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# Boosting exciton dissociation in anion and cation co-doped polymeric semiconductor for selective oxidation reaction†

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The inherently low dielectric properties and weak shielding effect of polymeric semiconductors cause excitons to dominate their photoexcitation process, which greatly restricts the photocatalytic performances mediated by charge carriers. Here, an anion and cation co-doping strategy was proposed to weaken the binding energy of excitons by forming distinct positive and negative charge regions, where the charge asymmetry produced an external potential to drive exciton dissociation. Using polymeric carbon nitride as a typical model framework, we show that the incorporation of anions ( $\text{Cl}^-$ ,  $\text{Br}^-$ ,  $\text{I}^-$ ) and cations ( $\text{Na}^+$ ,  $\text{K}^+$ ) could create a significant spatial separation of electrons and holes, thereby promoting exciton dissociation. Specifically,  $\text{K}^+$  and  $\text{Cl}^-$  co-doped polymeric carbon nitride could effectively promote the dissociation of excitons into hot carriers, contributing to the outstanding efficiency in hot-electron-involved photocatalytic processes, such as the generation of superoxide radicals ( $\text{O}_2^{\cdot-}$ ) and the oxidation of phenylboric acid under visible light. This work presents a practical approach for promoting excitons dissociation through the introduction of charge asymmetry.

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## Introduction

Nowadays, with the rapid development of social productivity, human demand for energy has far exceeded historical levels, and the utilization of clean energy is in great demand.<sup>1–3</sup> Photocatalysis is considered to be a sustainable way to convert solar energy into chemical energy, whose efficiency is closely related to the concentration of photoinduced charge carriers in catalysts.<sup>4–6</sup> As one of the most important types of photocatalysts, polymeric semiconductors have attracted great attention for their properties such as low biotoxicity, tunable band structure, and ease of structural modification.<sup>7,8</sup> However, the strong Coulomb interaction and the weak dielectric shielding effect of polymeric semiconductors make excitons the dominant photogenerated species, where only a few of them can dissociate into free carriers to participate in the artificial photocatalytic process. Meanwhile, a high concentration of excitons in the system leads to the occurrence of exciton–exciton annihilation, resulting in the consumption of photoinduced species.<sup>9,10</sup> Thus, the dissociation of excitons into free electrons and holes would be a practical strategy to enhance the

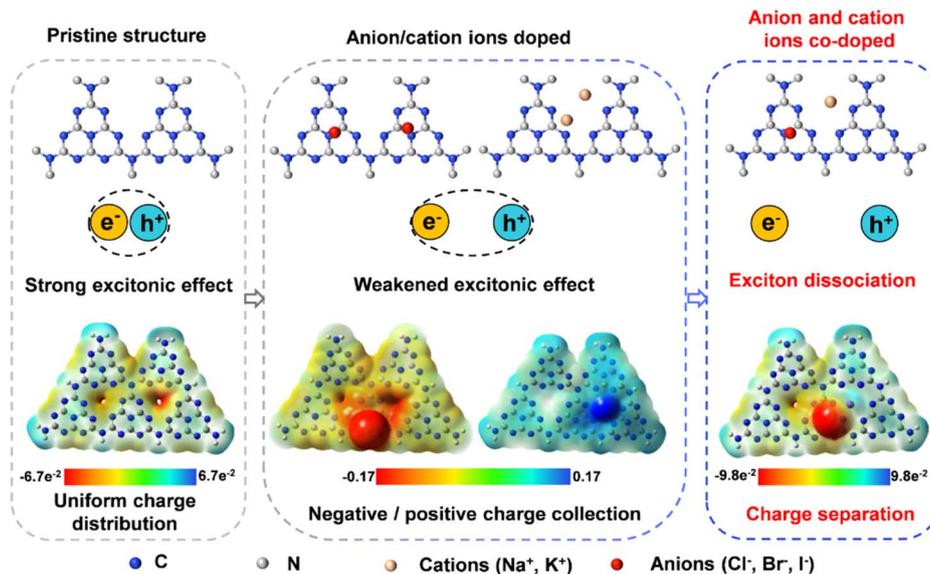
quantum efficiency of photocatalysts, and thus, it is in demand for oxidation–reduction reactions involving charge carriers.<sup>11,12</sup>

As excitons are electrically neutral quasiparticles that are formed by Coulomb interactions between the electrons and holes, the introduction of charge asymmetry generates an external potential to drive their dissociation.<sup>13–15</sup> Among the various strategies, elemental doping is an efficient way to break the local charge symmetry of photocatalysts by introducing external ions with various levels of electronegativity.<sup>16</sup> In addition, defective structures can serve as reactive sites to improve the separation efficiency of photo-generated charge carriers.<sup>17</sup> Both the defective structures in the construction and elemental doping methods could enhance the concentration of photo-generated charge carriers by creating different structural units. Although single-element doping may partially promote exciton dissociation into hot carriers, the possible formation of a steady charge-transfer state prevents further separation of carriers.<sup>18,19</sup> Herein, using typical polymeric carbon nitride (PCN) as an example, we proposed an anion and cation co-doping method to promote the exciton dissociation by introducing strong charge asymmetry. As depicted in Scheme 1, the pristine carbon nitride with uniform distribution of the surface electrostatic potential indicates the symmetry distribution of charges, further confirming its potentially strong excitonic effect. In addition, doping with either individual anions or cations in polymeric carbon nitride results in the collection of negative or positive charges in the PCN framework, partially weakening the binding of electrons and holes. As for co-doping,

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Scheme 1 Simulated electrostatic potential of polymeric carbon nitride under various doping conditions.

anions ( $\text{Cl}^-$ ,  $\text{Br}^-$ ,  $\text{I}^-$ ) and cations ( $\text{Na}^+$ ,  $\text{K}^+$ ) located at different sites of the PCN framework exhibit a distinct distribution of negative and positive charges, where the as-formed electrostatic potential facilitates the dissociation of excitons.

To further investigate this, density functional theory (DFT) calculations were conducted to study the influence of incorporating  $\text{K}^+$  and  $\text{Cl}^-$  ions for co-doping on the exciton dissociation of the polymeric carbon nitride matrix. Fig. 1a and b depict the spatial distributions of the highest occupied molecular orbital (HOMO) and the lowest unoccupied molecular orbital (LUMO) of the polymeric carbon nitride. The uniform distributions and significant overlaps on the tri-*s*-triazine structure rule out the possibility of the dissociation of excitons within the framework. In contrast, a significant spatial separation between the highest occupied molecular orbital (HOMO) and lowest unoccupied molecular orbital (LUMO) was observed in the  $\text{K}^+$  and  $\text{Cl}^-$  co-doped model (Fig. 1c and d). Specifically,  $\text{Cl}^-$ -containing units with electron-rich features dominate the distribution of the highest occupied molecular orbital (HOMO), while a  $\text{K}^+$ -

containing unit dominates the distribution of the lowest unoccupied molecular orbital (LUMO). Such spatial separation of electrons and holes between the two distinct motifs indicates the potential dissociation of excitons in the  $\text{K}^+$ ,  $\text{Cl}^-$  co-doped polymeric carbon nitride system.

## Results and discussion

In this study, PCN co-doped with anions ( $\text{Cl}^-$ ,  $\text{Br}^-$ ,  $\text{I}^-$ ) and cations ( $\text{Na}^+$ ,  $\text{K}^+$ ) was synthesized through a modified thermal polymerization of thiourea and the corresponding doping element-containing compounds (See detail in ESI<sup>†</sup>). Using the typical  $\text{K}^+$ ,  $\text{Cl}^-$  co-doped polymeric carbon nitride (denoted as  $\text{KCl-CN}$ ) as an example, a range of  $\text{KCl-CN}$  samples with varying contents of doping element and carbon nitride samples doped with individual  $\text{K}^+/\text{Cl}^-$  (denoted as  $\text{K-CN}$  and  $\text{Cl-CN}$ , respectively) were synthesized for comparison. The powdered X-ray diffraction (PXRD) patterns of  $\text{KCl-CN}$  (Fig. 2a and S1<sup>†</sup>) exhibit a weakened (100) peak and (002) peak, indicating a decrease in crystallinity and an increase in structural distortion of PCN after elemental doping.<sup>20</sup> As shown in Fig. 2b, the  $\text{KCl-CN}$  spectrum exhibits a similar Fourier transform infrared (FT-IR) spectrum as that of PCN, indicating that both retained essentially tri-*s*-triazine ring units. However, compared to the PCN, the FT-IR spectrum of  $\text{KCl-CN}$  exhibits a prominent peak at  $2181\text{ cm}^{-1}$ , which can be attributed to the stretching vibration of cyano groups ( $-\text{C}\equiv\text{N}$ ) formed from the amine groups ( $-\text{NH}_2$ ) in the heptazine rings.<sup>21</sup> Furthermore,  $\text{KCl-CN}$  shows two new absorption peaks at  $992$  and  $1154\text{ cm}^{-1}$ , which are attributed to the ion-dipole interaction of heptazine with alkali ions ( $\text{K}^+$ ) and halide ions ( $\text{Cl}^-$ ) over  $\text{KCl-CN}$ . Solid-state  $^{13}\text{C}$  NMR spectra (Fig. 2c) show that  $\text{KCl-CN}$  presents two new peaks at  $122.7$  and  $171.6\text{ ppm}$ , which are likely associated with the C (3) atom in the cyano group ( $-\text{C}\equiv\text{N}$ ) and the C (4) atom in the polymeric carbon nitride structure.<sup>22,23</sup> In addition, elemental

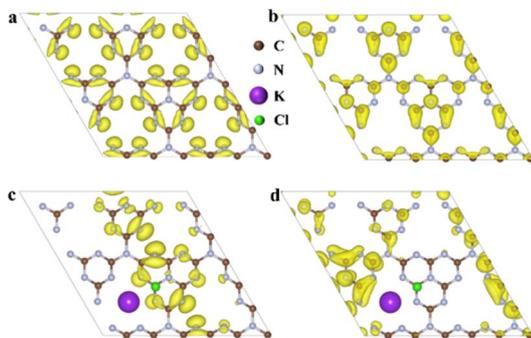


Fig. 1 The highest occupied molecular orbital (HOMO) and lowest unoccupied molecular orbital (LUMO) distributions in PCN (a and b), and  $\text{K}^+$ ,  $\text{Cl}^-$  co-doped CN (c and d), respectively.



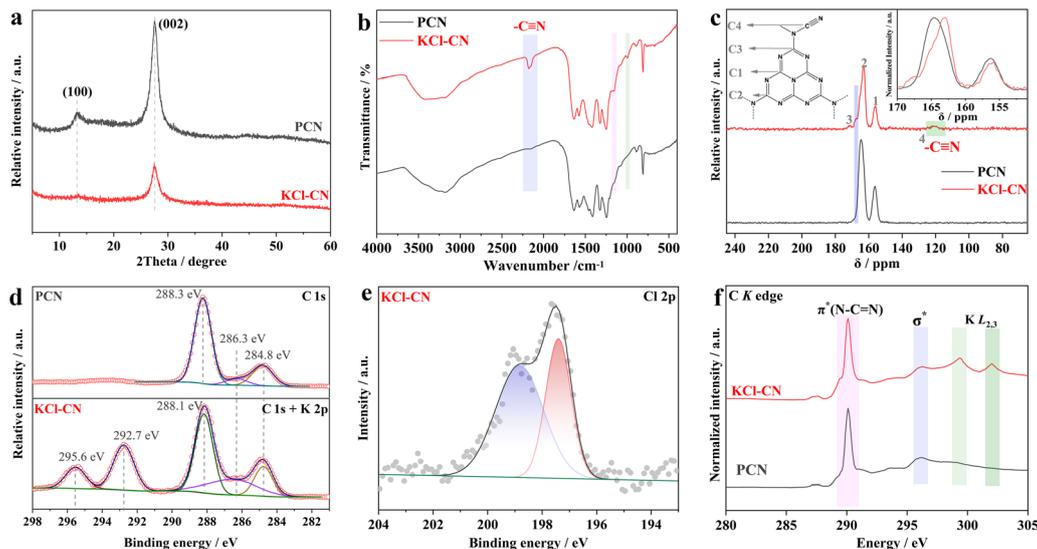


Fig. 2 (a) Powder XRD patterns, (b) FT-IR spectra, (c) solid-state  $^{13}\text{C}$  NMR spectra (inset: normalization intensity of solid-state  $^{13}\text{C}$  NMR spectra) and (d) C 1s XPS spectra of PCN and KCl-CN; (e) Cl 2p spectrum of KCl-CN; (f) C K-edge XANES curves of PCN and KCl-CN.

mapping with the energy-dispersive X-ray spectra (EDS) (Fig. S4†) illustrates the uniform distribution of  $\text{K}^+$  and  $\text{Cl}^-$  in the KCl-CN. X-ray photoelectron measurements (XPS) were performed to further explore the elements and chemical states of samples. In the high-resolution C 1s-K 2p XPS spectra (Fig. 2d), KCl-CN exhibits two distinct new peaks at 295.6 and 292.7 eV, corresponding to the  $\text{K } 2\text{p}_{3/2}$  and  $\text{K } 2\text{p}_{1/2}$  peaks of the K element. Meanwhile, KCl-CN exhibits a significant Cl  $2\text{p}_{3/2}$  and Cl  $2\text{p}_{1/2}$  signal in the Cl 2p spectrum (Fig. 2e). As compared to PCN, the peak corresponding to the  $\text{C-NH}_x$  at 288.3 eV in KCl-CN exhibits an obvious shift toward low energy, owing to the formation of cyano groups ( $-\text{C}\equiv\text{N}$ ) with a strong electron-withdrawing effect during the polymerization process. Similar low-energy shifts were observed in the N 1s spectra (Fig. S5b†).<sup>24</sup> Interestingly, the binding energy of  $\text{K}^+$  and  $\text{Cl}^-$  in KCl-CN is lower and higher than that of the KCl, respectively, implying the  $\text{K}^+$  and  $\text{Cl}^-$  are incorporated into PCN and bonded with the surrounding C and N atoms.<sup>25</sup> In the synchrotron X-ray absorption near-edge structure (XANES) spectra (Fig. 2f), as compared with PCN, KCl-CN presented new peaks at 298.2 and 302.5 eV, corresponding to  $\text{KL}_{2,3}$ .<sup>26,27</sup> Based on these results, it is safe to conclude that  $\text{K}^+$  and  $\text{Cl}^-$  were successfully introduced into PCN. At the same time, the other anions, *i.e.*,  $\text{Br}^-$  and  $\text{I}^-$ , and cations, *i.e.*,  $\text{Na}^+$  and  $\text{K}^+$ , were also successfully co-doped into the PCN framework (Fig. S6 and S7†).

The introduction of  $\text{K}^+$  and  $\text{Cl}^-$  into PCN is anticipated to have a significant impact on its excitonic effects, where spectroscopic measurements were carried out to investigate the photoexcitation processes involved. Compared PCN, KCl-CN narrows the band gap and enhances the absorption capacity of visible light, as illustrated in ultraviolet-visible (UV-vis) spectra and the corresponding Tauc plots in Fig. 3a. Based on their light absorption, the excitation wavelength for photoluminescence (PL) measurements was selected at 365 nm. As shown in Fig. 3b, the photoluminescence intensity of KCl-CN is lower than that

of PCN, indicating a significant reduction in the population of singlet excitons in KCl-CN. Time-resolved fluorescence spectroscopy was carried out to monitor the corresponding lifetime

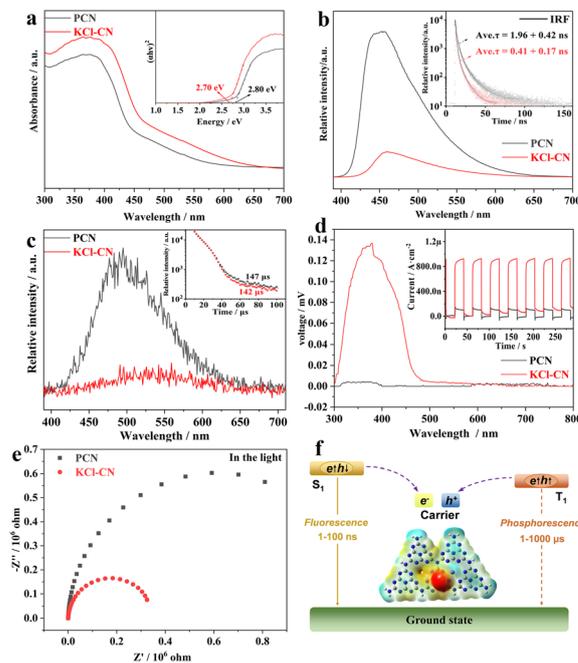


Fig. 3 (a) UV-vis spectra (inset: corresponding Tauc plots), (b) photoluminescence spectra (inset: time-resolved fluorescence spectra), (c) phosphorescence spectra (inset: time-resolved phosphorescence spectra), (d) surface photovoltage (inset: periodic on/off photocurrent response curves at a constant potential of approximately 0.4 V vs. Ag/AgCl reference electrode in 0.5 M  $\text{Na}_2\text{SO}_4$  under the light on/off condition) and (e) EIS Nyquist plots under visible light conditions of PCN and KCl-CN (EIS was performed over a frequency range of 0.01 Hz to  $10^6$  Hz with an AC voltage amplitude of 5 mV); (f) schematic illustration of the exciton dissociation process of anion and cation co-doped samples.



of emission peaks. The radiative decay results (Fig. 3b inset and Table S1†) could be described as a triple-exponential fitting, yielding lifetimes of 1.96 ns for PCN and 0.41 ns for KCl-CN. The reduction in the radiation emission lifetime can be understood by the reduction in the population of the singlet excitons in KCl-CN.<sup>28,29</sup> Phosphorescence spectra (Fig. 3c) were employed to investigate long-lived photoinduced species, revealing that PCN exhibits a more pronounced phosphorescence emission compared to KCl-CN. Furthermore, time-resolved phosphorescence kinetics spectra show that the phosphorescence lifetime of KCl-CN is shorter than that of PCN, indicating a distinct reduction in the population and lifetime of triplet excitons in KCl-CN.<sup>18</sup> Considering the competitive relationship between excitons and hot carriers, the reduced number of excitons is beneficial to enhance the generation of charge carriers. Electrochemical measurements were carried out to further verify this conjecture. As shown in Fig. 3d, KCl-CN exhibits a higher surface photovoltage intensity than that of PCN, indicating efficient carrier generation due to co-doping with  $K^+$  and  $Cl^-$ . Furthermore, KCl-CN exhibits a 6.1~fold higher response on periodic on/off photocurrent to PCN under a 400 nm cut-off filter at a 0.4 V bias voltage. As evidenced by the electrochemical impedance spectroscopy (EIS) results in Fig. 3e, KCl-CN possesses a lower electrochemical impedance than the PCN under illumination, leading to improved electron transport.<sup>30,31</sup>

The above results confirm that  $K^+$  and  $Cl^-$  co-doping has significant positive effects on boosting exciton dissociation into hot carriers under light excitation. Similarly, other anion ( $Cl^-$ ,  $Br^-$ ,  $I^-$ ) and cation ( $Na^+$ ,  $K^+$ ) co-doped polymeric carbon nitride samples with enhanced exciton dissociation exhibit greatly enhanced periodic on/off photocurrent compared to that of the

corresponding individual anion/cation-doped counterpart (Fig. S12†). Thus, anion ( $Cl^-$ ,  $Br^-$ ,  $I^-$ ) and cation ( $Na^+$ ,  $K^+$ ) co-doped polymeric carbon nitride samples promote the dissociation process of excitons, which can be understood by the processes illustrated in Fig. 3f.

Boosting hot carrier generation and transfer in the anion and cation co-doped samples leads to greatly enhanced photocatalytic activities, which were systematically studied using KCl-CN as an example. The Mott-Schottky curves (Fig. S14†) demonstrate the *n*-type semiconducting characteristics of the samples, where the flat potentials of PCN and KCl-CN were measured to be  $-1.14$  and  $-1.04$  V vs. NHE, respectively. Consequently, the band structure of PCN and KCl-CN (Fig. S15†) indicates that both samples are capable of driving the reduction in ground-state oxygen ( $O_2$ ) to superoxide radical ( $O_2^{\cdot-}$ ). Nitrotetrazolium blue chloride (NBT) was used as a feature probe to evaluate the  $O_2^{\cdot-}$  generation ability of samples, which can react with  $O_2^{\cdot-}$  to produce endoperoxide. As shown in Fig. 4a, the time-dependent NBT oxidation absorption curves demonstrate that KCl-CN can quickly react with NBT, as compared to the PCN, indicating enhanced  $O_2^{\cdot-}$  generation of KCl-CN. Furthermore, electron spin resonance (ESR) measurements were used to prove the existence of photogenerated reactive oxygen species (ROS); 5,5-dimethyl-1-pyrroline-*N*-oxide (DMPO) was used as a trapping agent. As shown in Fig. 4b, compared to PCN, KCl-CN typically exhibits a stronger signal of  $O_2^{\cdot-}$ . Given its capabilities in  $O_2^{\cdot-}$  generation, the KCl-CN is anticipated to be an efficient catalyst for the aerobic oxidation of organic molecules, such as the selective oxidation of phenylboronic acid to corresponding phenols and derivatives. As is known, phenols and their derivatives are important aromatic chemicals that are widely employed in industrial, medicinal,

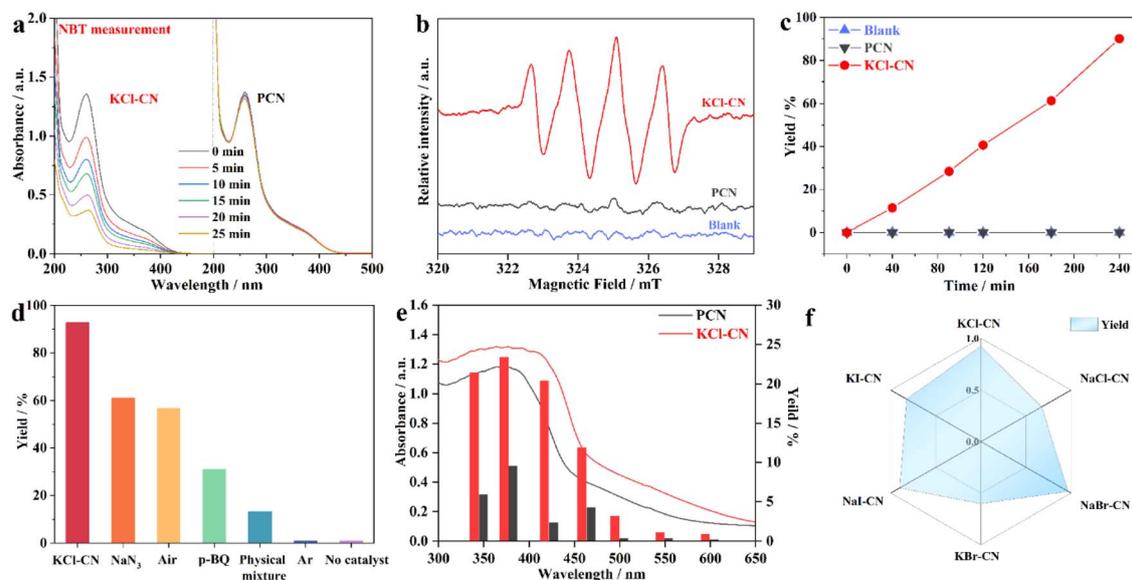
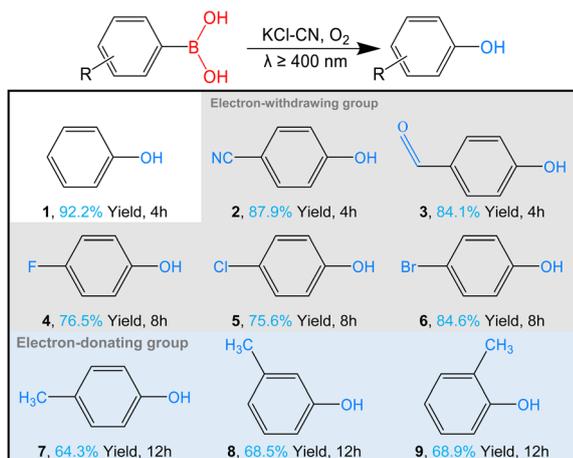


Fig. 4 (a) Time-dependent NBT degradation curves, (b) ESR trapping spectra of PCN and KCl-CN in the presence of DMPO in methanol, (c) time-dependent phenylboronic acid oxidation curves of PCN and KCl-CN, (d) different reaction conditions for phenylboronic acid oxidation spectrum in KCl-CN, (e) illumination wavelength-dependent phenylboronic acid oxidation curves of PCN and KCl-CN, and (f) different catalysts for phenylboronic acid oxidation spectrum. All of the above tests and experiments were performed under  $\lambda \geq 400$  nm.



**Table 1** Oxidation of phenylboronic acid derivatives in KCl–CN. Reaction conditions: catalyst (10 mg), phenylboronic acid (0.1 mmol), methanol (1 mL), acetonitrile (1 mL), xenon lamp (300 W), 298 K and O<sub>2</sub> (1 atm)



and cosmetic domains.<sup>32,33</sup> As depicted in Fig. 4c, KCl–CN exhibited a robust yield in the conversion of phenylboronic acid to phenol under visible light, while the yields of PCN were negligible. Control experiments conducted under various atmospheric conditions indicated that the oxidation reaction depends on the activation of oxygen (Fig. 4d). The addition of p-BQ (a scavenger for O<sub>2</sub><sup>•−</sup>) results in a significant reduction of yield during the oxidation process of phenylboronic acid.

In addition, Fig. 4e exhibits wavelength-dependent phenylboronic acid oxidation, indicating that KCl–CN possesses much higher photocatalytic phenylboronic acid oxidation ability than PCN. Fundamentally, Fig. 4f indicates that PCN co-doped with anions and cations has a superior oxidation capacity of phenylboronic acid, and thus, it shows greatly enhanced phenylboronic acid oxidation yield compared to the corresponding individual anions (Cl<sup>−</sup>, Br<sup>−</sup>, I<sup>−</sup>)/cations (Na<sup>+</sup>, K<sup>+</sup>)-doped counterpart (Fig. S18†). At last, the catalytic activity and structural configuration of KCl–CN were maintained during five consecutive cycling experiments, as evidenced using PXRD and FT-IR characterizations (Fig. S20 and S21†).

With the optimal reaction conditions in our hand, we explored another phenylboronic acid with different substitutions to evaluate the advantages and scope of photocatalytic selective oxidation reactions over the KCl–CN catalyst. In general, KCl–CN demonstrated exceptional performance, regardless of the presence of electron-withdrawing (2–6, Table 1) or -donating groups (7, Table 1), as well as the positions of the substituents (7–9, Table 1). Overall, KCl–CN demonstrated the ability to catalyze the production of a wide range of phenols under visible light, indicating its potential for application in selective photocatalytic oxidation reactions.

## Conclusions

An anion (Cl<sup>−</sup>, Br<sup>−</sup>, I<sup>−</sup>) and cation (Na<sup>+</sup>, K<sup>+</sup>) co-doping method has been proposed to achieve efficient exciton dissociation in

polymeric semiconductors, where boosting the charge carrier generation and photocatalytic efficiency have been demonstrated. Specifically, using the anion and cation co-doped polymeric carbon nitride as an example, we show that the K<sup>+</sup> and Cl<sup>−</sup> co-doped polymeric carbon nitride could create a significant spatial separation of electrons and holes in the formation of charge asymmetry. In addition, all the samples co-doped with anions and cations exhibit enhanced exciton dissociation compared to the corresponding individual anion/cation-doped counterparts. As a result, the anion and cation co-doped samples can readily activate molecular oxygen to form superoxide radicals (O<sub>2</sub><sup>•−</sup>) under visible light, which subsequently react with phenylboronic acid to produce the corresponding phenol. This work provides a practical approach to efficiently promote exciton dissociation and shows the great advantage of co-doping with anions and cations in the photocatalytic process.

## Data availability

Additional experimental details and data are provided in the ESI,† including XRD, EDS, XPS, XANES, SEM, TEM, BET, Mott–Schottky plots, EIS Nyquist plots, and ESR.

## Author contributions

J. G. performed the experiments and collected the data. L. L. and Z. L. conducted and analyzed the DFT calculations. J. Z. assisted in performing the experiments. H. W. and X. Z. cowrote the manuscript. H. W., X. Z. and Y. X. supervised the project. All authors approved the final version of the manuscript.

## Conflicts of interest

The authors declare that they have no competing interests.

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## References

- 1 S. Shafiee and E. Topal, *Energy Policy*, 2009, **37**, 181–189.



- 2 D. G. Vlachos and S. Caratzoulas, *Chem. Eng. Sci.*, 2010, **65**, 18–29.
- 3 Q. Wang and K. Domen, *Chem. Rev.*, 2020, **120**, 919–985.
- 4 H. Kisch, *Acc. Chem. Res.*, 2017, **50**, 1002–1010.
- 5 A. Kumar Singh, C. Das and A. Indra, *Coord. Chem. Rev.*, 2022, **465**, 214516.
- 6 G. Grando, G. Sportelli, G. Filippini, M. Melchionna and P. Fornasiero, *NanoTrends*, 2023, **4**, 100028.
- 7 Y. Wang, A. Vogel, M. Sachs, R. S. Sprick, L. Wilbraham, S. J. A. Moniz, R. Godin, M. A. Zwijnenburg, J. R. Durrant, A. I. Cooper and J. Tang, *Nat. Energy*, 2019, **4**, 746–760.
- 8 J. Dong, Z. Gong, Y. Chen, G. Hao, W. Zhou, J. Li, M. Yang, R. Deng and L. Wang, *Sci. China Mater.*, 2023, **66**, 3176–3188.
- 9 H. Wang, S. Jin, X. Zhang and Y. Xie, *Angew. Chem., Int. Ed.*, 2020, **59**, 22828–22839.
- 10 Y. Li, S. Li, L. Liu, Z. Zhang and W. Ma, *Appl. Catal., B*, 2023, **321**, 122025.
- 11 S. Lin, B. Wu, Q. Li, X. Xiao, M. Zheng, J. Liu, Y. Xie and B. Jiang, *Sci. China Mater.*, 2023, **66**, 4680–4688.
- 12 A. Avateev, I. Ghosh, B. König and M. Antonietti, *Angew. Chem., Int. Ed.*, 2018, **57**, 15936–15947.
- 13 G. D. Scholes and G. Rumbles, *Nat. Mater.*, 2006, **5**, 683–696.
- 14 M. Dvorak, S. H. Wei and Z. Wu, *Phys. Rev. Lett.*, 2013, **110**, 016402.
- 15 L. Yu, H. Li, H. Shang, P. Xing, B. Zhou, Z. Chen, X. Liu, H. Zhang, Y. Shi and L. Zhang, *ACS Nano*, 2023, **17**, 15077–15084.
- 16 K. H. Eckstein, H. Hartleb, M. M. Achsnich, F. Schöppler and T. Hertel, *ACS Nano*, 2017, **11**, 10401–10408.
- 17 X. N. Yu, S. F. Ng, L. K. Putri, L. L. Tan, A. R. Mohamed and W. J. Ong, *Small*, 2021, **17**, 2006851–2006896.
- 18 P. Zhang, L. Li, J. Zhao, H. Wang, X. Zhang and Y. Xie, *Precis. Chem.*, 2023, **1**, 40–48.
- 19 C. K. Lee, L. Shi and A. P. J. Willard, *Phys. Chem. Lett.*, 2016, **7**, 2246–2251.
- 20 X. Li, G. Fang, T. Wu, Q. Tian, Q. Yang and Z. Chen, *Appl. Surf. Sci.*, 2023, **635**, 157717.
- 21 X. Li, H. Mai, X. Wang, Z. Xie, J. Lu, X. Wen, S. P. Russo, D. Chen and R. A. Caruso, *J. Mater. Chem. A*, 2024, **12**, 5204–5214.
- 22 B. Jürgens, E. Irran, J. Senker, P. Kroll, H. Müller and W. Schnick, *J. Am. Chem. Soc.*, 2003, **125**, 10288–10300.
- 23 H. Ou, P. Yang, L. Lin, M. Anpo and X. Wang, *Angew. Chem., Int. Ed.*, 2017, **56**, 10905–10910.
- 24 Y. Xu, M. Fan, W. Yang, Y. Xiao, L. Zeng, X. Wu, Q. Xu, C. Su and Q. He, *Adv. Mater.*, 2021, **33**, 2101455.
- 25 J. Stoch and M. Ladecka, *Appl. Surf. Sci.*, 1988, **31**, 426–436.
- 26 A. W. Gillespie, A. Diochon, B. L. Ma, M. J. Morrison, L. Kellman, F. L. Walley, T. Z. Regier, D. Chevrier, J. J. Dynes and E. G. Gregorich, *Biogeochemistry*, 2013, **117**, 337–350.
- 27 Y. Xu, M. Fan, W. Yang, Y. Xiao, L. Zeng, X. Wu, Q. Xu, C. Su and Q. He, *Adv. Mater.*, 2021, **33**, 2101455.
- 28 A. Indra, R. Beltrán-Suito, M. Müller, R. P. Sivasankaran, M. Schwarze, A. Acharjya, B. Pradhan, J. Hofkens, A. Brückner, A. Thomas, P. W. Menezes and M. Driess, *ChemSusChem*, 2021, **14**, 306.
- 29 H. Wang, X. Sun, D. Li, X. Zhang, S. Chen, W. Shao, Y. Tian and Y. Xie, *J. Am. Chem. Soc.*, 2017, **139**, 2468–2473.
- 30 A. Indra, A. Acharjya, P. W. Menezes, C. Merschjann, D. Hollmann, M. Schwarze, M. Aktas, A. Friedrich, S. Lochbrunner, A. Thomas and M. Driess, *Angew. Chem., Int. Ed.*, 2017, **56**, 1653.
- 31 V. Kumar, V. Vyas, D. Kumar, A. Kushwaha and A. Indra, *Chem. Sci.*, 2024, **15**, 13218–13226.
- 32 R. Bal, M. Tada, T. Sasaki and Y. Iwasawa, *Angew. Chem., Int. Ed.*, 2006, **45**, 448452.
- 33 K. Dong, Y. Yao, H. Li, H. Li, S. Sun, X. He, Y. Wang, Y. Luo, D. Zheng, Q. Liu, Q. Li, D. Ma, X. Sun and B. Tang, *Nat. Synth.*, 2024, **3**, 763–773.

