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Induced abundant oxygen vacancies in $\text{Sc}_2\text{VO}_{5-\delta}/\text{g-C}_3\text{N}_4$ heterojunctions for enhanced photocatalytic degradation of levofloxacin†

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$\text{Sc}_2\text{VO}_{5-\delta}/\text{g-C}_3\text{N}_4$ heterojunctions (SVCs) with abundant oxygen vacancies (OVs) were synthesized by ultrasonic exfoliation combined with the thermal etching method. The structures, OVs and spatial separation of the photogenerated carriers were systematically characterized. The results manifested that the SVCs were successfully constructed via the strong interaction between $\text{g-C}_3\text{N}_4$ (CN) and $\text{Sc}_2\text{VO}_{5-\delta}$ (SV). The SVCs possessed a higher concentration of OVs than that of pristine CN and SV. The formation of the SVC heterostructures and the optimization of the OVs were the two major factors to accelerate the separation of the charge carriers and finally to improve the photocatalysis performance. The as-prepared 10%SVC (containing 10 wt% of SV) catalyst exhibited the highest OV concentration and the best photocatalytic performance. The levofloxacin (LVX) photodegradation activity showed a positive correlation with the OV concentration. The photocatalytic degradation efficiencies were 89.1, 98.8 and 99.0% on 10%SVC for LVX, methylene blue (MB) and rhodamine B (RhB), respectively. These photodegradation processes followed the pseudo first order kinetic equation. The apparent rate constant (k_{app}) of LVX degradation on 10%SVC was 11.0 and 7.5 times that of CN and SV. The h^+ , $\cdot\text{OH}$ and $\cdot\text{O}_2^-$ were the major reactive species in the photodegradation process.

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1. Introduction

The development of modern industry and the consumption of fossil fuel have caused serious environmental pollution, which is severely harmful to human health and the sustainable development of society.¹ Traditional techniques have been utilized to remove organic pollutants, including dyes and antibiotics.^{2–4} However, these traditional methods suffer from drawbacks, such as not being suitable for low concentration pollutants, high operation costs, low removal efficiency and secondary pollution.

In the recent few decades, semiconductor based photocatalysis has been regarded as the most promising technology for the elimination of low concentration contaminants.^{5–7} The photocatalytic technology has mild reaction conditions, low cost, superior recoverability and less secondary pollution and *etc.* Therefore, photocatalysis technology has been extensively investigated for the removal of dyes and antibiotics.^{8–11} The significant progress has been made in this field recently.¹² Various semiconductor materials with superior photocatalytic

performance in wastewater treatment have aroused extensively research interests.^{13–15} Nevertheless, the low quantum efficiency and the narrow visible light response range have always been the major obstacles for the practical application of the photocatalytic technology.

Rare earth vanadates are a type of semiconductor materials with novel magnetic, optical, electronic and catalytic performances,¹⁶ which have been widely applied as laser hosts, phosphors, polarizers and catalysts.¹⁷ Especially, the electron on 3d orbital of V can be excited by the visible light.¹⁸ Rare earth vanadates are consequently considered as the promising candidates for highly efficient visible-light photocatalysts. Several rare earth vanadates, including CeVO_4 ,¹⁹ DyVO_4 ,²⁰ EuVO_4 ,²¹ GdVO_4 ,¹⁷ HoVO_4 ,²² LaVO_4 ,¹⁸ NdVO_4 ,²³ PrVO_4 ,²⁴ SmVO_4 ,²⁵ TmVO_4 ,²¹ YVO_4 ,²⁶ were successfully prepared and the visible light driven photocatalytic activities were investigated. Nevertheless, the low visible light utilization and the high charge recombination are still the key factors of low photocatalytic efficiency of rare earth vanadates.²⁷ Doping,²⁸ metal-substitution,²⁹ morphology control,¹⁹ introduction of surface OVs^{30,31} and construction of heterojunction^{18,24,25,27} have been adopted to meliorate its photocatalytic quantum efficiency. Among them, surface OVs engineering can effectively regulate the electronic structure and charge migration, produce the active surface phases, and ameliorate the adsorption of the contaminants and intermediates in the photocatalytic

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degradation process.³² As a contrast, the bulk OVVs are the deep charge traps, which can expedite the recombination of the photogenerated electrons and holes. The photocatalytic performance is hence reduced by the bulk OVVs. Accurate introduction of surface OVVs is crucial to improve the photocatalytic activity. According to previous literature, the control of synthesis temperature, substitution, doping, chemical etching and surface reduction are the effectual approaches to tune the OVVs of the photocatalyst.³³ Thereinto, the control of calcination temperature is a convenient method to engineer the OVVs concentration on the surface or at the interface of the binary or ternary oxides catalysts during the post treatment process. It is believed that the calcination temperature influences the crystallinity, morphology, surface area and OVVs concentration. In general, the calcination temperature range between 350 to 500 °C is advantageous to achieve the optimal surface OVVs concentration of the oxides.³³

The construction of heterojunction is another feasible and effective way to facilitate the spatial separation and the recombination rate of photogenerated charge carriers.^{34–37} Many rare earth vanadates based heterojunctions have been established to promote its photocatalytic degradation of contaminants. For instance, LaVO₄/g-C₃N₄ heterojunction was fabricated and the visible light driven photocatalytic performance was assessed. 99% of trichloroethylene was mineralized on 3% LaVO₄/g-C₃N₄, which were 3.89 and 4.67 times of LaVO₄ and g-C₃N₄, respectively.¹⁸ In another work, the integration of UiO-66-NH₂ with SmVO₄ enhanced the light absorption, carrier lifetime and specific surface area of the composite. SmVO₄/UiO-66-NH₂ eliminated 50% of tetracycline in 90 min under solar light.²⁵ La₂O₃/CeVO₄@halloysite nanotubes composites also successfully fabricated by hydrothermal method. The construction of heterojunction between La₂O₃ and CeVO₄ inhibited the recombination of photogenerated charge carriers.²⁷ 87.1% of tetracycline in 60 min under visible light irradiation on the optimized composite, which was 3.89 times higher than that of raw CeVO₄. LaVO₄/BiOBr heterojunction was also prepared by hydrothermal method. The introduction of LaVO₄ heightened the solar energy harvesting and reduced the transfer resistance of the photogenerated charge carriers. 95.4% and 87.1% of acetone and toluene degraded under visible light irradiation over 3% LaVO₄/BiOBr, which were 2.6 and 5.3 times higher than that of pristine BiOBr.³⁸ As mentioned above, the spatial separation of the photogenerated charge and the photocatalytic activity can be effectively enhanced through constructing rare earth vanadates based heterojunction.

In this study, Sc₂VO_{5-δ} with abundant OVVs was loaded on the ultrathin g-C₃N₄ nanosheets to form the Sc₂VO_{5-δ}/g-C₃N₄ heterojunctions. Transmission electron microscopy (TEM), high resolution TEM (HRTEM), scanning electron microscopy (SEM), Fourier-transform infrared spectra (FTIR) and X-ray photoelectron spectroscopy (XPS) results clarified the successful formation of SVCs and its ultrathin structure. Ultraviolet-visible diffuse reflectance spectra (DRS) and XPS VB spectra revealed it was type II heterojunction. Electron paramagnetic resonance (EPR) confirmed the existence of OVVs and XPS O1s spectra determined the OVVs concentration. Transient photocurrent

response (TPC), electrochemical impedance spectroscopy (EIS) and photoluminescence spectra (PL) assessed the promotion of the separation efficiency and the inhibition of the recombination of the photogenerated charge carriers in SVCs. The influence of OVVs and SV content on the photocatalytic degradation efficiency of LVX, MB and RhB was investigated. Due to the introduction of surface OVVs and the formation of heterojunction, the as-prepared SVCs manifested highly enhanced photocatalytic degradation efficiency. This study supplied a new paradigm for further exploring the synergistic effect of surface OVVs and heterostructure on the photocatalytic activity.

2. Experiment

2.1 Synthesis of SVCs

2.1.1 Preparation of CN nanosheets. 10 g dicyandiamine and 10 g NH₄Cl was firstly mixed and annealed for 4 h at 550 °C. The obtained yellow CN solid was ground. 100 mg bulk CN powder dispersed in 200 mL ultrapure water and followed 4 h of ultrasonic treatment. The precipitation was discarded by centrifugation. The supernatant was evaporated by lyophilization to acquire light yellow CN nanosheets.

2.1.2 Preparation of SVCs. For the preparation of 10%SVC, 200 mg CN nanosheets, 34.0 mg Sc(NO₃)₃·6H₂O and 30.5 mg NH₄VO₃ were dispersed in 100 mL ultrapure water and continuously stirred for 4 h. CN nanosheets adsorbing Sc and V ions was obtained after the water was evaporated. Then, the powder was calcined at 500 °C for 2 h. After cooling down to room temperature, the product was collected and ground for the following photocatalytic degradation reaction. 2%SVC, 5% SVC and 20%SVC, containing 2, 5 and 20 wt% of SV respectively, were prepared under the same reaction conditions. In order to investigate the effect of calcination temperature on the OVVs, SV was synthesized at 400, 500 and 600 °C for 4 h and named as SV40, SV50 and SV60. 10%SVC was prepared at 400, 450, 500 and 550 °C for 2 h and designated as 10%SVC40, 10%SVC45, 10%SVC50, 10%SVC55, respectively.

2.2 Characterization

X-ray diffraction (XRD) patterns were tested on Rigaku SmartLab diffractometer. FTIR spectra were obtained on Nicolet NEXUS 470 spectrometer. TEM and HRTEM images were recorded on JEOL-2100F transmission electron microscope. SEM, energy dispersion spectrum (EDS) and elemental mapping images were measured on JSM-4800F scanning electron microscope. XPS spectra were measured on Thermo ESCALAB 250XI system. EPR spectra were recorded on Bruker E500 spectrometer. The N₂ adsorption-desorption isotherms and BET specific surface area were analyzed by ASAP 2460 Micrometrics instrument. DRS spectra were determined on Shimadzu UV-2401 spectrophotometer. UV-vis spectra were conducted on Varian Cary 50 UV-vis spectrophotometer.

2.3 Photoelectrochemical measurement

The photoelectrochemical experiments of CN, SVCs and SV were implemented on CHI 660E electrochemical workstation. Pt and



Ag/AgCl were utilized as the counter electrode and reference electrode. For the preparation of working electrode, 5 mg CN, SVCs or SV was mixed firstly with 0.2 mL Nafion in 1.8 mL ethanol by sonication and then dropped on 1 cm² of FTO glass. TPC and EIS of the catalysts were detected in 0.2 M Na₂SO₄ electrolyte solution.

2.4 Photocatalytic degradation reaction

100 mg of CN, SVCs or SV was dispersed in 200 mL of 12.0 mg L⁻¹ LVX, 5.0 mg L⁻¹ MB or 5.0 mg L⁻¹ RhB, respectively. The solution was continuously stirred in the dark for 30 min to reach adsorption desorption equilibrium of the contaminants on the catalysts. The photocatalytic degradation of contaminants was evaluated under the irradiation of 40 W white LED. The average intensity of the visible light was about 20 mW cm⁻² (CEL-NP2000-2, CEALIGHT, Beijing). 5 mL solution was fetched out at intervals and the catalyst was removed through 0.22 μm PTFE filter membrane. The contaminant concentration in degradation solution was detected by UV-vis spectrophotometer at 294 nm (LVX), 664 nm (MB) and 552 nm (RhB). The temperature was fixed at 25 °C in the photocatalysis process. The degradation experiments were repeated three times. The apparent quantum yield of LVX degradation was estimated according to the literature.³⁹ The stability of 10%SVC was appraised by 4 cycles of photocatalytic degradation experiments. 10%SVC was centrifuged and desorbed in deionized water for several times after each cycle experiment to eliminate absorbed contaminant. Then, 10%SVC was separated and freeze-dried for the usage in next cycle experiment. The reproducibility of 10%SVC was evaluated by repeating the above process twice. NaNO₃, EDTA, isopropyl alcohol (IPA) and *p*-benzoquinone (pBQ) were selected as the scavengers to trap e⁻, h⁺, ·OH and ·O₂⁻, respectively.

3. Results and discussion

The XRD patterns of CN, SVCs and SV were shown in Fig. 1. Two characteristic diffraction peaks at 13.0° and 27.6° were clearly observed in the XRD patterns of CN and SVCs. It respectively corresponded to (100) and (002) crystal planes of CN, suggesting the existence of CN in the as-prepared SVCs. The peaks at 13.0° originated from the in-plane repeating structural units and the peaks at 27.6° were attributed to the inter-planar structure stacking of aromatic systems.⁴⁰ The broad feature of the peaks indicated low crystallinity of the samples. All diffraction peaks of SV were in accord well with tetragonal phase Sc₂VO₅ (PDF#28-1014). The main peaks at 21.6°, 26.4°, 31.6°, 36.5°, 41.0°, 44.4°, 46.9°, 48.6°, 52.6°, 52.9°, 62.6° and 63.2° could be clearly observed, which was respectively correspond to (112), (201), (203), (220), (223), (224), (320), (322), (400) and (217) crystal plane of tetragonal phase. The diffraction peaks of SV could not be detected in 2%SVC and 5%SVC. It was mainly due to the relative lower dosage of SV in SVCs, which could not initiate the chemical skeleton and structure changes of CN. It was similar to the previous result.⁴¹ The diffraction peak at 31.6° of SV was observed in 10%SVC and 20%SVC. The weak and broad

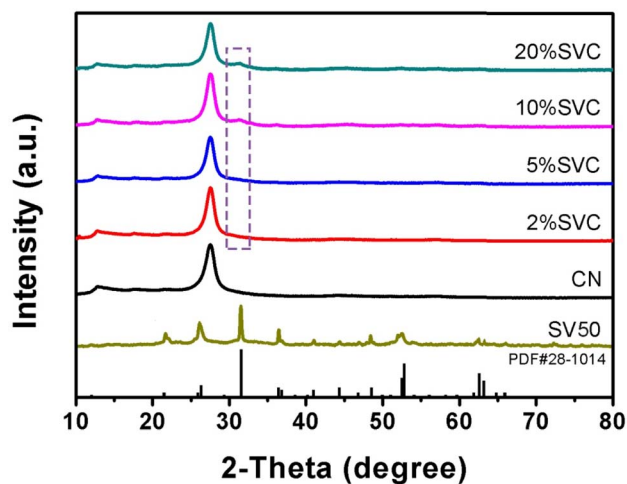


Fig. 1 XRD patterns of CN, 2%SVC, 5%SVC, 10%SVC, 20%SVC and SV.

diffraction peaks at 31.6° of 10%SVC and 20%SVC demonstrated the low crystallinity of SV in SVCs. Hence, the XRD result confirmed the successful preparation of SVCs with certain of amorphous phase.

As reported in the literature, the amorphous phase in the catalyst can significantly influence its crystallinity and further affect its surface OVs.³³ Actually, exorbitant crystallinity resulted in lower concentration of surface OVs, which was not conducive to the separation of photogenerated charge. Conversely, excessive low crystallinity caused more bulk OVs, which was also detrimental to the improvement of photocatalytic performance. In general, the calcination temperature range between 350 to 500 °C is advantageous to achieve the optimal surface OVs concentration. Thus, the appropriate crystallinity of SVCs could be conveniently generated by optimizing the calcination temperature. And then the optimal surface OVs concentration was achieved by regulating the crystallinity of the catalyst. To corroborate the influence of calcination temperature on crystallinity and OVs concentration of the catalyst, 10%SVC was synthesized at 400, 450, 500 and 550 °C respectively. The crystallinity was investigated by XRD patterns and the surface OVs were detected by XPS O1s spectra. As the XRD results depicted in Fig. S1,[†] the diffraction peak at 31.6° appeared stronger and sharper with the increase of the calcination temperature, indicating the crystallinity of 10%SVC was gradually ameliorated.

The morphologies of bulk CN, CN nanosheets and SVCs were determined by SEM. As displayed in Fig. 2a, the bulk CN exhibited blocky aggregate structure, in which lamellar structure of CN nanosheets could not be observed. The characteristic layered structure of CN was clearly revealed in Fig. 2b, indicating that CN nanosheets could effectively be generated by the exfoliation effect in the ultrasonic treatment and the thermal etching in the reheating synthetic procedure. As shown in Fig. 2c and d, 10%SVC had the structural feature of CN nanosheets. The SEM result clearly revealed that SV nanoparticles were located on CN nanosheets (Fig. 2c). 2%SVC, 5%SVC and 20%SVC also presented the same layered stacking structure (Fig. S2[†]). The thin-layer structure of SVCs could increase the



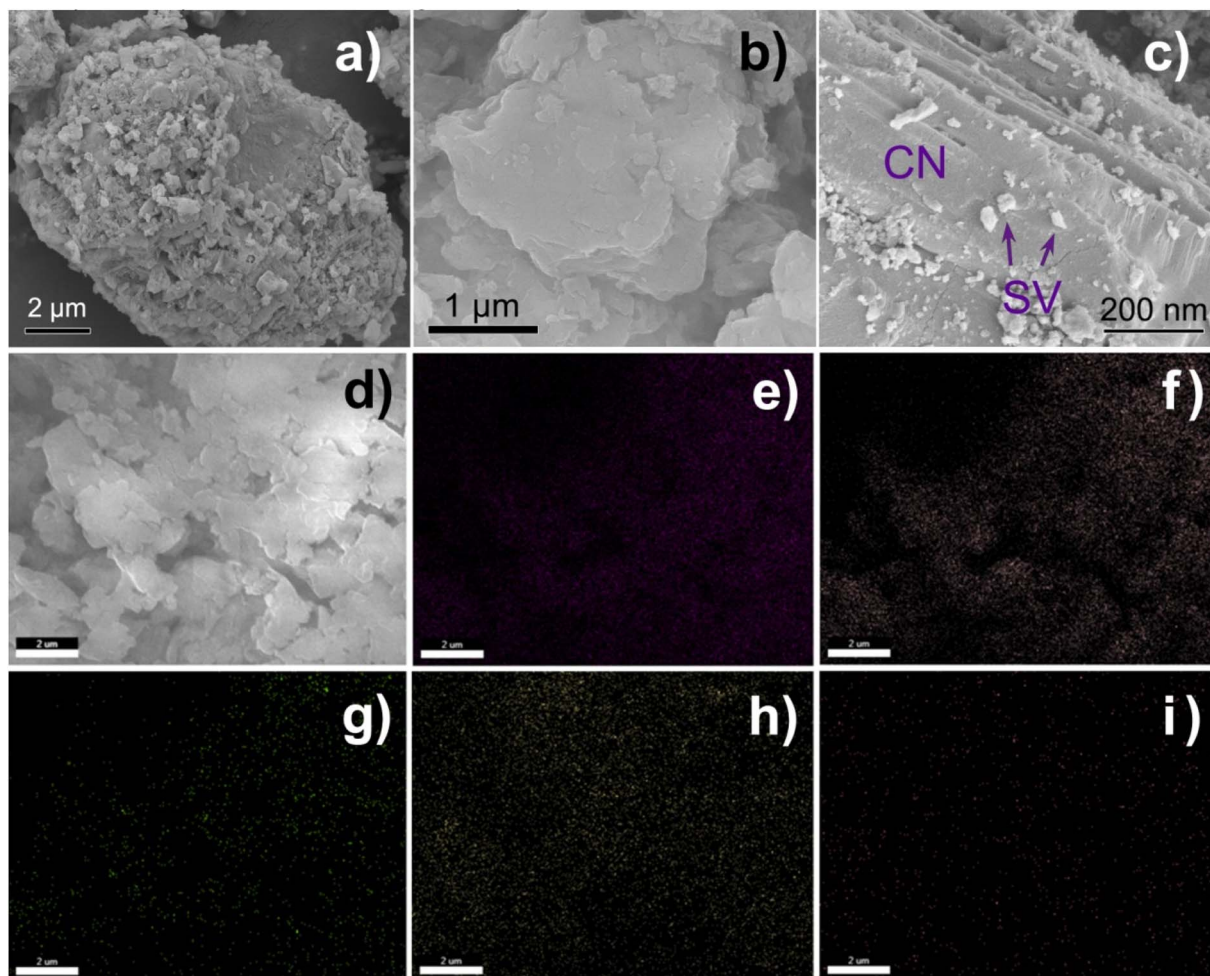


Fig. 2 SEM images of (a) bulk CN, (b) CN nanosheets and (c), (d) 10%SVC, and the element mapping of (e) C, (f) N, (g) O, (h) Sc and (i) V. Scale bar = 2 μm .

specific surface area and reaction active sites, which was instrumental in accelerating photocatalytic degradation reaction. The EDS spectra and elemental mapping techniques were employed to appraise the chemical composition and the element distribution of SVCs. 10%SVC was selected as the target heterojunction. C, N, O, Sc and V were identified by EDS in selected region (Fig. S3[†]). The corresponding weight ratio of C, N, O, Sc and V in 10%SVC was listed in Fig. S3[†]. 9.54% of SV mass percentage in 10%SVC was calculated from the EDS result. It was consistent with the dosage of $\text{Sc}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$ and NH_4VO_3 in the precursor. The distribution of C, N, O, Sc and V in 10%SVC was assessed by the elemental mappings shown in Fig. 2e–i. The uniformity of the element mapping images suggested that Sc, V and O were distributed evenly in 10%SVC. This result demonstrated that SV was loaded homogeneously on CN nanosheets.

The morphology and structure of CN, SVCs and SV were further characterized by TEM and HRTEM. The results were displayed in Fig. 3 and S4[†]. The ultrathin layered structures of CN nanosheets with wrinkles in 10%SVC were distinctly observed in Fig. 3a and b. SV exhibited irregular granular

morphology with a certain degree of aggregation. The particle size of SV was about 25 to 70 nm estimated from the TEM images. SV nanoparticles were loaded on CN nanosheets. TEM images evidently clarified that the intimate contact between CN nanosheets and SV nanoparticles, meaning the successful formation of SVCs heterostructure. The lattice fringes of SV in 10%SVC were definitely displayed in HRTEM (Fig. 3c). The d-spacing of 0.28 nm was indexed to the (203) lattice planes of tetragonal SV, which was in accordance with the XRD result discussed in Fig. 1. The similar layered structure of CN and irregular granular morphology of SV in 2%SVC, 5%SVC and 20%SVC could also be observed in Fig. S4[†]. These TEM results sufficiently confirmed the formation of SVC heterojunction.

FTIR and Raman spectra were conducted to investigate the chemical structure of as-prepared catalysts. Fig. 4a manifested the FTIR spectra of CN nanosheets and SVCs. Four typical adsorption bands were observed in all spectra. The adsorption peaks at 806 and 886 cm^{-1} were assigned to breathing vibration and N–H deformation vibration mode of CN s-triazine, respectively.⁴² A characteristic adsorption band in the range of 1200–1700 cm^{-1} was ascribed to the stretching vibrations of C–N and



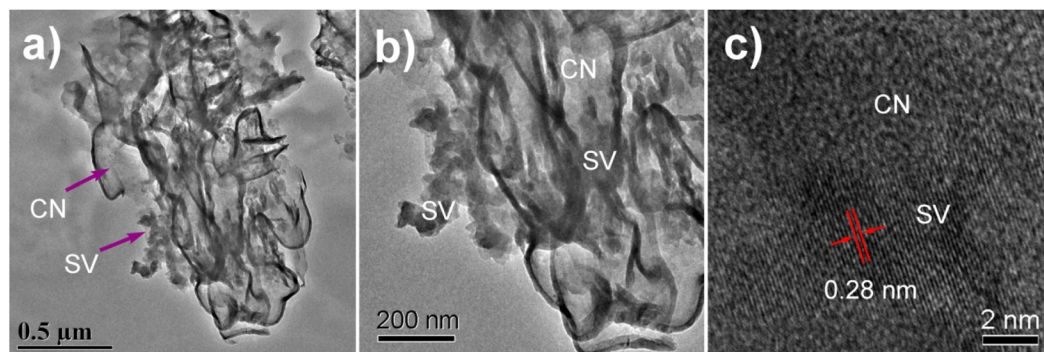


Fig. 3 (a), (b) TEM and (c) HRTEM images of 10%SVC.

C=N of aromatic CN heterocycles.⁴³ A weak adsorption band at 2174 cm^{-1} exhibited in 5%SVC, 10%SVC and 20%SVC, which was originated from the terminal cyano groups. Similar adsorption band generally could be found in many CN-based composites.⁴⁴ A broad adsorption band at $2900\text{--}3500\text{ cm}^{-1}$ was attributed to N-H stretching vibration of residual unpolymerized amino groups and O-H stretching related to physically adsorbed water molecule.⁴⁴ All the peak positions of 2%SVC, 5% SVC, 10%SVC and 20%SVC in the FTIR spectra did not shift compared with CN. It was consistent with the XRD results, suggesting that the formation of SVCs heterojunction did not vary its chemical skeleton and structure of CN nanosheets.

The Raman spectra of CN nanosheets and SVCs were depicted in Fig. 4b. The characteristic peaks of CN were presented at $486, 588, 704, 749, 976, 1108, 1250, 1315, 1480, 1565$ and 1630 cm^{-1} . It was consistent with the Raman results reported in the literature.^{43,45,46} In detail, the peak at 486 cm^{-1} was originated from the heptazine ring breathing mode,⁴⁶ and the 588 cm^{-1} band caused by the ring bending vibration.⁴³ The peaks centered at $704, 749$ and 976 cm^{-1} were respectively assigned to in plane bending vibration,⁴⁵ out-of-plane bending vibration of graphitic domain and the breathing mode of triazine ring.⁴⁷ Peaks around 1250 and 1630 cm^{-1} were corresponded to the C=N stretching vibration mode of triazine ring.⁴³ The $1315, 1480$ and 1565 cm^{-1} bands were distributed to

semi-circle stretching,⁴³ the bending vibration of $-\text{NH}_2$ group and the graphitic G band.⁴⁸ As shown in Fig. 4b, the Raman peaks of CN at 749 and 1630 cm^{-1} were shifted to 762 and 1614 cm^{-1} after the SVC heterojunction was formed. Meanwhile, the intensity of 1225 cm^{-1} peak was gradually strengthened with the increase of SV content in SVCs. These results clarified that the intimate contact between CN nanosheets and SV nanoparticles. It was agreed with the TEM and HRTEM results and signified the successful formation of SVCs heterostructure, which was advantageous to the spatial separation of the photogenerated electrons and holes during the photocatalytic degradation reaction.⁴⁹

The surface chemical composition and chemical state of CN, 10%SVC and SV were analyzed by XPS spectra. As illustrated in Fig. 5a, the XPS survey spectrum of 10%SVC revealed the existence of C, N, O, Sc and V, further demonstrating the successful formation of SVC heterojunction. In Fig. 5b, four peaks at the binding energy of $284.8, 285.8, 288.3$ and 288.5 eV were observed in the high resolution C1s XPS spectrum of pristine CN. These peaks were respectively designated to the graphitic C-C bond of polluted carbon, C-OH, C=N and O=C-O groups.⁵⁰ The C1s peaks shifted to $284.9, 286.2, 288.2$ and 288.7 eV after the formation of SVC heterojunction. The N1s peaks of pristine CN located at $398.8, 399.3, 400.9$ and 404.7 eV (Fig. 5c), corresponding to the pyridinic nitrogen (C=N-C),

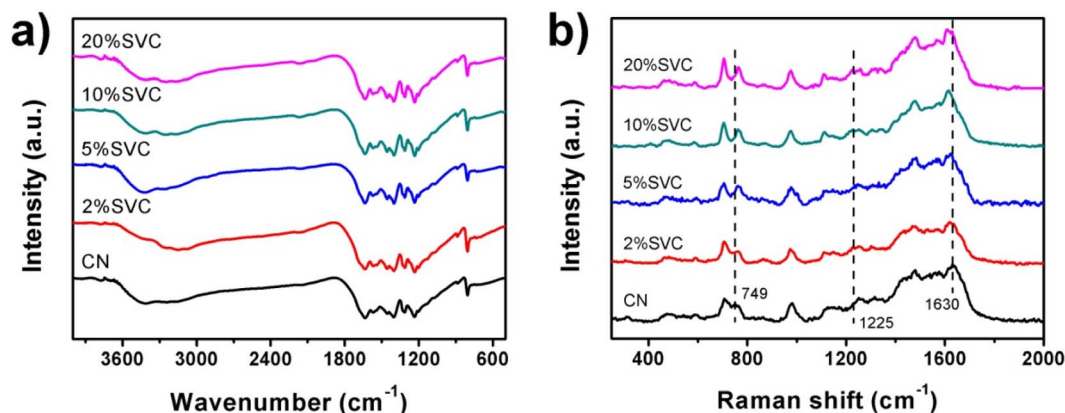


Fig. 4 (a) FTIR and (b) Raman spectra of CN, 2%SVC, 5%SVC, 10%SVC and 20%SVC.



pyrrolic nitrogen ($N-(C)_3$), surface amino nitrogen ($C-NH_2$) and the charge effect in the heterocycles.⁵¹ In contrast with pristine CN, the N1s peaks of 10%SVC were located at 398.7, 399.4, 400.8 and 404.8 eV. The variations of the binding energy implied the change of electron cloud density of C and N atoms.⁴¹ It might be attributed to the strong interaction between SV and CN. These XPS results supplied the convincing evidence to corroborate the formation of SVC heterojunctions. The Sc2p XPS spectra of SV and 10%SVC supported this verdict. As displayed in Fig. S5,[†] pure SV had the peaks at 398.4 and 403.1 eV relating to Sc2p_{3/2} and Sc2p_{1/2}, which could originate from the Sc–O bond.^{52,53} These two deconvoluted peaks shifted to 398.0 and 402.6 eV for 10%SVC, respectively. The chemical shift towards lower binding energy was observed after CN was incorporated with SV. The increase of the electron cloud density of Sc might be the reason for this reduction of the binding energy.⁵⁴ This result further indicated the strong interaction between SV nanoparticles and CN nanosheets.

The O1s spectra of 10%SVC and SV were depicted in Fig. 5d. For SV, three deconvoluted peaks at 529.8, 530.6 and 531.8 eV were obtained, respectively attributing to the lattice oxygen (OL), surface oxygen vacancies (OV) and surface adsorption oxygen (OA). The peak area percentage of OL, OV and the ratio of OV/OL in SVCs and SV samples were estimated and listed in the Table S1.[†] The result distinctly revealed that the surface OVs concentration in 10%SVC was higher than other SVCs and pure

SV. The peak intensity of 10%SVC at 529.8 eV related to the lattice oxygen was lower than that of SV. It was presumably resulted from the partial coverage of SV by CN nanosheets, which also led to the decrease of the XRD diffraction peak intensity of SV in 2%SVC and 5%SVC.⁴¹ Furthermore, the O1s binding energies of SV40 and SV50 were lower than that of values reported previously, which might be associated with its lower crystallinity. In order to investigate the effect of calcination temperature on the surface OVs, the precursors were calcinated at 400, 500 and 600 °C for 4 h to synthesize SV. The corresponding O1s spectra of SV40, SV50 and SV60 were comparatively exhibited in Fig. S6.[†] The deconvoluted peaks of SV60 located at the binding energy of 529.8, 531.5 and 532.8 eV, which was very consistent with the literature.⁵⁵ It indicated that SV60 had high crystallinity. The peak area percentage of OL, OV and OV/OL values were presented in Table S1.[†] The result manifested that the OVs concentration in SV was in negative correlation with the calcination temperature. It was reasonably inferred that the OVs concentration in SV was bound up with the content of amorphous phase.³³ The calcination temperature was the crucial and convenient parameter to regulate the amorphous phase and the OVs concentration.

A series of 10%SVC samples were prepared at 400, 450, 500 and 550 °C for 2 h to further confirm the influence of calcination temperature on the OVs concentration. The surface OVs concentration was calculated based on XPS O1s spectra and the

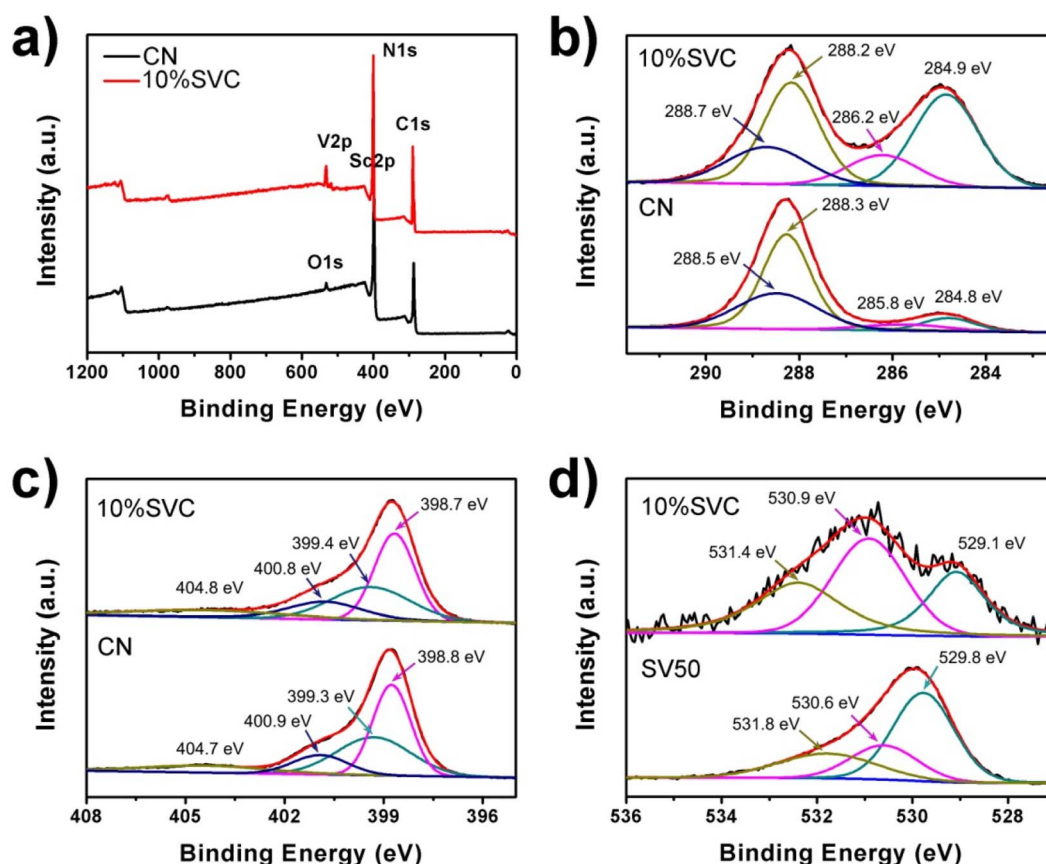


Fig. 5 (a) XPS survey spectra, high resolution XPS spectra of (b) C1s, (c) N1s and (d) O1s of CN, 10%SVC and SV50.



results were listed in Table S2.† As displayed in Table S2,† the surface OV concentration of 10%SVC was increased with the rise of calcination temperature. It reached the maximum when the calcination temperature was 500 °C. Conversely, the surface OV concentration then decreased while the calcination temperature exceeded 500 °C. These results signified that the optimal surface OV concentration was achieved at 500 °C, which are adequately clarified that the calcination temperature can effectively control the crystallinity and the surface OV concentration of the catalyst.³³ EPR spectra can provide the direct evidence to confirm the presence of OV in CN, SV and 10%SVC.⁵⁶ As shown in Fig. 6a, the typical EPR signals of OV at $g = 2.003$ were observed. This signal could be attributed to the electrons trapped by the OV. The weak EPR signal of pristine SV indicated that SV sample had low OV concentration. Meanwhile, 10% SVC exhibited strong signal at $g = 2.003$, suggesting that the OV were successfully introduced into 10%SVC.

UV-vis DRS of CN, SV and SVCs were measured to assess its optical properties. Fig. 6b revealed that the absorption edge of CN was about 457 nm. It was equivalent to 2.71 eV of band gap energy (E_g), which implied that CN nanosheets had certain visible light absorption capacity. This result was similar to the E_g of pure CN in the literature.¹⁴ SVCs samples exhibited almost identical absorption edge, indicating that the incorporation of SV and CN did not change the band gap energy. SV possessed higher absorbance in the UV and visible light region than that

of CN and SVCs samples. The absorption edge of SV was around 617 nm, corresponding to 2.01 eV of E_g . Conversely, the visible light absorption of SVCs has not been effectively improved. In addition, the small shoulder was found in the visible light region in DRS spectra of SVCs, which might derive from the abundant OV in the catalysts.⁵⁶ The E_g of pristine SV and CN were determined by the Kubelka–Munk function. As shown in Fig. 6c, the E_g of SV and CN were respectively 2.01 and 2.71 eV, which were consistent with the DRS results. The VB positions of SV and CN were +1.60 and +1.19 eV according to the XPS VB spectra depicted in Fig. 6d. The corresponding CB positions of SV and CN could be consequently estimated based on the formula: $E_g = E_{VB} - E_{CB}$. It was -0.41 and -1.52 eV, respectively.

The visible light driven photocatalytic degradation activities of LVX, MB and RhB over as-prepared CN, SVCs and SV samples were investigated by using a 40 W White LED as the light source. The photodegradation kinetic curves of LVX on CN, SVCs and SV were illustrated in Fig. 7a. 10%SVC exhibited best photocatalytic performance among these samples. 89.1% of LVX was eliminated within 80 min over 10%SVC. The apparent quantum yield for LVX degradation was 2.01%, which was 4.3 and 17.8 times of pristine CN and SV, respectively. Compared to CN and SV, all SVC heterojunctions presented enhanced photocatalytic activities, indicating that the formation of the SVC heterostructures and the optimization of the surface OV concentration had accelerated the separation of the photogenerated

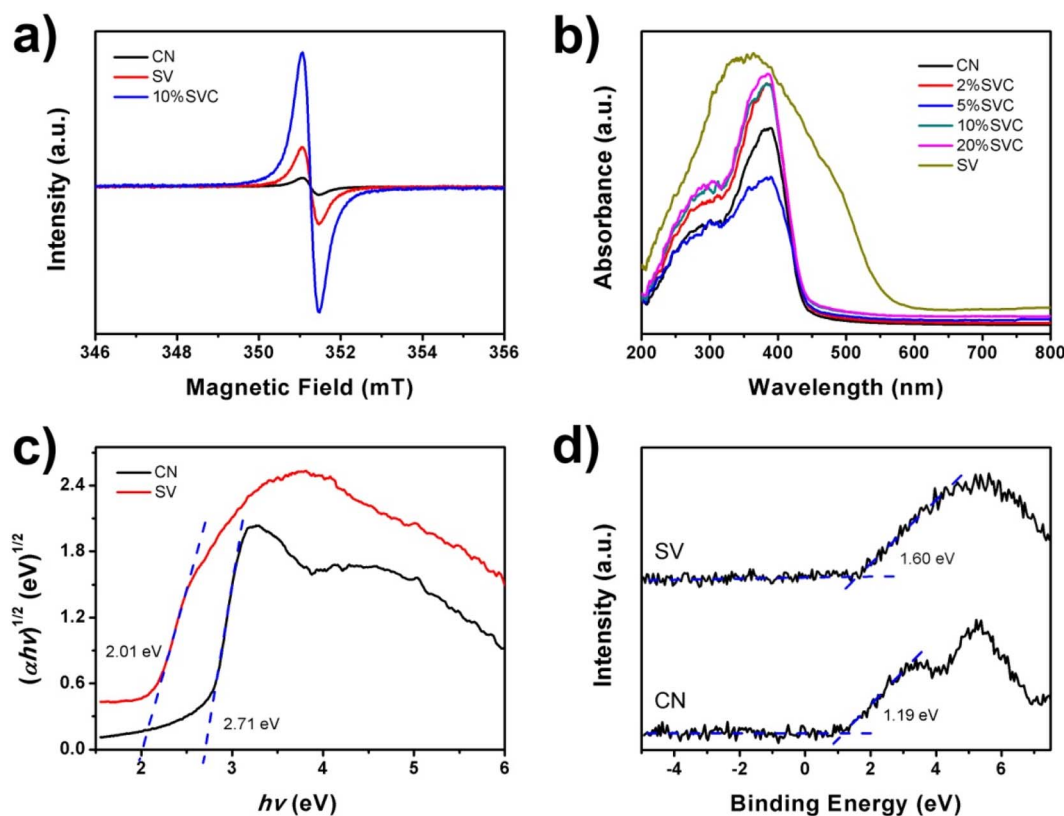


Fig. 6 (a) EPR spectra of CN, SV and 10%SVC, (b) UV-vis diffuse reflectance spectra of CN, SV and SVCs, (c) XPS VB spectra and (d) band gap energies of CN and SV.



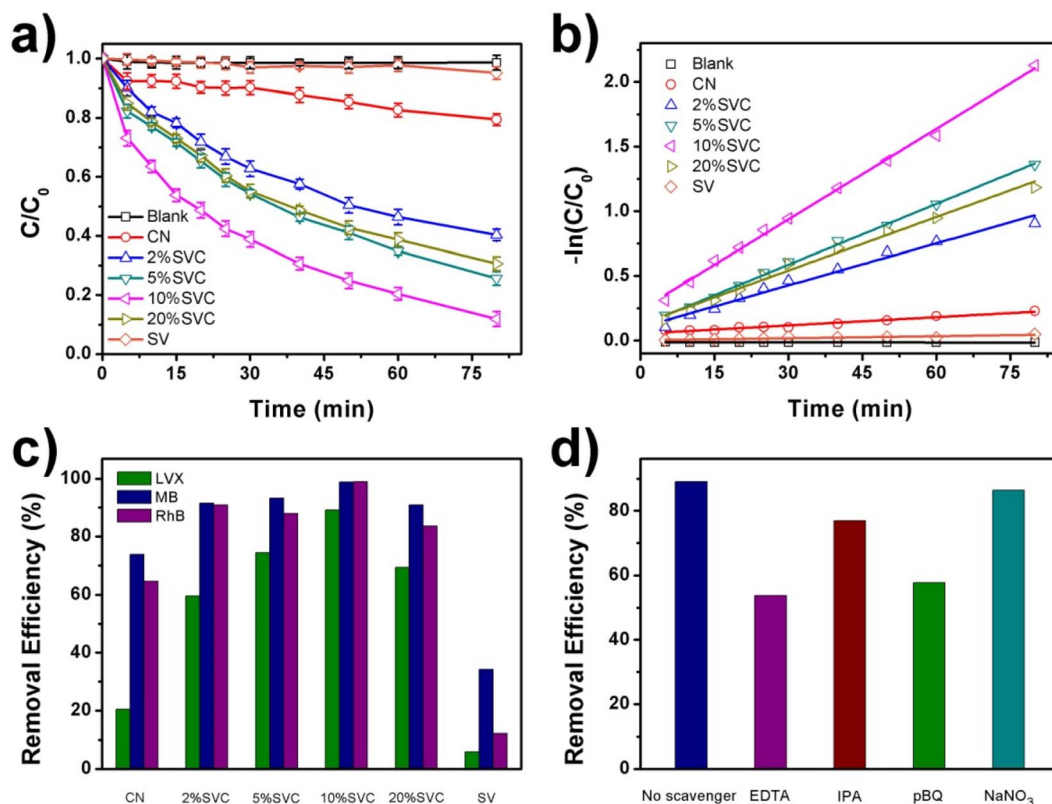


Fig. 7 (a) Photocatalytic degradation of LVX over CN, SVCs and SV under visible light irradiation, (b) corresponding first-order kinetics fitting plots, (c) removal efficiency of LVX, MB and RhB over 10%SVC and (d) the effect of radical scavengers on the photocatalytic degradation of LVX over 10%SVC [experimental conditions: $V = 200$ mL; $LVX = 12.0$ mg L^{-1} ; catalyst dosage = 0.5 g L^{-1}].

charge carriers and finally improved its photocatalysis performance. MB and RhB were also selected as the target pollutants to further verify the excellent visible light driven photocatalytic activities of SVC heterojunctions. The photodegradation kinetic curves of MB and RhB on CN, SVCs and SV were displayed in Fig. S7a and S8a.† 10%SVC also manifested highest degradation efficiency. 98.8% of MB and 99.0% of RhB were removed, which was 2.9 and 8.1 times of pure SV, respectively. SVC heterojunctions all emerged higher photocatalytic activities than that of CN and SV. Furthermore, the photocatalytic degradation efficiency of MB, RhB and LVX on 10%SVC was compared with various recent reported results. As depicted in Table S4,† its degradation activity was higher than that of literature values, signifying that 10%SVC is probably a valuable catalyst of practical application in the environmental sewage treatment.

The photodegradation data of LVX over the as-prepared catalysts followed the pseudo first order kinetic equation (Fig. 7b). The order of the apparent rate constants (k_{app}) was as follows: 0.00213 min^{-1} (CN) $<$ 0.00313 min^{-1} (SV) $<$ 0.0108 min^{-1} (2%SVC) $<$ 0.0138 min^{-1} (20%SVC) $<$ 0.0156 min^{-1} (5%SVC) $<$ 0.0234 min^{-1} (10%SVC). The k_{app} of 10%SVC was 11.0 and 7.5 times of pristine CN and SV, respectively. The photodegradation reaction data of MB and RhB over CN, SVCs and SV were also fitted by the pseudo first order kinetic equation (Fig. S7b and S8b†) and the k_{app} was listed in Table S3.† It was observed that the k_{app} of MB degradation on 10%SVC was 2.5 and 8.3 times of

pristine CN and SV. Meanwhile, the k_{app} of RhB degradation on 10%SVC was 3.4 and 40.0 times of pristine CN and SV. In addition, the enhanced photocatalytic activities of SVC heterojunctions could be confirmed in Fig. 7c. The removal efficiencies of LVX, MB and RhB over CN, SVCs and SV lucidly revealed that SVCs possessed much higher photocatalytic performance than that of pristine CN and SV under the same degradation conditions.

Furthermore, we synthesized a series of 10%SVC samples at different calcination temperature and characterized the surface OV by O1s XPS spectra. The results demonstrated that the surface OV of 10%SVC were significantly affected by the calcination temperature. Herein, LVX photocatalytic degradation experiments over this series of 10%SVC samples were conducted to analyze the effect of surface OV on the photocatalytic performance. The results were exhibited in Fig. S9.† It was observed that the photodegradation performance of 10% SVC increased firstly and then decreased with the calcination temperature, the optimum removal efficiency was acquired when the calcination temperature reached 500 °C. Comparing with the surface OV concentration of 10%SVC samples from O1s XPS spectra results (Table S2†), in which 10%SVC50 possessed the optimal surface OV concentration. The LVX photocatalytic degradation performance was positive correlation with the surface OV concentration in 10%SVC. Therefore, combining the photocatalytic experiments, O1s XPS and EPR



spectra results, we could reasonably infer that the formation of the SVC heterostructures and the optimization of the surface OV's concentration were two major factors to accelerate the separation of the photogenerated charge carriers and finally to improve its photocatalysis performance. The photodegradation of LVX on 10%SVC was implemented for four cycles to assess the structural and photocatalytic stability of the catalyst. The recycling experiments and the FTIR, XRD results of 10%SVC before and after the photodegradation reaction were displayed in Fig. S10.† The negligible decrease of photodegradation efficiency signified the superior photocatalytic stability of 10%SVC. Meanwhile, the variations were hardly observed in the XRD and FTIR results, indicating 10%SVC possessed excellent structural stability.

To identify the reactive species involved in the photocatalytic degradation of LVX on 10%SVC, NaNO₃, EDTA, IPA, and pBQ were selected as the scavengers to trap e⁻, h⁺, ·OH and ·O₂⁻, respectively. The radical scavenger experiments were implemented under the same photodegradation conditions. As the results exhibited in Fig. 7d, the photodegradation efficiency of LVX on 10%SVC were remarkably reduced by the addition of EDTA and pBQ, indicating that h⁺ and ·O₂⁻ were the main reactive species participated in the LVX photodegradation process. Furthermore, the addition of IPA also slightly inhibited the photodegradation of LVX, suggesting ·OH played the minor role in the degradation process. The presence of NaNO₃ hardly affected the degradation, which proved that e⁻ did not directly involved in the photodegradation reaction of LVX.

ESR spectra can provide a direct evidence to authenticate the participation of the ·O₂⁻ and ·OH in the photodegradation

reaction of LVX. In this technique, DMPO was utilized as the capture agent for the measurement of ·O₂⁻ and ·OH with the presence of CN, 10%SVC and SV. As shown in Fig. 8a, the DMPO··O₂⁻ signal of 10%SVC was clearly stronger than that of pristine SV and CN, revealing that more ·O₂⁻ radicals were generated with the existence of 10%SVC. The DMPO··O₂⁻ signal of SV was extremely weak, indicating the poor yield of ·O₂⁻. Similar results were observed in the ESR spectra of DMPO··OH with the presence of CN, 10%SVC and SV (Fig. 8b). 10%SVC had stronger DMPO··O₂⁻ and DMPO··OH signals, demonstrating the formation of the SVC heterostructures and the optimization of the surface OV's concentration was advantageous to accelerate the production of the ·O₂⁻ and ·OH radicals, even if the photocatalytic activity of pristine CN and SV was finite. It might result from the higher spatial separation efficiency of the photogenerated charge in SVCs. It also was proved by the photodegradation of LVX, MB and RhB. Actually, in these photodegradation experiments, CN exhibited better photocatalytic activity than SV, which was very in accordance with the ESR results.

To further appraise the spatial separation efficiency of the photogenerated charge carriers, TPC spectra, EIS Nyquist plots and photoluminescence spectra (PL) of CN, 10%SVC and SV were measured. TPC spectra of these samples were illustrated in Fig. 8c. The photocurrent response curves of CN, 10%SVC and SV was stable and repeatable within 5 cycles of the on-off switching period, suggesting the high photo-stability of as-prepared catalysts. 10%SVC manifested higher photocurrent density than that of pristine CN and SV. The higher photocurrent implied the faster migration and the slower recombination of the photogenerated charge carriers, which finally resulted in

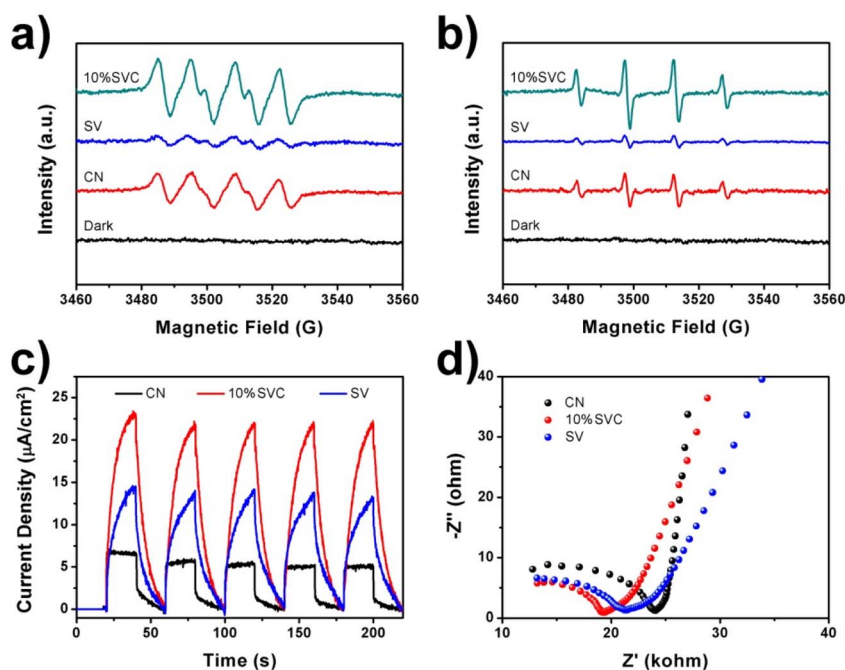


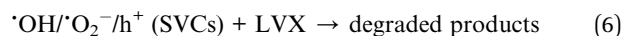
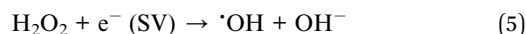
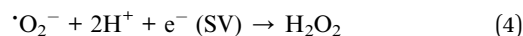
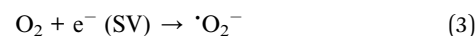
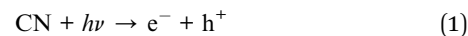
Fig. 8 ESR spectra of (a) DMPO··O₂⁻ in methanol and (b) DMPO··OH in water with the presence of CN, 10%SVC and SV, (c) TPC spectra and (d) EIS Nyquist plots of CN, 10%SVC and SV.



better photocatalytic performance.⁵⁷ Therefore, the enhanced photodegradation activity of LVX, MB and RhB on 10%SVC can attribute to the higher spatial separation efficiency of the electron–hole pairs in 10%SVC. The formation of the SVC heterostructures and the optimization of the surface OV concentration might be two major factors to accelerate the separation of the charge carriers. EIS Nyquist plots of CN, 10% SVC and SV were depicted in Fig. 8d. It clearly showed that 10% SVC had much smaller arc radius than pristine CN, meaning that 10%SVC possessed lower charge transfer resistance⁵⁸ and consequently contributed to reduce recombination rate of photogenerated carriers. Furthermore, the PL spectra of CN, 10%SVC and SV were displayed in Fig. S11.† The PL intensity decreased as compared to CN when the 10%SVC heterojunction formed. It was another evidence to indicate that the photo-generated carrier separation efficiency of 10%SVC was higher than that of CN.⁵⁹ It could therefore effectively enhance photocatalytic degradation activity of LVX.

According to the results analyzed above, the schematic diagram of energy band structures of CN, SV and the possible mechanism of the photodegradation of LVX on 10%SVC were proposed and schemed in Fig. 9. The E_{VB} of SV and CN was +1.60 and +1.19 eV, corresponded to the E_{CB} of −0.41 and −1.52 eV, respectively. Therefore, the SVCs samples were type II heterojunction. The photogenerated electrons and holes were firstly generated in SVCs under the stimulation of the visible light (eqn (1) and (2)). In consideration of SVCs were belonged to type II heterojunction, the photogenerated electrons on the CB of CN migrated onto the CB of SV, and the photogenerated holes on the VB of SV transferred to the VB of CN. The VB potential of SV was negative than the redox potentials of $\cdot\text{OH}/\text{OH}^-$ (1.99 eV vs. NHE) and $\cdot\text{OH}/\text{H}_2\text{O}$ (2.27 eV vs. NHE),⁶⁰ revealing that H_2O and OH^- could not be oxidized directly to form $\cdot\text{OH}$ by the holes on the VB of CN. Comparatively, the CB potential of SV was negative enough than the redox potential of $\text{O}_2/\cdot\text{O}_2^-$ (−0.33 eV vs. NHE). Therefore, the photogenerated electrons could reduce O_2 to generate $\cdot\text{O}_2^-$

(eqn (3)). The $\cdot\text{O}_2^-$ could subsequently react with H^+ to produce H_2O_2 (+0.68 eV vs. NHE) (eqn (4)). And H_2O_2 was then reduced by the photogenerated electrons to get $\cdot\text{OH}$ ⁶¹ (eqn (5)). Finally, $\cdot\text{OH}$, $\cdot\text{O}_2^-$ and h^+ triggered the photodegradation of LVX (eqn (6)).



4. Conclusion

In summary, the $\text{Sc}_2\text{VO}_{5-3}/\text{g-C}_3\text{N}_4$ heterojunctions with abundant OVs were successfully manufactured. The surface OVs of the catalyst can be effectively regulated by adjusting the calcination temperature. XRD, TEM, HRTEM, SEM, FTIR, Raman and XPS results clarified the successful formation of SVCs and its ultrathin structure. DRS and XPS VB spectra revealed it was type II heterojunction. EPR confirmed the existence of OVs and XPS O1s spectra determined the OVs concentration. PL, TPC and EIS assessed the promotion of the separation efficiency and the inhibition of the recombination of the photogenerated charge carriers in SVCs. The influence of OVs and SV content on the photocatalytic degradation efficiency of LVX, MB and RhB was investigated. Due to the introduction of surface OVs and the formation of heterojunction, the as-prepared SVCs manifested highly enhanced photocatalytic degradation efficiency. The as-prepared 10%SVC exhibited best photocatalytic performance. 89.1% of LVX, 98.8% of MB and 99.0% of RhB were effectively eliminated and the photodegradation data followed the pseudo first order kinetic equation. The k_{app} of LVX degradation on 10%SVC was 11.0 and 7.5 times of pristine CN and SV. The h^+ , $\cdot\text{OH}$ and $\cdot\text{O}_2^-$ were the major reactive species in the photodegradation process. This study supplied a new paradigm for further exploring the synergistic effect of surface OVs and heterostructure on the photocatalytic activity.

Author contributions

Jian Feng: conceptualization, methodology, supervision, writing – review & editing, resources, funding acquisition. Liyao Zu: investigation. Hongrong Yang: investigation. Yuanyuan Zheng: investigation. Ziying Chen: investigation. Wei Song: investigation. Ran Zhao: investigation. Li Wang: investigation, data curation, visualization. Xia Ran: investigation, visualization. Bo Xiao: project administration.

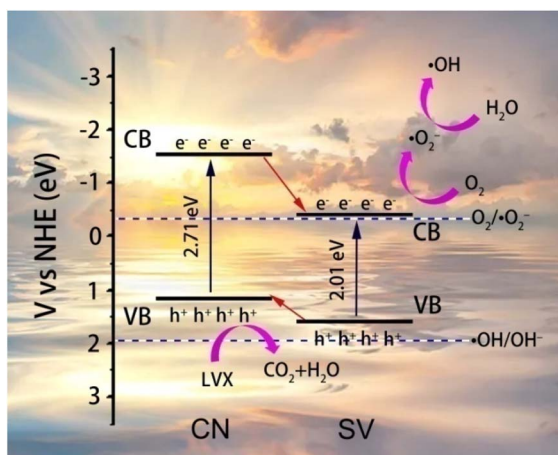


Fig. 9 The possible mechanism of the photodegradation of LVX on 10%SVC.



Conflicts of interest

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

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