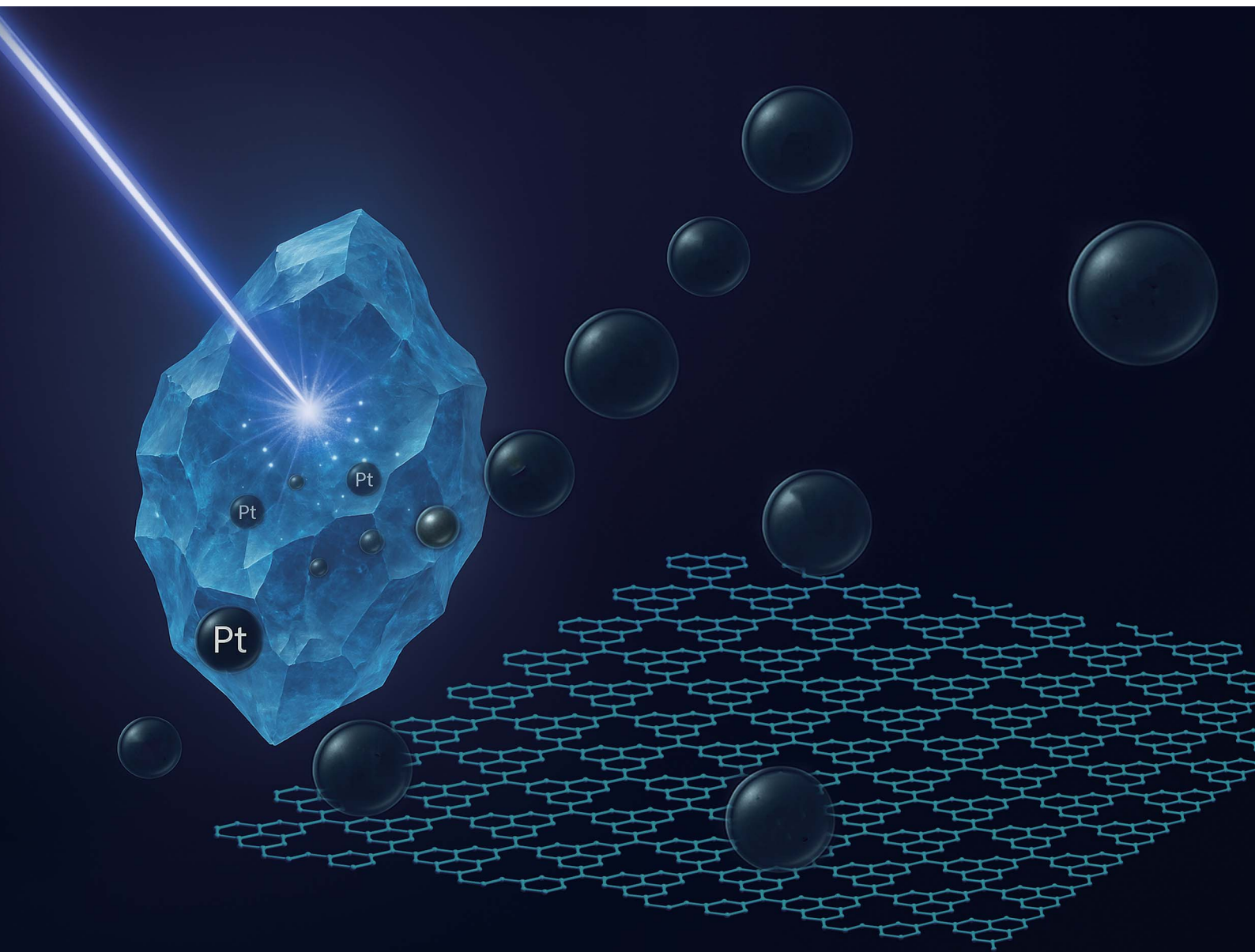


# Nanoscale Advances

Volume 7  
Number 9  
7 May 2025  
Pages 2385–2762

rsc.li/nanoscale-advances



ISSN 2516-0230

**COMMUNICATION**

Yongming Fu, Jie Ma *et al.*  
Pulsed laser synthesis of free-standing Pt single atoms  
in an ice block for enhancing photocatalytic hydrogen  
evolution of g-C<sub>3</sub>N<sub>4</sub>

Cite this: *Nanoscale Adv.*, 2025, 7, 2474Received 13th January 2025  
Accepted 25th March 2025

DOI: 10.1039/d5na00043b

rsc.li/nanoscale-advances

# Pulsed laser synthesis of free-standing Pt single atoms in an ice block for enhancing photocatalytic hydrogen evolution of g-C<sub>3</sub>N<sub>4</sub>†

Yongming Fu,<sup>ID</sup>\*<sup>ab</sup> Qianyu Lu,<sup>ab</sup> Jianhong Wang,<sup>ab</sup> Na Sun,<sup>ab</sup> Jinjun Gao,<sup>ab</sup>  
Peng Chen,<sup>ab</sup> Jizhou Wu,<sup>ID</sup><sup>abc</sup> and Jie Ma<sup>\*abc</sup>

This study reports an innovative synthesis method of a Pt/g-C<sub>3</sub>N<sub>4</sub> single atom catalyst for enhancing photocatalytic hydrogen evolution. The method involves the synthesis of free-standing Pt single atoms within an H<sub>2</sub>PtCl<sub>6</sub> ice block using a pulsed laser reduction process, followed by transferring them onto few-layer g-C<sub>3</sub>N<sub>4</sub> through electrostatic adsorption at low temperature. This approach eliminates the need for high-energy lasers and porous support materials during laser solid-phase synthesis. The photocatalytic activities of Pt/g-C<sub>3</sub>N<sub>4</sub> synthesized under various laser conditions are evaluated to optimize the synthesis parameters. The optimal Pt/g-C<sub>3</sub>N<sub>4</sub> catalyst demonstrates a significantly higher photocatalytic hydrogen evolution capability (320 μmol h<sup>-1</sup>), 129 times that of pure g-C<sub>3</sub>N<sub>4</sub> (2.2 μmol h<sup>-1</sup>). This work expands the laser-solid phase synthesis method, offering a promising route for the production of single atom catalysts with simple operation, clear synthetic pathways, low cost, and environmental friendliness.

the most significant challenge in SAC is the aggregation of single atoms arising from their high surface free energy, which significantly compromises their unique catalytic properties and ultimately limits the practical applications of SAC.<sup>5</sup>

To address this challenge, researchers are actively exploring innovative approaches to enhance the dispersion of single atoms. One effective approach is the spatial confinement strategy, which involves the encapsulation of single atoms within molecular cages to prevent their migration.<sup>6</sup> Unlike the traditional spatial confinement techniques that depend on the use of porous materials, Wu *et al.* proposed a novel ice-phase confinement method.<sup>7</sup> In this approach, Pt ions in an H<sub>2</sub>PtCl<sub>6</sub> aqueous solution are stabilized and dispersed by forming H–Pt–OH complexes with water molecules. The solution is subsequently frozen into ice blocks, where the crystalline structure of water molecules restricts the thermal movement of Pt atoms during the photoreduction process. Although this innovative technique effectively suppresses the aggregation of Pt single atoms, the low photon density of conventional light sources, such as mercury lamps, limits the quantity of synthesized Pt single atoms.

Laser reduction methods have been developed for one-step synthesis of SAC, capitalizing on the localized high temperature and rapid cooling effects generated by high-energy laser pulses interacting with the support material.<sup>8–12</sup> The high photon density in a single laser pulse significantly increases the probability of interactions between photons and metal precursors, thereby enhancing the yield of single atoms. Furthermore, the brief duration of the laser pulse limits the formation of single atoms at any given time, effectively minimizing the aggregation and nucleation of single atoms. For instance, Liu *et al.* proposed a laser solid-phase synthesis technique that facilitates the simultaneous reduction of metal precursors and graphene oxide in one step.<sup>10</sup> Zou *et al.* introduced a laser planting strategy in which high-energy laser pulses decompose metal precursors into single atoms while concurrently inducing vacancy defects on the surface of support materials for anchoring single atoms through electrostatic interactions.<sup>13</sup> Compared to the conventional photochemical reduction methods, laser reduction methods

## 1. Introduction

Single atom catalysts (SAC) have garnered significant attention for various photocatalytic and electrocatalytic applications, including CO<sub>2</sub> reduction, water splitting, CO oxidation, and N<sub>2</sub> fixation.<sup>1–3</sup> In SAC, isolated atoms are anchored on the surface of a support material through electron exchange interactions, forming dispersed and localized active sites. The distinctive structure of SAC allows for precise control over catalytic reactions at the atomic level, maximizing catalytic efficiency, selectivity, and atomic utilization.<sup>4</sup> Despite their notable advantages,

\*School of Physics and Electronic Engineering & Institute of Laser Spectroscopy, State Key Laboratory of Quantum Optics and Quantum Optics Devices, Shanxi University, Taiyuan 030006, China. E-mail: fuyongming@sxu.edu.cn; mj@sxu.edu.cn

<sup>b</sup>Xinzhou Institute of Innovation Ecosystem, Shanxi University, Xinzhou 034000, China  
<sup>c</sup>Collaborative Innovation Center of Extreme Optics, Shanxi University, Taiyuan 030006, China

† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d5na00043b>



eliminate the need for high-temperature treatments and additional reductants, enabling the one-step synthesis of high-loading metal single atoms under mild conditions. However, the laser synthesis methods rely on high pulsed energy to introduce vacancy defects on the support material surface for stabilizing single atoms, resulting in a strong dependence on the high-power laser and the specific characteristics of the support material.

In this letter, we present an innovative method for synthesizing SAC by integrating the freeze-phase photochemical method with the laser reduction technique. Free-standing Pt single atoms are synthesized within the  $\text{H}_2\text{PtCl}_6$  ice block through a pulsed laser reduction process, followed by transferring them onto few-layer  $\text{g-C}_3\text{N}_4$  at low temperature. The 355 nm pulsed laser directly induces the reduction of the Pt precursor, eliminating the need for high-energy lasers and porous support materials. Freezing the precursor solution facilitates the confinement of metal atoms by water molecules, ensuring stable dispersion and preventing nucleation during the formation of Pt single atoms. Two-dimensional  $\text{g-C}_3\text{N}_4$  with an intrinsic six-fold cavity is chosen as the support material,<sup>14</sup> enabling the electrostatic adsorption of Pt single atoms by Pt–N interaction and further suppressing atom agglomeration. Furthermore, the gradual thawing of the ice block reduces the concentration of dissociative single atoms in the suspension, promoting their successful anchoring onto the  $\text{g-C}_3\text{N}_4$ . The photocatalytic hydrogen evolution capabilities of Pt single atoms anchored on  $\text{g-C}_3\text{N}_4$  ( $\text{Pt/g-C}_3\text{N}_4$ ) synthesized under different conditions are evaluated to investigate the effect of the pulse repetition frequency and scanning speed. This method offers a promising route for the production of SAC, featuring simple operation, clear synthetic pathways, low cost, and environmental sustainability.

## 2. Experimental

Free-standing Pt single atoms were directly synthesized through a laser freeze-phase reduction method using a home-made laser scanning set (Fig. S1†). 10 mL of  $\text{H}_2\text{PtCl}_6$  aqueous solution ( $1 \text{ mg mL}^{-1}$ ) was transferred to a stainless-steel cup with a diameter of 3 cm and rapidly frozen in liquid nitrogen to form a cylindrical ice block with a thickness of 20 mm (Fig. S2a†). The ice block was then placed under a commercially available low-power pulsed laser (P-Plus2-355-5, Huaray, China) with a wavelength of 355 nm. The laser was set at a current of 1 A, a repetition frequency of 50 kHz, and a pulse width of 50 ns. A lens with a focal length of 190 mm was used to focus the laser beam onto the upper surface of the ice block, achieving a focused spot diameter of 25  $\mu\text{m}$  (Fig. S2b†). A two-dimensional galvo scanner was employed to control the laser spot for continuous horizontal scanning throughout the ice block surface at a scanning speed of 700  $\text{mm s}^{-1}$ . To ensure the entire volume of the ice block was exposed to the laser rather than just at the surface, a linear stage was used to control the vertical scanning of the laser beam with a speed of 28  $\text{mm min}^{-1}$ . To compensate the power loss at greater depths caused by the absorption of the ice block (Fig. S3†), multiple repeated scans were conducted on the ice block from top to bottom. During the scanning process, the

ice block was maintained in liquid nitrogen to prevent melting. After a scanning duration of 140 min, free-standing Pt single atoms were obtained in the ice block.

The Pt single atoms were anchored onto few-layer  $\text{g-C}_3\text{N}_4$  through strong electrostatic adsorption at low temperature. Bulk  $\text{g-C}_3\text{N}_4$  powder was synthesized by thermal polymerization of urea. A few-layer  $\text{g-C}_3\text{N}_4$  suspension was prepared by dispersing 50 mg of bulk  $\text{g-C}_3\text{N}_4$  powder in 10 mL of deionized water, followed by ultrasonic treatment at a power of 400 W for 2 h. The ice block containing Pt single atoms was put into the suspension under vigorous stirring for 48 h. The temperature was maintained at 2 °C to slow the ice melting. Subsequently, the 20 mL suspension was centrifuged 5 times to remove any residual Pt species. Finally, the resulting  $\text{Pt/g-C}_3\text{N}_4$  powder was dried using a freeze-drying process.

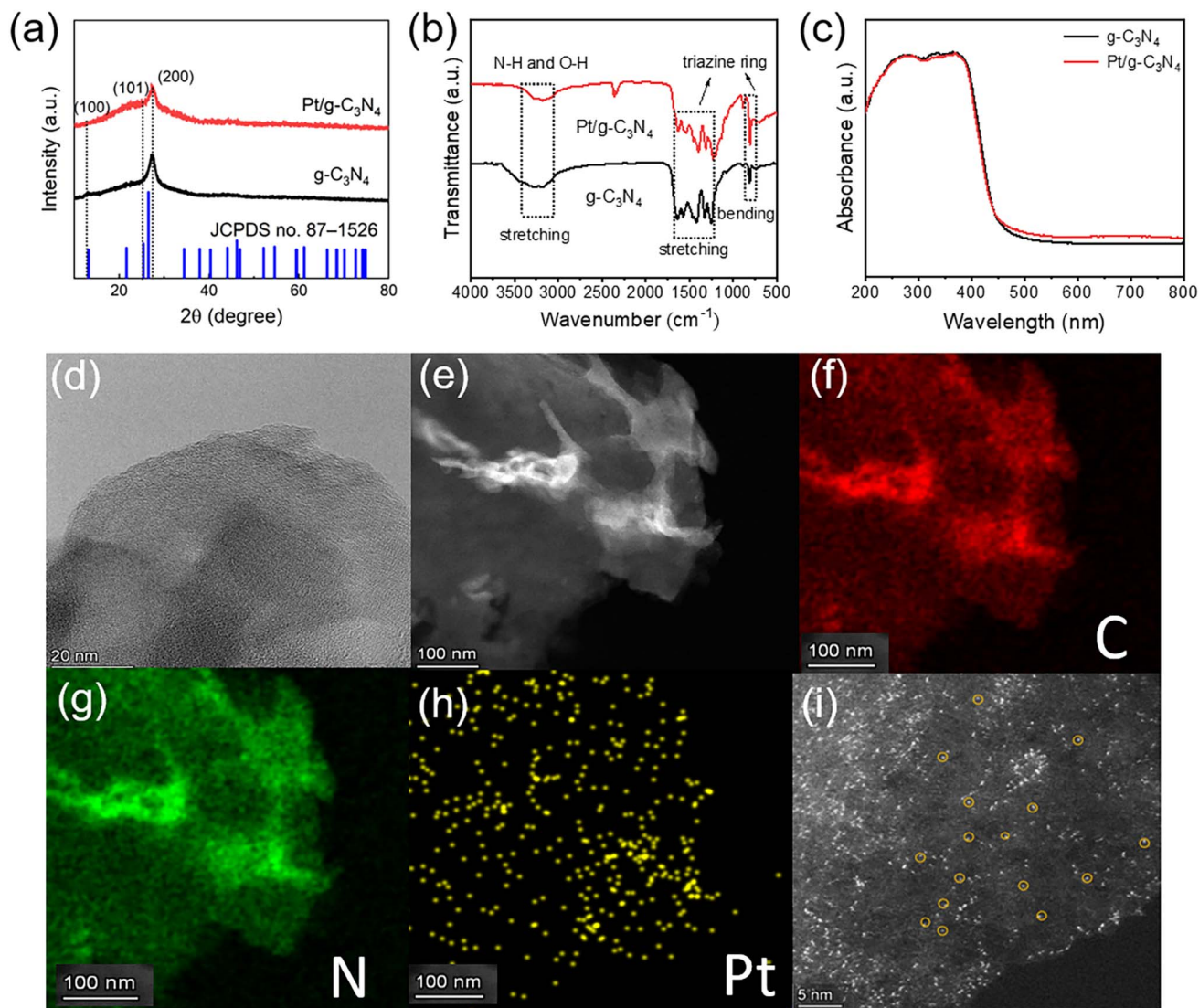
The Pt content in  $\text{Pt/g-C}_3\text{N}_4$  was quantified using inductively coupled plasma-optical emission spectroscopy (ICP-OES, 5510, Agilent, USA). The surface morphology was characterized with a transmission electron microscope (TEM, JEM-F200, JEOL, Japan). The identification of crystalline phases, chemical functional groups, and surface chemical states was performed using X-ray diffraction (XRD, Rigaku Miniflex 600, Japan), Fourier transform infrared spectroscopy (FTIR, Nicolet iS20, Thermo Scientific, USA), and X-ray photoelectron spectrometry (XPS, K-Alpha, Thermo Scientific, USA), respectively. Diffuse reflectance spectroscopy (DRS, UH5700, Hitachi, Japan) was employed to acquire the UV-visible absorption spectroscopy data. The steady and time-resolved photoluminescence (PL) spectra were measured using a photoluminescence spectrometer (FLS1000, Edinburgh, UK). High angle annular dark-field scanning transmission electron microscopy (HAADF-STEM, JEM-ARM200F, JEOL, Japan) images were captured to confirm the atomic dispersion of Pt atoms. X-ray absorption fine spectroscopy (XAFS) measurement of the Pt element was conducted at the XAFS beamline of the Shanghai Synchrotron Radiation Facility (SSRF), where the experimental data were acquired in fluorescence mode using an ionization chamber.

The photocatalytic water splitting reactions were conducted using a photocatalytic activity evaluation system (CEL-PAEM-D8Plus, CEALIGHT, China). In a typical experiment, 50 mg of  $\text{Pt/g-C}_3\text{N}_4$  catalyst was suspended in 80 mL of deionized water and 20 mL of triethanolamine. Prior to the photocatalytic test, the closed-loop system was evacuated for 20 min to eliminate the air in the reaction system. The generated  $\text{H}_2$  gas was analyzed every 30 min using gas chromatography (CG-7920, CEALIGHT, China) with Ar gas as the carrier gas. During the photocatalytic test, a 300 W xenon lamp was used to provide the simulated light source. The system was irradiated for 4 h to assess the  $\text{H}_2$  evolution performance of SAC. The reaction temperature was maintained at 4 °C using cooling water.

## 3. Results and discussion

Fig. 1a presents the XRD patterns of few-layer  $\text{g-C}_3\text{N}_4$  and  $\text{Pt/g-C}_3\text{N}_4$ . The characteristic diffraction peaks at 12.95°, 25.31°, and 27.38° correspond to the (100), (101), and (002) crystallographic planes of  $\text{g-C}_3\text{N}_4$  (JCPDS file no. 87-1526), respectively.<sup>15</sup> The





**Fig. 1** Microstructures of few-layer  $g\text{-C}_3\text{N}_4$  and  $\text{Pt}/g\text{-C}_3\text{N}_4$ . (a) XRD patterns. (b) FTIR spectra. (c) DRS spectra. (d) High-resolution TEM image of  $\text{Pt}/g\text{-C}_3\text{N}_4$ . (e–h) Elemental mapping profiles of  $\text{Pt}/g\text{-C}_3\text{N}_4$  under the dark-field of HAADF-STEM (e), C (f), N (g), and Pt (h). (i) High-resolution HAADF-STEM image under dark-field to clearly demonstrate Pt single atoms.

diminished intensity and enhanced angle of the (002) peak suggests a reduced long-range order and enlarged interface distance in the stacking of  $g\text{-C}_3\text{N}_4$  layers, which aligns with the characteristics of few-layer structure.<sup>16</sup> The diminished intensity of the (002) peak suggests a reduced long-range order in the stacking of  $g\text{-C}_3\text{N}_4$  layers, which aligns with the characteristics of few-layer structure. As depicted in the FTIR spectra (Fig. 1b), the two samples exhibit similar vibrational peak bands corresponding to N–H stretching ( $3100\text{--}3300\text{ cm}^{-1}$ ), C=N or C–N stretching ( $1200\text{--}1700\text{ cm}^{-1}$ ), and triazine breathing modes ( $811\text{ cm}^{-1}$ ) in heptazine rings, indicating that the introduction of Pt species does not significantly affect the molecular backbone of  $g\text{-C}_3\text{N}_4$ .<sup>17</sup> DRS results demonstrate that the absorption intensity in the UV region of the  $\text{Pt}/g\text{-C}_3\text{N}_4$  is comparable to that of few-layer  $g\text{-C}_3\text{N}_4$  (Fig. 1c). By extrapolating the intercept of the  $x$ -axis in the linear region of Tauc plots, the band gaps of  $g\text{-C}_3\text{N}_4$  and  $\text{Pt}/g\text{-C}_3\text{N}_4$  are calculated to be 2.65 and 2.66 eV, respectively,

suggesting that the bandgap is not effectively affected by Pt incorporation. The formation of Pt single atoms anchored on few-layer  $g\text{-C}_3\text{N}_4$  is then morphologically studied. Fig. 1d is the TEM image of  $\text{Pt}/g\text{-C}_3\text{N}_4$ . There are no obvious Pt nanoparticles on the surface of few-layer  $g\text{-C}_3\text{N}_4$ . However, the homogeneous distribution of C, N, and Pt elements (Fig. 1e–h) suggests that trace amounts of Pt atoms exist in the form of isolated single atoms on the surface of  $g\text{-C}_3\text{N}_4$ , which is further demonstrated by the densely dispersed bright spots (Fig. 1i) in HAADF-STEM analysis under dark-field.<sup>18</sup> These results verify the successful synthesis of free-standing Pt single atoms through the laser freeze-phase reduction method and subsequent transfer onto few-layer  $g\text{-C}_3\text{N}_4$  through low-temperature electrostatic adsorption. The mass fraction of Pt single atoms in  $\text{Pt}/g\text{-C}_3\text{N}_4$  is measured to be 2.37% by HAADF-STEM and 1.19% by ICP-OES, indicating a relatively high loading mass of Pt single atoms on few-layer  $g\text{-C}_3\text{N}_4$ .



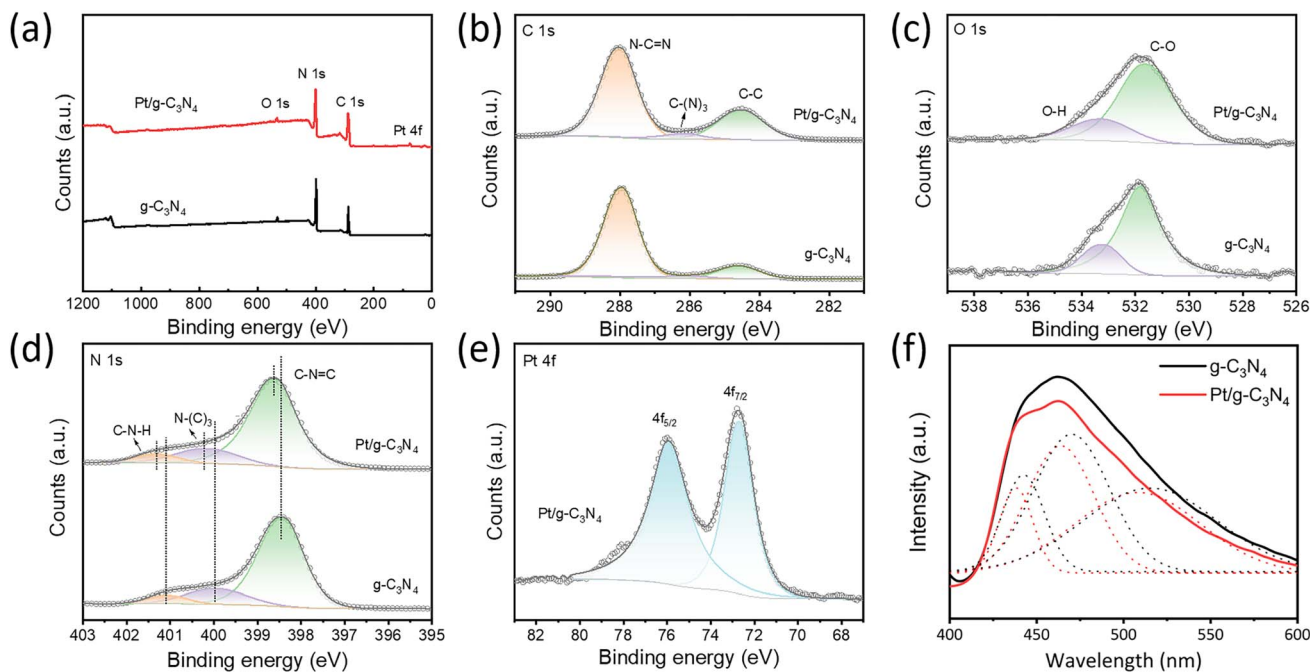


Fig. 2 Chemical state of few-layer  $g\text{-C}_3\text{N}_4$  and  $\text{Pt}/g\text{-C}_3\text{N}_4$ . (a–e) XPS spectra of survey (a), C 1s (b), O 1s (c), N 1s (d), and Pt 4f (e). (f) PL spectra.

The impact of Pt single atoms on the surface states of few-layer  $g\text{-C}_3\text{N}_4$  is studied by XPS, as illustrated in Fig. 2. The survey spectra indicate the presence of C, N, and O in the  $g\text{-C}_3\text{N}_4$  and Pt in the  $\text{Pt}/g\text{-C}_3\text{N}_4$  (Fig. 2a). The high-resolution C 1s spectrum of  $g\text{-C}_3\text{N}_4$  is well fitted to three distinct peaks at 284.6, 286.1, and 287.9 eV, corresponding to the C–C, C–(N)<sub>3</sub>, and N–C=N bonds, respectively (Fig. 2b). The  $g\text{-C}_3\text{N}_4$  and  $\text{Pt}/g\text{-C}_3\text{N}_4$  also show similar O 1s spectra, which are divided into two peaks at 531.82 and 533.18 eV, corresponding to the C–O and O–H bonds formed during the thermal polymerization process (Fig. 2c).<sup>19</sup> The N 1s spectrum of  $g\text{-C}_3\text{N}_4$  exhibits three peaks classified as pyridinic N (398.40 eV), pyrrolic N (399.95 eV) and amino groups (401.40 eV). For  $\text{Pt}/g\text{-C}_3\text{N}_4$ , the binding energies shift towards higher values (Fig. 2d), which is indicative of the formation of N coordination bonds with Pt single atoms.<sup>20</sup> In the Pt 4f spectrum (Fig. 2e), the two symmetrical peaks located at 72.76 and 75.94 eV correspond to the  $4f_{7/2}$  and  $4f_{5/2}$  states of Pt species, suggesting an oxide state ranging between +2 and +4.<sup>21</sup> PL spectra reveal that both few-layer  $g\text{-C}_3\text{N}_4$  and  $\text{Pt}/g\text{-C}_3\text{N}_4$  exhibit broad emission peaks within 400–600 nm (Fig. 2f). Deconvoluted by Gaussian fitting, the two spectra can be similarly divided into three emission centers belonging to pathways of  $\pi^* \rightarrow \pi$ ,  $\sigma^* \rightarrow \text{LP}$ , and  $\pi^* \rightarrow \text{LP}$  transitions, respectively.<sup>22,23</sup> A red-shift is observed in  $\text{Pt}/g\text{-C}_3\text{N}_4$ , indicating the bandgap narrowing of the  $\text{sp}^2$  C–N hybridize state. Moreover, the PL intensity of  $\text{Pt}/g\text{-C}_3\text{N}_4$  is lower than that of  $g\text{-C}_3\text{N}_4$ , suggesting that the incorporation of Pt single atoms effectively suppresses the recombination of photogenerated charge carriers, thereby enhancing the photocatalytic performance.<sup>24</sup>

The precise coordination environment and chemical state of Pt atoms in  $\text{Pt}/g\text{-C}_3\text{N}_4$  are elucidated through synchrotron-based X-ray absorption near-edge structure (XANES) and extended X-

ray absorption fine structure (EXAFS) (Fig. 3).<sup>25</sup> The energy of the white line peak for Pt species in  $\text{Pt}/g\text{-C}_3\text{N}_4$  is 11 567.5 eV, positioned between reference points of Pt foil and  $\text{PtO}_2$  (Fig. 3a and S4†). The Pt single atom interacts with the pyridinic N and facilitates the transfer of electrons from the Pt atom to  $g\text{-C}_3\text{N}_4$ , resulting in an oxidation state of  $\text{Pt}^{\delta+}$  ( $0 < \delta < 4$ ).<sup>26</sup> The  $k^2$ -weighted  $R$ -space Fourier transformed EXAFS (FT-EXAFS) spectrum of  $\text{Pt}/g\text{-C}_3\text{N}_4$  exhibits a prominent peak around 1.77 Å, corresponding to the first-shell Pt–C/N scattering path (Fig. 3b and Table S1†). The absence of signals at 2.54 Å for the Pt–Pt bond and 1.62 Å for the Pt–O bond further confirms the Pt single atom feature, consistent with the previous HAADF-STEM observations. To ascertain the coordinating atom with the Pt single atom, the EXAFS spectrum of  $\text{Pt}/g\text{-C}_3\text{N}_4$  is fitted, showing that the Pt single atom is anchored to  $g\text{-C}_3\text{N}_4$  through the Pt–N pathway with a coordination number of  $3.8 \pm 0.2$  and a bond length of 2.25 Å (Fig. 3c). The wavelet transformed EXAFS (WT-EXAFS) provides a method for simultaneously analyzing information in both  $R$ -space and  $k$ -space resolution.<sup>27</sup> As shown in Fig. 3d, the maximum intensity of  $\text{Pt}/g\text{-C}_3\text{N}_4$  centered round  $5.42 \text{ \AA}^{-1}$  in  $k$ -space corresponds to a resolution of 1.77 Å in  $R$ -space, which is assigned to the Pt–N coordination structure. Note that the intensities centered at 10.08 and  $5.06 \text{ \AA}^{-1}$  associated with Pt–Pt and Pt–O coordination are not observed in the  $\text{Pt}/g\text{-C}_3\text{N}_4$ . The coordination analyses substantiate the oxidation state of the Pt single atoms and their strong interaction with N atoms in an isolated state. The results indicate that the Pt single atoms synthesized by laser freeze-phase reduction are stably anchored in the six-fold cavities of  $g\text{-C}_3\text{N}_4$  through electrostatic adsorption. The electron-rich N atoms in the  $g\text{-C}_3\text{N}_4$  framework can provide sufficient coordination sites,<sup>28</sup> and the unique structure with six-fold cavity provides most favorable anchored



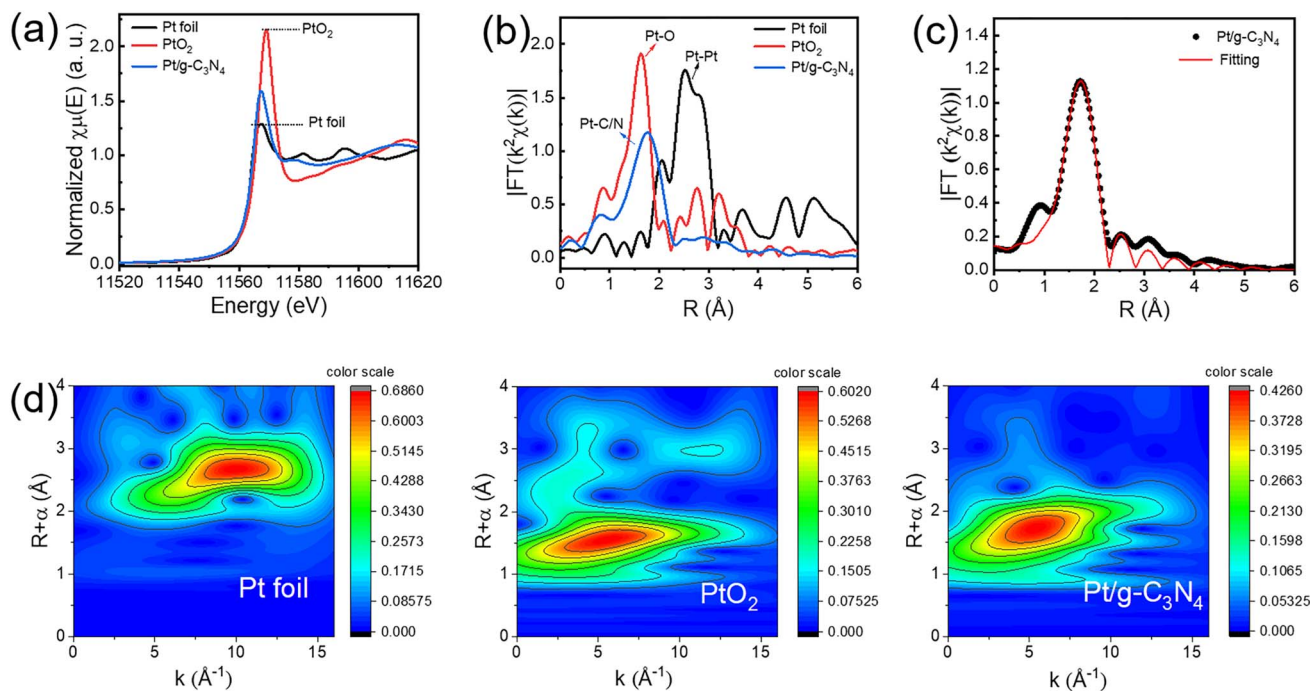


Fig. 3 XAFS results of Pt/g-C<sub>3</sub>N<sub>4</sub> compared with Pt foil and PtO<sub>2</sub>. (a) XANES spectra. (b) FT-EXAFS spectra. (c) FT-EXAFS fitting through Pt–N. (d) WT-EXAFS spectra.

sites for embracing metal single atoms through the formation of metal–nitrogen bonds.<sup>29–31</sup> Furthermore, the deformed wrinkle space of g-C<sub>3</sub>N<sub>4</sub> helps in stabilizing single-atom Pt in the six-fold cavity.<sup>32</sup> This anchoring mechanism is vital for maintaining the dispersion and stability of Pt single atoms, which is essential for enhancing the photocatalytic performance of Pt/g-C<sub>3</sub>N<sub>4</sub>.

To comprehensively examine the influence of synthetic conditions on the photocatalytic hydrogen evolution performance of few-layer g-C<sub>3</sub>N<sub>4</sub>, the free-standing Pt single atoms are synthesized under different laser parameters. These parameters can be generalized by considering the energy density and pulse duration, which are critical factors in laser–material interactions. Based on previous studies involving pulsed laser synthesis of single atoms,<sup>8,10,13</sup> three crucial parameters of horizontal scanning speed, vertical scanning speed, and pulse repetition frequency are focused on. The located energy density can be calculated based on the laser power, spot size, and scanning speed, while the pulse duration determines the thermal effects and cooling rates, which are critical for the synthesis of Pt single atoms. Under a pulse repetition frequency of 20 kHz, it is observed that the speed of 700 mm s<sup>−1</sup> achieves the highest hydrogen production with horizontal scanning speeds ranging from 100 to 900 mm s<sup>−1</sup> (Fig. 4a). The pulsed laser breaks the continuous horizontal scanning into a “point-to-point” scanning mode, which prevents a single scan from covering all Pt species within the ice block. The distance between adjacent points is mainly determined by the horizontal scanning speed. Given that the total scanning time is fixed to 140 min, a faster horizontal scanning speed results in more

repeated scans, leading to a higher coverage rate of the laser spot on the Pt species, and consequently, an increased generation of Pt single atoms. However, excessively high speed may reduce the interaction time between the laser beam and the Pt precursor, negatively impacting the formation of Pt single atoms. Similarly, the pulse repetition frequency, evaluated from 20 to 70 kHz, reveals an optimal frequency of 50 kHz for maximizing hydrogen evolution (Fig. 4b). When the average power and pulse width of the pulsed laser remain constant, the pulse frequency is inversely proportional to the pulse energy. The formation of Pt single atoms primarily occurs during the pulse duration. A lower pulse frequency allows for a longer cooling time, resulting in reduced thermal accumulation of the laser, which helps to prevent the aggregation of Pt single atoms. However, excessively low frequencies result in excessively high pulse energy, intensifying the thermal motion of Pt atoms during a single pulse, which can lead to cluster formation. The vertical scanning speed is also investigated, indicating that a vertical scanning speed of 28 mm min<sup>−1</sup> achieves the best photocatalytic performance (Fig. 4c). The vertical scanning speed is significantly lower than the horizontal speed, primarily influencing the spacing between Pt single atom layers. Due to the recoating effect of pulsed laser scanning, if the vertical scanning speed is too slow, the same ice layer may be scanned repeatedly, causing atom aggregation. Under the optimal conditions—700 mm s<sup>−1</sup> for horizontal scanning speed, 28 mm min<sup>−1</sup> for vertical scanning speed, and 50 kHz for pulse repetition frequency—the hydrogen production rate of the Pt/g-C<sub>3</sub>N<sub>4</sub> reaches 320 μmol h<sup>−1</sup>, representing a remarkable increase of 129 times compared to that of pure g-C<sub>3</sub>N<sub>4</sub> (2.2 μmol h<sup>−1</sup>),



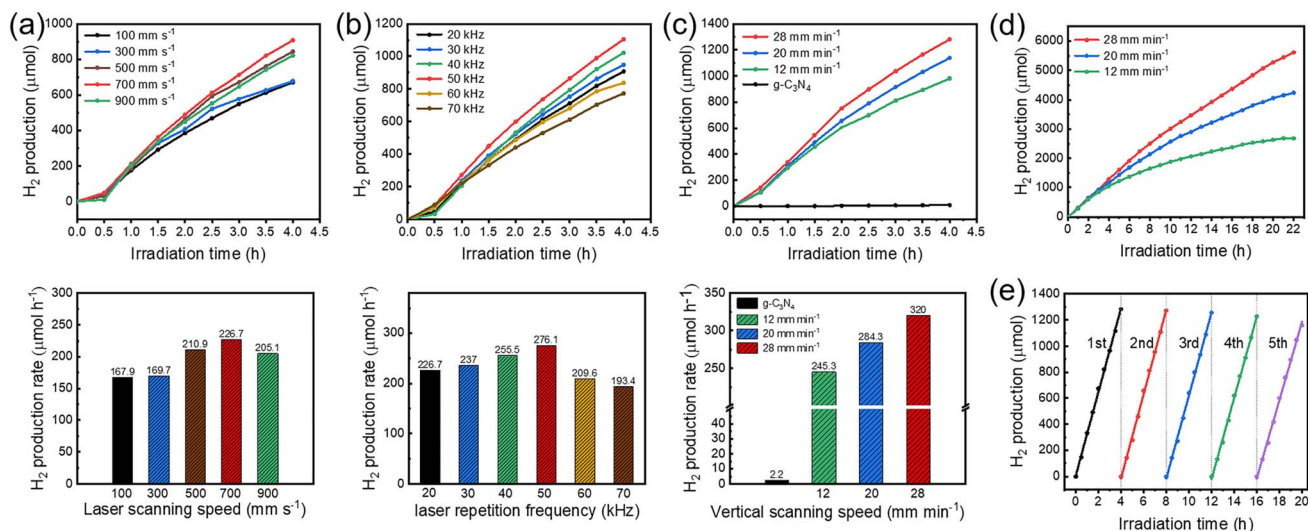


Fig. 4 The photocatalytic hydrogen evolution performance of Pt/g-C<sub>3</sub>N<sub>4</sub> under different synthetic conditions. (a) Effect of laser scanning speed. (b) Effect of laser repetition frequency. (c) Effect of vertical scanning speed. (d) Prolonged photocatalytic test over 22 h. (e) Photocatalytic capabilities within five cycles.

demonstrating a substantial enhancement in photocatalytic activity. The significant enhancement in photocatalytic hydrogen evolution is primarily attributed to the synergistic effects of improved electron transfer and increased reaction sites. Pt single atoms form Pt–N coordination with N atoms in g-C<sub>3</sub>N<sub>4</sub>, leading to the formation of N 2p–Pt 5d hybrid orbitals at the interface.<sup>17</sup> The Pt–N coordination reduces the interfacial charge transfer resistance, which significantly suppresses the recombination of photogenerated electron–hole pairs.<sup>33,34</sup> Moreover, the incorporation of Pt single atoms effectively lowers the Gibbs free energy for H\* adsorption. When H\* is adsorbed on the nitrogen atoms, the charge redistribution within the Pt–N coordination structure further decreases the Gibbs free energy for H\* adsorption,<sup>31</sup> providing more active sites for the photocatalytic hydrogen evolution reaction.

It is worth noting that the concentration of Pt single atoms also affects the final loading mass. As the size of the ice block keeps constant and is strictly determined by the stainless-steel cup, the concentration of Pt single atoms is simply modulated by changing the H<sub>2</sub>PtCl<sub>6</sub> concentration in the precursor solution. The photocatalytic hydrogen evolution of Pt/g-C<sub>3</sub>N<sub>4</sub> with different Pt loading is shown in Fig. S5.† When the H<sub>2</sub>PtCl<sub>6</sub> concentration increases from 0.1 to 1 mg mL<sup>-1</sup>, the rising photocatalytic activity is mainly due to the increased number of Pt single atoms by laser synthesis. Once the concentration exceeds 1 mg mL<sup>-1</sup>, the decreasing photocatalytic activity may be attributed to two possible reasons. One is that the excessive H<sub>2</sub>PtCl<sub>6</sub> concentration enhances the absorption of the UV laser, resulting in lower generation of Pt single atoms in the ice block. The other possibility is that an excessive number of Pt single-atoms synthesized within the ice tend to form clusters due to extensive collision during low-temperature adsorption. Further research is required to determine the exact cause in future works. Furthermore, the stability and sustainability of the optimized Pt/g-C<sub>3</sub>N<sub>4</sub> are evaluated. A prolonged photocatalytic test over 22 h displays a slight

reduction in hydrogen evolution following the initial 9 h, after which the rate stabilizes in the next 13 h (Fig. 4d). After 5 consecutive cycles, the hydrogen evolution capability only decreases by 9.6% (Fig. 4e). To analyze the mechanism for the slight decrease in catalytic performance, the Pt/g-C<sub>3</sub>N<sub>4</sub> is characterized again after five cycles. The Pt content obtained by ICP-OES is 1.15%, which remains essentially consistent with the initial Pt/g-C<sub>3</sub>N<sub>4</sub>, indicating that Pt single atoms exhibit excellent anti-leaching properties. The HAADF-STEM image (Fig. S6†) shows that the Pt remains atomically dispersed after five reaction cycles, and no obvious change is observed in the FTIR spectrum (Fig. 5a). By comparing the XPS spectra before and after the catalytic reaction (Fig. 5b–e), the fine spectra of C 1s (Fig. 5b) and O 1s (Fig. 5c) remain almost completely unchanged, while the peaks of N 1s (Fig. 5d) and Pt 4f (Fig. 5e) both shift to lower binding energies. This suggests that the Pt single atoms have gained electrons and been partially reduced during the catalytic process, leading to a decrease in valence state. Slow-scan XRD (0.5° min<sup>-1</sup>) is employed to detect whether the reduction of Pt single atoms leads to the formation of metallic Pt. As shown in Fig. 5f, the low scan rate reduces the noise in the XRD pattern, clearly indicating that no peak corresponding to metallic Pt is present. Therefore, after prolonged catalytic reactions, the Pt atoms with lower valence state remain in an isolated state. The reduction in Pt valence state is attributed to the weakening of the metal–support interaction between the Pt single atoms and the g-C<sub>3</sub>N<sub>4</sub> support,<sup>35</sup> which can cause a gradual decline in hydrogen production performance. These results indicate that although anchoring Pt single atoms on g-C<sub>3</sub>N<sub>4</sub> *via* electrostatic adsorption provides good stability, the interaction between Pt and the g-C<sub>3</sub>N<sub>4</sub> support tends to weaken over prolonged photocatalytic reaction time, a factor that should be addressed in future studies.

The improved laser synthesis method exhibits several distinct advantages over conventional laser synthesis approaches. Firstly, this method eliminates the need for high-



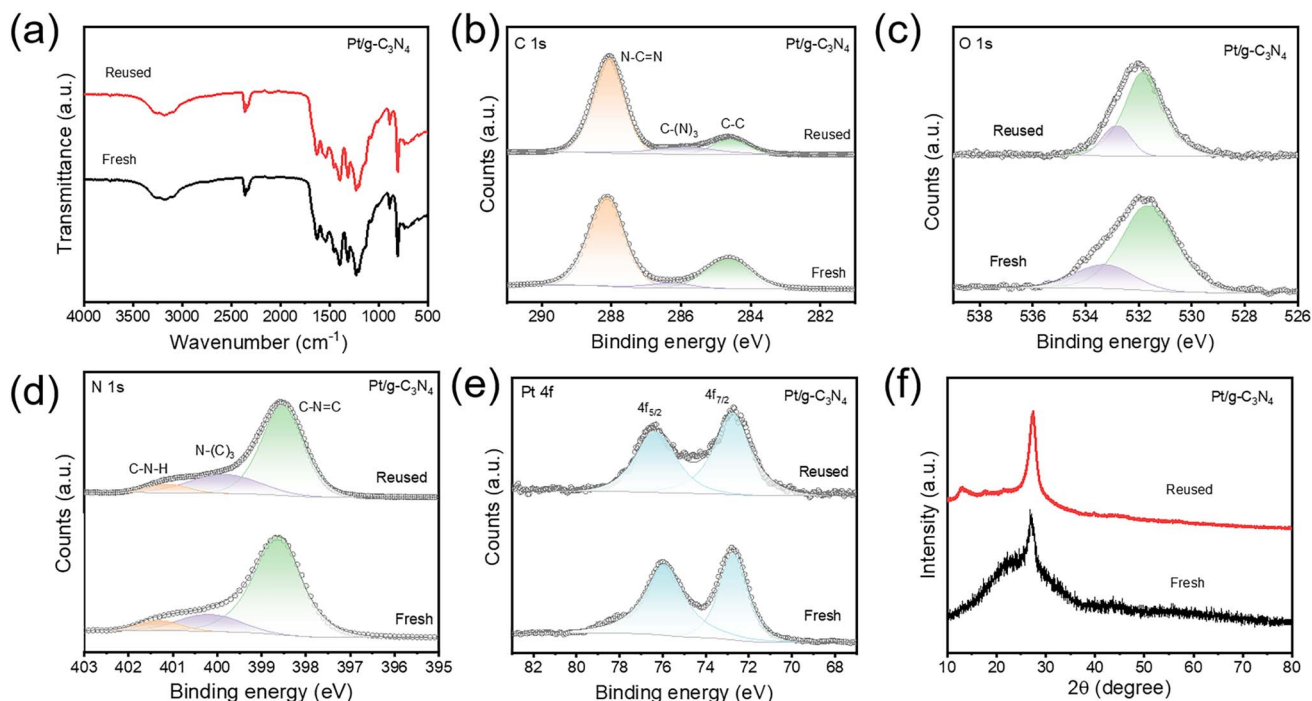


Fig. 5 The characterization of Pt/g-C<sub>3</sub>N<sub>4</sub> after five photocatalytic cycles. (a) FTIR spectra. (b–e) XPS spectra of C 1s (b), O 1s (c), N 1s (d), and Pt 4f (e). (f) XRD patterns.

energy lasers, which are typically required in traditional laser-based synthesis methods, thereby reducing equipment costs and power consumption. Secondly, this method uses water as the solvent. Compared to laser liquid-phase synthesis methods that employ organic solvents, it reduces solvent costs and the risk of environmental pollution. Thirdly, this method operates under low-temperature conditions, without the necessity for additional reductants or high-temperature treatments, aligning with green chemistry principles and minimizing energy consumption. Furthermore, this method is feasible for synthesizing highly dispersed single-atom catalysts, enhancing the utilization efficiency of noble metals. These advantages make the proposed method economically viable for large-scale production.

## 4. Conclusions

In summary, a novel synthesis strategy that involves synthesizing free-standing Pt single atoms and subsequently anchoring them on g-C<sub>3</sub>N<sub>4</sub> has been successfully demonstrated. This innovative approach significantly boosts the photocatalytic hydrogen evolution performance. The integration of the freeze-phase photochemical method with the laser reduction technique facilitates the direct reduction of Pt precursors without the need for creating defects on the support material, overcoming the limitations associated with traditional high-energy laser synthesis methods. The resulting Pt/g-C<sub>3</sub>N<sub>4</sub> SAC exhibits excellent photocatalytic activity and stability, highlighting the effectiveness of this synthesis approach in stabilizing single atoms and enhancing catalytic performance. This laser

synthesis method also demonstrates the potential for anchoring single atoms on a variety of substrates, which will be further investigated in the future. This work provides a simple and eco-friendly method for producing high-performance SACs.

## Data availability

The data that support the findings of this study are available from the corresponding author upon reasonable request.

## Conflicts of interest

The authors have no conflicts to disclose.

## Acknowledgements

This work was supported by the National Natural Science Foundation of China (Grant No. 62325505, 62020106014, 62175140, 62475138), the Shanxi Province Graduate Student Research Innovation Project (Grant No. 2024KY102), and the Xinzhou Key Research and Development Project (Grant No. 20240102).

## References

- 1 B. Qiao, A. Wang, X. Yang, L. F. Allard, Z. Jiang, Y. Cui, J. Liu, J. Li and T. Zhang, "Single-atom catalysis of CO oxidation using Pt<sub>1</sub>/FeO<sub>x</sub>", *Nat. Chem.*, 2011, 3, 634–641.
- 2 A. Wang, J. Li and T. Zhang, "Heterogeneous single-atom catalysis", *Nat. Rev. Chem.*, 2018, 2, 65–81.



- 3 X.-F. Yang, A. Wang, B. Qiao, J. Li, J. Liu and T. Zhang, "Single-Atom Catalysts: A New Frontier in Heterogeneous Catalysis", *Acc. Chem. Res.*, 2013, **46**, 1740–1748.
- 4 L. Liu and A. Corma, "Metal Catalysts for Heterogeneous Catalysis: From Single Atoms to Nanoclusters and Nanoparticles", *Chem. Rev.*, 2018, **118**, 4981–5079.
- 5 S. Ji, Y. Chen, X. Wang, Z. Zhang, D. Wang and Y. Li, "Chemical Synthesis of Single Atomic Site Catalysts", *Chem. Rev.*, 2020, **120**, 11900–11955.
- 6 K. Wang, X. Wang and X. Liang, "Synthesis of High Metal Loading Single Atom Catalysts and Exploration of the Active Center Structure", *ChemCatChem*, 2021, **13**, 28–58.
- 7 H. Wei, K. Huang, D. Wang, R. Zhang, B. Ge, J. Ma, B. Wen, S. Zhang, Q. Li, M. Lei, C. Zhang, J. Irawan, L.-M. Liu and H. Wu, "Iced photochemical reduction to synthesize atomically dispersed metals by suppressing nanocrystal growth", *Nat. Commun.*, 2017, **8**, 1490.
- 8 N. Fu, X. Liang, X. Wang, T. Gan, C. Ye, Z. Li, J.-C. Liu and Y. Li, "Controllable Conversion of Platinum Nanoparticles to Single Atoms in Pt/CeO<sub>2</sub> by Laser Ablation for Efficient CO Oxidation", *J. Am. Chem. Soc.*, 2023, **145**, 9540–9547.
- 9 K. Khan, T. Liu, M. Arif, X. Yan, M. D. Hossain, F. Rehman, S. Zhou, J. Yang, C. Sun, S.-H. Bae, J. Kim, K. Amine, X. Pan and Z. Luo, "Laser-Irradiated Holey Graphene-Supported Single-Atom Catalyst towards Hydrogen Evolution and Oxygen Reduction", *Adv. Energy Mater.*, 2021, **11**, 2101619.
- 10 Y. Peng, J. Cao, Y. Sha, W. Yang, L. Li and Z. Liu, "Laser solid-phase synthesis of single-atom catalysts", *Light:Sci. Appl.*, 2021, **10**, 168.
- 11 H. Yuan, D. Jiang, Z. Li, X. Liu, Z. Tang, X. Zhang, L. Zhao, M. Huang, H. Liu, K. Song and W. Zhou, "Laser Synthesis of PtMo Single-Atom Alloy Electrode for Ultralow Voltage Hydrogen Generation", *Adv. Mater.*, 2024, **36**, 2305375.
- 12 S. Zhang, Q. Jiang, T. Shi, Q. Sun, Y. Ye, Y. Lin, L. R. Zheng, G. Wang, C. Liang, H. Zhang and H. Zhao, "Laser Irradiation in Liquid to Release Cobalt Single-Atom Sites for Efficient Electrocatalytic N<sub>2</sub> Reduction", *ACS Appl. Energy Mater.*, 2020, **3**, 6079–6086.
- 13 X. Z. Wang, X. Pei, W. Liu, Y. Leng, X. Yu, C. Wang, L. Hu, Q. Su, C. Wu, Y. Yao, Z. Lin and Z. Zou, "Room-Temperature Laser Planting of High-Loading Single-Atom Catalysts for High-Efficiency Electrocatalytic Hydrogen Evolution", *J. Am. Chem. Soc.*, 2023, **145**, 13788–13795.
- 14 H. Miao, W. Zhang, T. Wang, Z. Yang and C. Kong, "g-C<sub>3</sub>N<sub>4</sub>-based nanocomposites for the photocatalytic degradation of VOCs: A review", *Prog. Nat. Sci.:Mater. Int.*, 2023, **33**, 407–424.
- 15 L. M. Oanh, L. T. Hang, N. D. Lai, N. T. Phuong, D. V. Thang, N. M. Hung, D. D. Bich and N. V. Minh, "Influence of annealing temperature on physical properties and photocatalytic ability of g-C<sub>3</sub>N<sub>4</sub> nanosheets synthesized through urea polymerization in Ar atmosphere", *Phys. B*, 2018, **532**, 48–53.
- 16 P. Wu, T. Zhou, Z. Tong, F. Xi, J. Lu, X. Li, W. Ma, S. Li and X. Yang, "Balancing charge carrier density and exciton recombination in defective g-C<sub>3</sub>N<sub>4</sub> for efficient photocatalytic hydrogen evolution", *Prog. Nat. Sci.:Mater. Int.*, 2025, **35**, 238–244.
- 17 Q. Zhang, M. Yue, P. Chen, Q. Ren, W. Kong, C. Jia, Q. Lu, J. Wu, Y. Li, W. Liu, P. Li, Y. Fu and J. Ma, "Accelerating photocatalytic hydrogen production by anchoring Pt single atoms on few-layer g-C<sub>3</sub>N<sub>4</sub> nanosheets with Pt-N coordination", *J. Mater. Chem. C*, 2024, **12**, 3437–3449.
- 18 P. Zhou, F. Lv, N. Li, Y. L. Zhang, Z. J. Mu, Y. H. Tang, J. P. Lai, Y. G. Chao, M. C. Luo, F. Lin, J. H. Zhou, D. Su and S. J. Guo, "Strengthening reactive metal-support interaction to stabilize high-density Pt single atoms on electron-deficient g-C<sub>3</sub>N<sub>4</sub> for boosting photocatalytic H<sub>2</sub> production", *Nano Energy*, 2019, **56**, 127–137.
- 19 Y. Jiang, Z. Sun, C. Tang, Y. Zhou, L. Zeng and L. Huang, "Enhancement of photocatalytic hydrogen evolution activity of porous oxygen doped g-C<sub>3</sub>N<sub>4</sub> with nitrogen defects induced by changing electron transition", *Appl. Catal., B*, 2019, **240**, 30–38.
- 20 H. Yang, N. Lu, J. Zhang, R. Wang, S. Tian, M. Wang, Z. Wang, K. Tao, F. Ma and S. Peng, "Ultra-low single-atom Pt on g-C<sub>3</sub>N<sub>4</sub> for electrochemical hydrogen peroxide production", *Carbon Energy*, 2023, **5**, e337.
- 21 M. Ou, S. Wan, Q. Zhong, S. Zhang and Y. Wang, "Single Pt atoms deposition on g-C<sub>3</sub>N<sub>4</sub> nanosheets for photocatalytic H<sub>2</sub> evolution or NO oxidation under visible light", *Int. J. Hydrogen Energy*, 2017, **42**, 27043–27054.
- 22 A. Sharma, M. Varshney, K. H. Chae and S. O. Won, "Mechanistic investigations on emission characteristics from g-C<sub>3</sub>N<sub>4</sub>, g-C<sub>3</sub>N<sub>4</sub>@Pt and g-C<sub>3</sub>N<sub>4</sub>@Ag nanostructures using X-ray absorption spectroscopy", *Curr. Appl. Phys.*, 2018, **18**, 1458–1464.
- 23 Y. Yuan, L. Zhang, J. Xing, M. B. Utama, X. Lu, K. Du, Y. Li, X. Hu, S. Wang, A. Genc, R. Dunin-Borkowski, J. Arbiol and Q. Xiong, "High-yield synthesis and optical properties of g-C<sub>3</sub>N<sub>4</sub>", *Nanoscale*, 2015, **7**, 12343–12350.
- 24 Y. Li, Z. He, L. Liu, Y. Jiang, W.-J. Ong, Y. Duan, W. Ho and F. Dong, "Inside-and-out modification of graphitic carbon nitride (g-C<sub>3</sub>N<sub>4</sub>) photocatalysts via defect engineering for energy and environmental science", *Nano Energy*, 2023, **105**, 108032.
- 25 X. Yang, J. Qin, Z. Dai, Y. Sun, H. Liu, X. Zheng and Z. Hu, "MOF-derived Fe based catalysts for efficiently Advanced Oxidation Processes: From single atoms to diatomic and nanoparticles", *Prog. Nat. Sci.:Mater. Int.*, 2023, **33**, 534–543.
- 26 Y. Li and A. I. Frenkel, "Deciphering the Local Environment of Single-Atom Catalysts with X-ray Absorption Spectroscopy", *Acc. Chem. Res.*, 2021, **54**, 2660–2669.
- 27 T. Lan, R. Yalavarthi, Y. Shen, M. Gao, F. Wang, Q. Hu, P. Hu, M. Beladi-Mousavi, X. Chen, X. Hu, H. Yang, E. Cortes and D. Zhang, "Polyoxometalates-Mediated Selectivity in Pt Single-Atoms on Ceria for Environmental Catalysis", *Angew. Chem., Int. Ed.*, 2025, **64**, e202415786.
- 28 S. Hu, P. Qiao, X. Liang, G. Ba, X. Zu, H. Hu, J. Ye and D. Wang, "Single-atom Pt–N<sub>4</sub> active sites anchored on porous C<sub>3</sub>N<sub>4</sub> nanosheet for boosting the photocatalytic CO<sub>2</sub> reduction with nearly 100% CO selectivity", *Appl. Catal., B*, 2024, **364**, 123737.
- 29 T. Mahvelati-Shamsabadi, K. C. Bhamu, S. Lee, T. T. Dang, V. H. Khoi, S. H. Hur, W. M. Choi, S. G. Kang, T. J. Shin



- and J. S. Chung, "Coordinatively unsaturated atomically dispersed  $\text{Pt}^{+2}\text{-N}_4$  sites on hexagonal nanosheet structure of  $\text{g-C}_3\text{N}_4$  for high-performance photocatalytic  $\text{H}_2$  production", *Appl. Catal., B*, 2023, **337**, 122959.
- 30 C. Yang, Z. Zhao and Q. Liu, "Mechanistic insight into the dispersion behavior of single platinum atom on monolayer  $\text{g-C}_3\text{N}_4$  in single-atom catalysts from density functional theory calculations", *Appl. Surf. Sci.*, 2021, **556**, 150697.
- 31 C. Yang, Z. Zhao and Q. Liu, "Regulating effect on photocatalytic water splitting performance of  $\text{g-C}_3\text{N}_4$  via confinement of single atom Pt based on energy band engineering: A first principles investigation", *Appl. Surf. Sci.*, 2022, **577**, 151916.
- 32 C. Yang, Z.-Y. Zhao, H.-T. Wei, X.-Y. Deng and Q.-J. Liu, "DFT calculations for single-atom confinement effects of noble metals on monolayer  $\text{g-C}_3\text{N}_4$  for photocatalytic applications", *RSC Adv.*, 2021, **11**, 4276–4285.
- 33 K. L. Zhou, Z. Wang, C. B. Han, X. Ke, C. Wang, Y. Jin, Q. Zhang, J. Liu, H. Wang and H. Yan, "Platinum single-atom catalyst coupled with transition metal/metal oxide heterostructure for accelerating alkaline hydrogen evolution reaction", *Nat. Commun.*, 2021, **12**, 3783.
- 34 Z. Zhang, Z. Zhang, C. Chen, R. Wang, M. Xie, S. Wan, R. Zhang, L. Cong, H. Lu, Y. Han, W. Xing, Z. Shi and S. Feng, "Single-atom platinum with asymmetric coordination environment on fully conjugated covalent organic framework for efficient electrocatalysis", *Nat. Commun.*, 2024, **15**, 2556.
- 35 F. Maurer, J. Jelic, J. Wang, A. Ganzler, P. Dolcet, C. Woll, Y. Wang, F. Studt, M. Casapu and J.-D. Grunwaldt, "Tracking the formation, fate and consequence for catalytic activity of Pt single sites on  $\text{CeO}_2$ ", *Nat. Catal.*, 2020, **3**, 824–833.

