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# Facile synthesis of crystalline viologen-based porous ionic polymers with hydrogen-bonded water for efficient catalytic CO<sub>2</sub> fixation under ambient conditions†

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In this work, we report a series of crystalline viologen-based porous ionic polymers (denoted VIP-X, X = Cl or Br), that have *in situ* formed dicationic viologens paired with halogen anions and intrinsic hydrogen-bonded water molecules, towards metal-free heterogeneous catalytic conversion of carbon dioxide (CO<sub>2</sub>) under mild conditions. The targeted VIP-X materials were facilely constructed *via* the Menshutkin reaction of 4,4'-bipyridine with 4,4'-bis(bromomethyl)biphenyl (BCBMP) or 4,4'-bis(chloromethyl)biphenyl (BBMBP) monomers. Their crystalline and porous structures, morphological features and chemical structures and compositions were fully characterized by various advanced techniques. The optimal catalyst VIP-Br afforded a high yield of 99% in the synthesis of cyclic carbonate by CO<sub>2</sub> cycloaddition with epichlorohydrin under atmospheric pressure (1 bar) and a low temperature (40 °C), while other various epoxides could be also converted into cyclic carbonates under mild conditions. Moreover, the catalyst VIP-Br could be separated easily and reused with good stability. The remarkable catalytic performance could be attributed to the synergistic effect of the enriched Br<sup>-</sup> anions and available hydrogen bond donors –OH groups coming from H-bonded water molecules.

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## Introduction

Currently, the chemical fixation and utilization of CO<sub>2</sub> has attracted increasing attention, because CO<sub>2</sub> is an environmentally unfavourable greenhouse gas as well as an abundantly available C1 resource.<sup>1</sup> Among various strategies for handling the above issue, the synthesis of high value-added cyclic carbonates by CO<sub>2</sub> cycloaddition with epoxides is one of the most efficient and atom-economic routes to achieve catalytic CO<sub>2</sub> fixation.<sup>2–5</sup> To date, various homogeneous and heterogeneous catalytic systems have been developed for CO<sub>2</sub> conversion.<sup>3–5</sup> In general, homogeneous catalysts can exhibit high catalytic activities under mild conditions but with complicated processes of separation, purification and recycling, which would limit their practical application.<sup>3,4</sup> In contrast, heterogeneous catalysts can solve these problems, but often the reactions are performed under high temperatures and/or CO<sub>2</sub> pressures to obtain the desired catalytic activities.<sup>5</sup> Meanwhile, there are also some problems in preparing efficient heterogeneous catalysts with available active sites *via* an easily-achieved

synthetic method. Therefore, it is necessary to develop a facile method to construct the desired heterogeneous catalysts toward CO<sub>2</sub> fixation under mild conditions.

Porous ionic polymers (PIPs) are a class of rising functional polymeric materials, which have aroused much attention owing to their intriguing characters including abundant ionic sites, controllable porous structures, chemical functionalities, and ion exchange properties.<sup>6,7</sup> In particular, ionic moieties of halogen-based PIPs have strong dipole quadrupole interactions with CO<sub>2</sub>, which endow intrinsic CO<sub>2</sub>-philicity for PIPs, while the specific halogen anions can be used as active nucleophiles for catalytic conversion of CO<sub>2</sub>.<sup>8,9</sup> Further, a lot of ionic liquids (ILs) or IL-like structures in the solid polymers have intrinsic hydrophilicity, which can chemically absorb water molecules nearby ionic moieties *via* hydrogen bond interactions between water and halogen anions (such as Cl<sup>-</sup> and Br<sup>-</sup>).<sup>10–13</sup> It was reported that a suitable amount of water can be regarded as hydrogen bond donors (HBD) co-catalysts for enhancing the coupling of CO<sub>2</sub>/epoxide by forming H-bonds with the oxygen atom of epoxides.<sup>4,14</sup> Therefore, we intend to explore the targeted synthesis of task-specific porous ionic polymers with halogen anions and H-bonded water molecules for enhancing the CO<sub>2</sub> activation and conversion.

Recently, different types of ionic polymers that containing diverse ionic moieties such as imidazolium,<sup>15–18</sup> pyridinium,<sup>19–21</sup>

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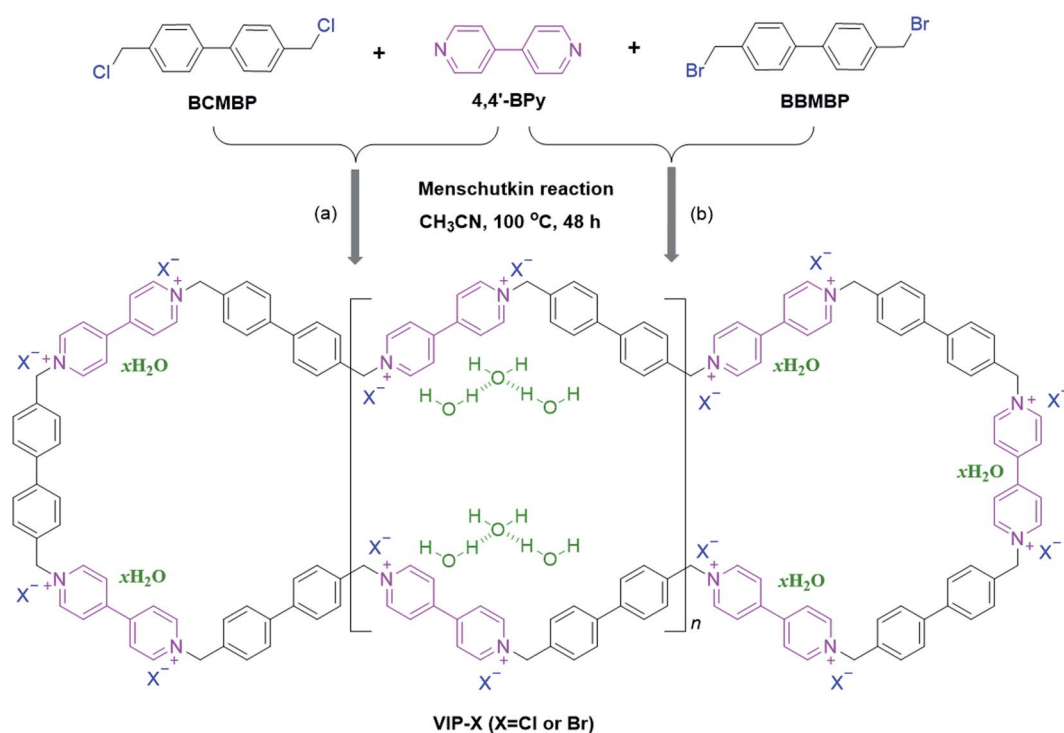
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viologen, <sup>22–25</sup> quaternary phosphonium, <sup>26–28</sup> and others, <sup>6,7,29–31</sup> have been reported. Amongst, viologen-based ionic polymers have been regarded to be one type of promising functionalized polymers towards diverse applications, due to their unique properties of adjustable redox states and viologen ionic active sites. <sup>22,23</sup> In terms of synthesis, a series of viologen-based ionic polymers have been prepared using various strategies including C–C coupling reactions (*e.g.*, Heck reaction, <sup>32</sup> and Sonogashira coupling reaction <sup>33,34</sup>), ionothermal trimerization, <sup>35</sup> imine condensation, <sup>36,37</sup> Zincke reaction, <sup>38–41</sup> and Menshutkin reaction. <sup>11,42–48</sup> Through coupling reactions and ionothermal reactions can afford viologen-based porous ionic polymers with high surface areas, these processes are complicated and environmental-unfriendly that require noble-metal catalysts or additives or high reaction temperatures. <sup>32–37</sup> Very recently, Zincke reaction has been used as a catalyst-free method to prepare viologen-based porous ionic networks with moderate surface areas. <sup>38–41</sup> However, this strategy is highly dependent on using high-cost multi-aniline-based monomers and viologen Zincke salts. Besides, this reaction is not an atomic economic reaction, which inevitably releases the toxic by-product 2,4-dinitroaniline. <sup>38,39</sup> Thus, we think the Menshutkin reaction (*i.e.*, nucleophilic substitution) can afford a convenient, catalyst-free, high-efficient and atomic economic route to achieve viologen ionic polymers. Nevertheless, direct synthesis of viologen-based ionic polymers with crystalline and porous structures is still a challenging task. In aspects of heterogeneous catalysis in the CO<sub>2</sub> conversion, only a few viologen-based ionic polymers prepared by various reactions have been employed in the

catalytic CO<sub>2</sub> fixation, but which often performed under harsh conditions at high temperatures or high CO<sub>2</sub> pressures. <sup>34,35,39</sup> Therefore, it is very desirable to develop a low-cost and facile method to prepare the targeted viologen-based porous ionic polymers for the CO<sub>2</sub> conversion at a low temperature and an atmospheric pressure.

Herein, we report a facile and direct synthesis of crystalline viologen-based porous ionic polymers (denoted VIP-X, X = Cl or Br) simultaneous with enriched halogen anions and H-bonded water molecules. As depicted in Scheme 1, the targeted VIP-X were fabricated by the Menshutkin reaction of 4,4'-bipyridine (4,4'-BPY) with commercially-available linkers BCMBP or BBMBP. It is noted that some viologen-based ionic polymers synthesized by the Menshutkin reaction usually have amorphous and non-porous structures with low surface areas. <sup>44,46–48</sup> The present VIP-Cl not only has an obvious crystalline ordered structure and but also behaves a flower-like sheet morphology with considerable surface area up to 56 m<sup>2</sup> g<sup>-1</sup>. Besides, the specific contents of stable H-bonded water molecules were found in the obtained VIP-X. Though chemically absorbed water is often observed in ILs and IL-derived solid materials, <sup>10–13</sup> few attentions have paid on the valuable role of these intrinsic H-bonded water molecules for enhancing HBD-based catalysis. In this work, the optimal VIP-Br exhibited remarkable heterogeneous catalytic activities in the CO<sub>2</sub> cycloaddition with various epoxides under mild conditions without adding any co-catalysts and solvents, which attributed to the synergistic catalysis of the abundant Br<sup>-</sup> anions and HBD co-catalyst H<sub>2</sub>O.



**Scheme 1** Synthetic route to crystalline viologen-based porous ionic polymers VIP-X (X = Cl or Br) with intrinsic H-bonded water molecules via the Menshutkin reaction of 4,4'-bipyridine with (a) BCMBP and (b) BBMBP under solvothermal conditions (*i.e.*, CH<sub>3</sub>CN, 100 °C, 48 h).



## Experimental

### Materials

Raw materials 4,4'-bipyridine, 4,4'-bis(chloromethyl)biphenyl, 4,4'-bis(bromomethyl)biphenyl and the required substrates and solvents are commercially available and used without further purification.

### Methods

$^{13}\text{C}$  cross-polarization/magic angle spinning (CP/MAS) NMR spectra was measured using a Bruker AVANCE III 600 spectrometer. Liquid-state  $^1\text{H}$  spectra of the product cyclic carbonates were performed on a Bruker DPX 500 spectrometer. Fourier transform infrared spectroscopy (FTIR) was performed on a Bruker Vertex 80 V instrument. Chemical compositions and states of the samples were determined by the X-ray photoelectron spectroscopy (XPS, Thermo ESCALAB 250Xi). Electron paramagnetic resonance (EPR) spectra were recorded on a Bruker EMX-10/12 spectrometer at the X-band at room temperature. CHN elemental analysis was conducted on an elemental analyzer Vario EL cube. Thermogravimetry analysis (TGA) was measured with a TA Q50 instrument at a heating rate of  $10\text{ }^\circ\text{C min}^{-1}$  under a  $\text{N}_2$  atmosphere. Scanning electron microscope (SEM) images were performed on the HITACHI S-4800 field emission scanning electron microscope. Transmission electron microscopy (TEM) images were performed on the JEOL JEM-2100F 200 kV field-emission transmission electron microscopes. Powder X-ray Diffraction (PXRD) patterns were collected on a Bruker D8 Advance powder diffractometer.  $\text{N}_2$  sorption experiments were measured using a Quantachrome autosorb iQ2 analyzer at 77 K, and the surface area of sample was calculated using the Brunauer–Emmett–Teller (BET) method and the pore size distribution was determined by the Barrett–Joyner–Halenda (BJH) model. All the samples were degassed at  $150\text{ }^\circ\text{C}$  for 6 h in a high vacuum before analysis.

### Synthesis of viologen-based ionic polymers

Viologen-based ionic polymers were synthesized by one-step Menshutkin reaction. Typically, 4,4'-BPy (1 mmol) and BCMBP (1 mmol) were dissolved in acetonitrile ( $\text{CH}_3\text{CN}$ , 20 mL). The static reaction was taken place in a 25 mL Teflon-lined autoclave in a constant temperature oven at  $100\text{ }^\circ\text{C}$  for 48 h. After reaction, the solid product was collected by filtration and washed with  $\text{CH}_3\text{CN}$ , water and ethanol for several times. By drying under vacuum at  $80\text{ }^\circ\text{C}$  for 12 h. The resultant ionic polymer was named as VIP-Cl (yellow powder solid, yield of 86%). Similarly, VIP-Br was prepared in a similar procedure by the replacement of BCMBP with BBMBP (yellow powder solid, yield of 88%). Besides, a series of VIP-Cl and VIP-Br solid materials were prepared in various solvents including 1,4-dioxane, *N*-methyl pyrrolidone (NMP) and *N,N*-dimethylformamide (DMF).

### Cycloaddition of $\text{CO}_2$ with epoxides

Using epichlorohydrin (ECH) as a typical substrate, the catalytic  $\text{CO}_2$  fixation reaction was carried out under mild conditions. In a typical run, ECH (2 mmol) and the catalyst VIP-Br (0.05 g) were

placed in a Schlenk tube connected with a  $\text{CO}_2$  balloon (1 bar). After then, the mixture was stirred for desired time at the target temperature. After reaction, ethyl acetate (3 mL) was added to the reaction mixture and stirred for 0.5 h, the solid catalyst was separated by centrifugation. The obtained filtrate was analysed by gas chromatograph (GC) to afford the yield and selectivity of the product. For other substrates, the crude products were obtained by concentrating under reduced pressure and then were directly analysed by  $^1\text{H}$  NMR spectroscopy to determine the yields of cyclic carbonates.

For the catalyst recycling experiments, the reaction was performed under the same reaction conditions each time using the recovered catalyst from the previous run. The reusability of the catalyst was tested in five-run cycling experiments. The solid catalyst was collected by centrifuged, washed with ethanol, dried in vacuum and then charged into the next run.

## Results and discussion

### Synthesis and characterization of VIP-X

As depicted in Scheme 1, viologen-based ionic polymers VIP-X ( $\text{X} = \text{Cl}, \text{Br}$ ) were facilely synthesized *via* the Menshutkin reaction of 4,4'-BPy with BCMBP or BBMBP under solvothermal conditions. For synthesis of VIP-Cl, various solvents including  $\text{CH}_3\text{CN}$ , 1,4-dioxane, NMP and DMF were investigated to afford a series of VIP-Cl (solvent) solid materials. These VIP-Cl series are insoluble in water and common organic solvents such as ethanol, ethyl acetate,  $\text{CH}_3\text{CN}$ , tetrahydrofuran (THF), DMF and dimethyl sulfoxide (DMSO), which were different from previously reported water or solvent-soluble viologen-based ionic salts,<sup>49</sup> oligomers,<sup>50</sup> and linear polymers or networks.<sup>37,51,52</sup>

Remarkably, compared with crystalline structures of 4,4'-BPy and BCMBP (Fig. S1A†), XRD patterns (Fig. 1) of the formed VIP-Cl materials lose most of original sharp peaks of monomers, but still display three obvious Bragg diffraction peaks around  $10.2^\circ$ ,  $21.3^\circ$  and  $25.9^\circ$ , and also appear broad peaks between  $23\text{--}25^\circ$ , indicating VIP-Cl materials are partially-crystalline ordered

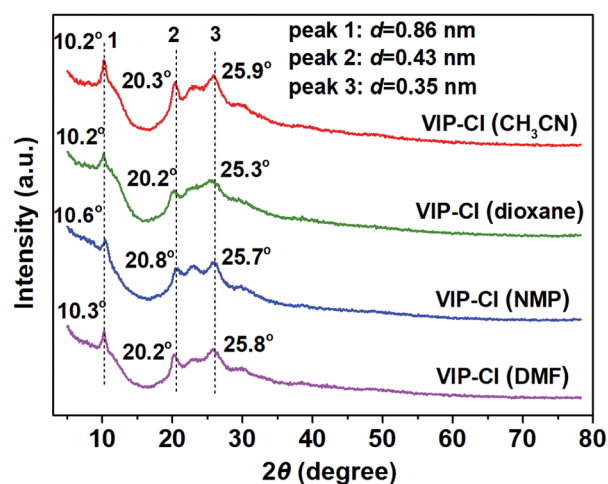


Fig. 1 XRD patterns of a series of VIP-Cl materials that were synthesized in different solvents.



polymers along with certain amorphous structures. The order crystallization is very rare in the reported viologen-based ionic polymers.<sup>38,45</sup> In detail, the sharp peak at  $10.2^\circ$  with a  $d$ -spacing value of 0.86 nm corresponds to molecular dimension of viologen moieties, which is in accordance with the molecular structure of 4,4'-BPy monomer (Fig. S2†). The raised broad peaks at around  $2\theta = 20.3^\circ$  and  $25.9^\circ$  with corresponding  $d$ -spacings of 0.43 and 0.35 nm, implying that viologen and biphenyl moieties are periodically arranged with certain spatial distances.<sup>11,53</sup> Unfortunately, our attempts to provide the most probably structure for VIP-Cl materials have failed, because of the lack of enough available cell parameters obtained from XRD patterns, which are similar to previously reported ionic covalent organic polymers.<sup>45</sup> These situations mainly due to the reason that VIP-Cl materials are partially-crystalline linear polymers but not long-range order crystalline frameworks, which are different from typical crystalline framework materials such as metal-organic frameworks and covalent organic frameworks with definite topological structures and cell parameters.<sup>24,25,29</sup>

Though the VIP-Cl samples obtained in different solvents have similar crystalline structures reflected in the XRD patterns, their porous characters are obviously different, which were determined by  $N_2$  adsorption/desorption measurements (Fig. 2 and S3†). For the samples of VIP-Cl (dioxane), VIP-Cl (NMP) and VIP-Cl (DMF), they exhibit type II isotherms (Fig. S3A†) with low  $N_2$  uptakes at 77 K, only giving low BET surface areas of  $5 \text{ m}^2 \text{ g}^{-1}$ ,  $2 \text{ m}^2 \text{ g}^{-1}$  and  $1.5 \text{ m}^2 \text{ g}^{-1}$ , respectively. Surprisingly, the sample VIP-Cl ( $\text{CH}_3\text{CN}$ ) has a larger surface area of  $56 \text{ m}^2 \text{ g}^{-1}$  and a total pore volume of  $0.33 \text{ cm}^3 \text{ g}^{-1}$ . Further, VIP-Cl ( $\text{CH}_3\text{CN}$ ) exhibit type IV isotherms (Fig. 2A) with an increasing uptake at the higher relative pressure ( $0.80 < P/P_0 < 0.99$ ) and an obvious H1 type hysteresis loop, indicative of a typical mesoporous polymer. The BJH pore size distribution (Fig. 2B) of VIP-Cl ( $\text{CH}_3\text{CN}$ ) confirm its mesoporous structure with two types of mesopores centered at 3.9 and 24.3 nm. Under the similar synthetic conditions in the above solvents, a series of VIP-Br solid materials were also obtained by replacing BCMBP with BBMBP. The sample VIP-Br prepared in  $\text{CH}_3\text{CN}$  has a moderate surface area of  $38 \text{ m}^2 \text{ g}^{-1}$  and pore volume of  $0.29 \text{ cm}^3 \text{ g}^{-1}$  (Fig. 2), which is higher than those of other VIP-Br samples (Fig. S3B†) but is a bit lower than that of VIP-Cl ( $\text{CH}_3\text{CN}$ ). These

results owe to the larger  $\text{Br}^-$  size than the one of  $\text{Cl}^-$ , which would occupy larger space nearby viologen ionic sites within the polymer. Also, the pore size distribution (Fig. 2B) of VIP-Br displays abundant smaller mesopores ranging from 3.9 to 13.4 nm by pairing with the larger  $\text{Br}^-$  anions. It is noted that VIP-Br series also have partially-crystalline structures with some observable peaks at  $10.5^\circ$ ,  $20.3^\circ$ ,  $22.6^\circ$  and  $26.0^\circ$  in the XRD patterns (Fig. S1C†), however, they display much lower degree of crystallinity than VIP-Cl series, indicating more amorphous form exists in the VIP-Br series. As a result, the samples VIP-Cl ( $\text{CH}_3\text{CN}$ ) and VIP-Br ( $\text{CH}_3\text{CN}$ ) that synthesized in  $\text{CH}_3\text{CN}$  have higher surface areas than samples prepared in other solvents, which would appear with the simplified names VIP-Cl and VIP-Br latter. To our knowledge, the prepared crystalline VIP-Cl possesses a relative high surface area of  $56 \text{ m}^2 \text{ g}^{-1}$ , superior to most of viologen-based polymers with low surface areas that prepared by the Menshutkin reaction,<sup>11,45–48</sup> comparable to viologen-based covalent organic networks that synthesized by the Zincke reaction,<sup>38–41</sup> but is still inferior to those viologen-based porous polymers that fabricated using various metal catalysts and additives<sup>32–35</sup> (see Table S1† for the detailed comparisons).

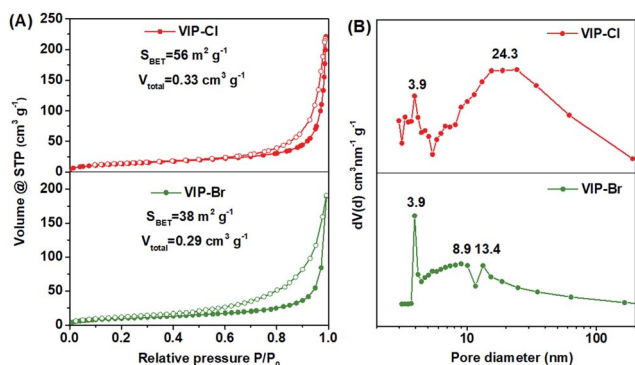


Fig. 2 (A)  $N_2$  adsorption–desorption isotherms and (B) BJH pore size distributions of VIP-Cl and VIP-Br synthesized in  $\text{CH}_3\text{CN}$ .

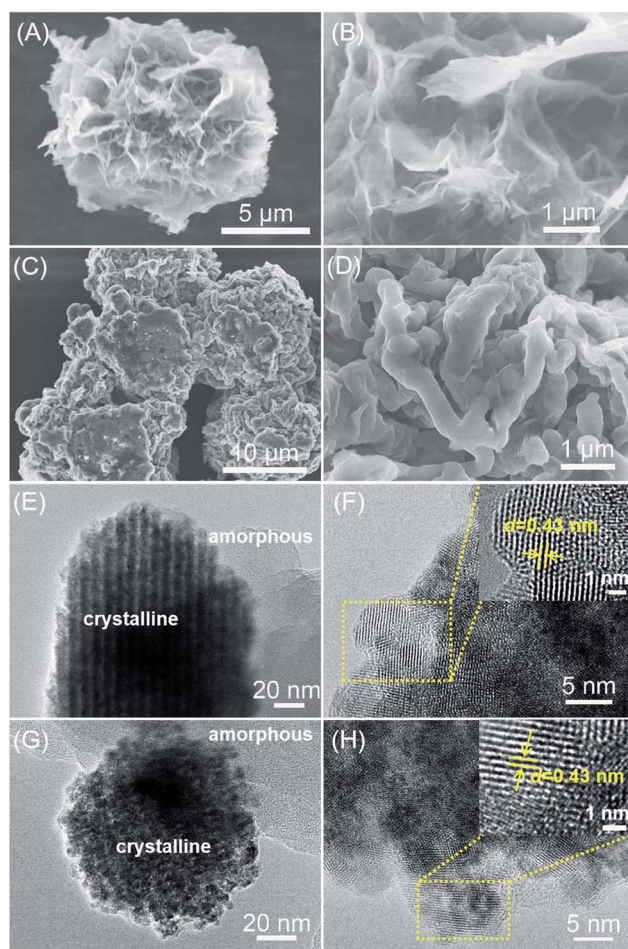


Fig. 3 SEM images of (A and B) VIP-Cl and (C and D) VIP-Br, and TEM images of (E and F) VIP-Cl and HRTEM images (G and H) VIP-Br.



The morphology of VIP-X (X = Cl or Br) was observed by the scanning electron microscopy (SEM) images. As shown in Fig. 3A–D, the sample VIP-Cl exhibits a hierarchical, flower-like ordered morphology that assembled by the interconnected nanosheets, while VIP-Br also has a flower-like shape consisted of cross-linked aggregates. TEM images exhibit that VIP-Cl and VIP-Br samples are composed of crystalline structures (the dark sections in Fig. 3E and G) and amorphous forms (the light sections in Fig. 3E and G), which are consistent with their partially-crystalline structures reflected by the XRD patterns. From the HRTEM images (Fig. 3F and H), the obvious lattice fringes with a spacing of 0.43 nm for VIP-Cl and VIP-Br were observed, which correspond to the layer *d*-spacing of 0.43 nm from the Bragg diffraction peaks at 20.3° in the XRD patterns. These ordered spatial arrays should be attributed to aromatic  $\pi$ - $\pi$  stacking interaction between the periodically repeated viologen and biphenyl units.<sup>11,53</sup>

The chemical structures and compositions of VIP-X were determined by solid-state <sup>13</sup>C NMR, FTIR, XPS and elemental analysis. In the <sup>13</sup>C NMR spectra (Fig. 4A) of VIP-Cl and VIP-Br, the obvious multi-peaks in the region of 144.1–126.1 ppm are attributable to aromatic carbon signals in the phenyl and bipyridinium rings of viologen cationic polymers.<sup>43,48</sup> The

carbon signals located at 61.0 ppm for VIP-Cl and 59.3 ppm for VIP-Br are assigned to methylene carbon atoms (–CH<sub>2</sub>–) linking the bipyridinium and benzene rings.<sup>43,48</sup> The above results of <sup>13</sup>C NMR confirm that viologen moieties and biphenyl-methylene units were successfully introduced into the polymers. In addition, FTIR spectra (Fig. 2B and C) of VIP-X were further performed to confirm the chemical structures of the targeted ionic polymers. The presence of viologen ionic moieties were demonstrated by the characteristic peaks at 1634 or 1639 and 1206 cm<sup>–1</sup> that assigned to the stretching vibrations of C=N<sup>+</sup> and C–N bonds in viologen units.<sup>32,39,48</sup> The successful incorporation of biphenyl-methylene groups was confirmed by the biphenyl aromatic C–H stretching vibrations at 3116, 3034 and 3027 cm<sup>–1</sup>, methylene C–H stretching vibrations at 2854 cm<sup>–1</sup>, the skeletal vibrations of the aromatic ring appeared at 1500–1443 cm<sup>–1</sup>, and obvious out-of-plane C–H bending vibration bands of adjacent hydrogens on aromatic ring at 789 cm<sup>–1</sup>.<sup>54</sup> No obvious wagging absorption peaks of –CH<sub>2</sub>Cl (1273 cm<sup>–1</sup>) and –CH<sub>2</sub>Br (1228 cm<sup>–1</sup>) groups,<sup>42,48,54</sup> stretching absorption peaks for C–Cl (676 cm<sup>–1</sup>) or C–Br bonds (596 cm<sup>–1</sup>) within the monomers BCMBP and BBMBP (Fig. 4B and C) were found in the formed polymers, indicating the formation of closed-loop ionic polymers with high degrees of polymerization,

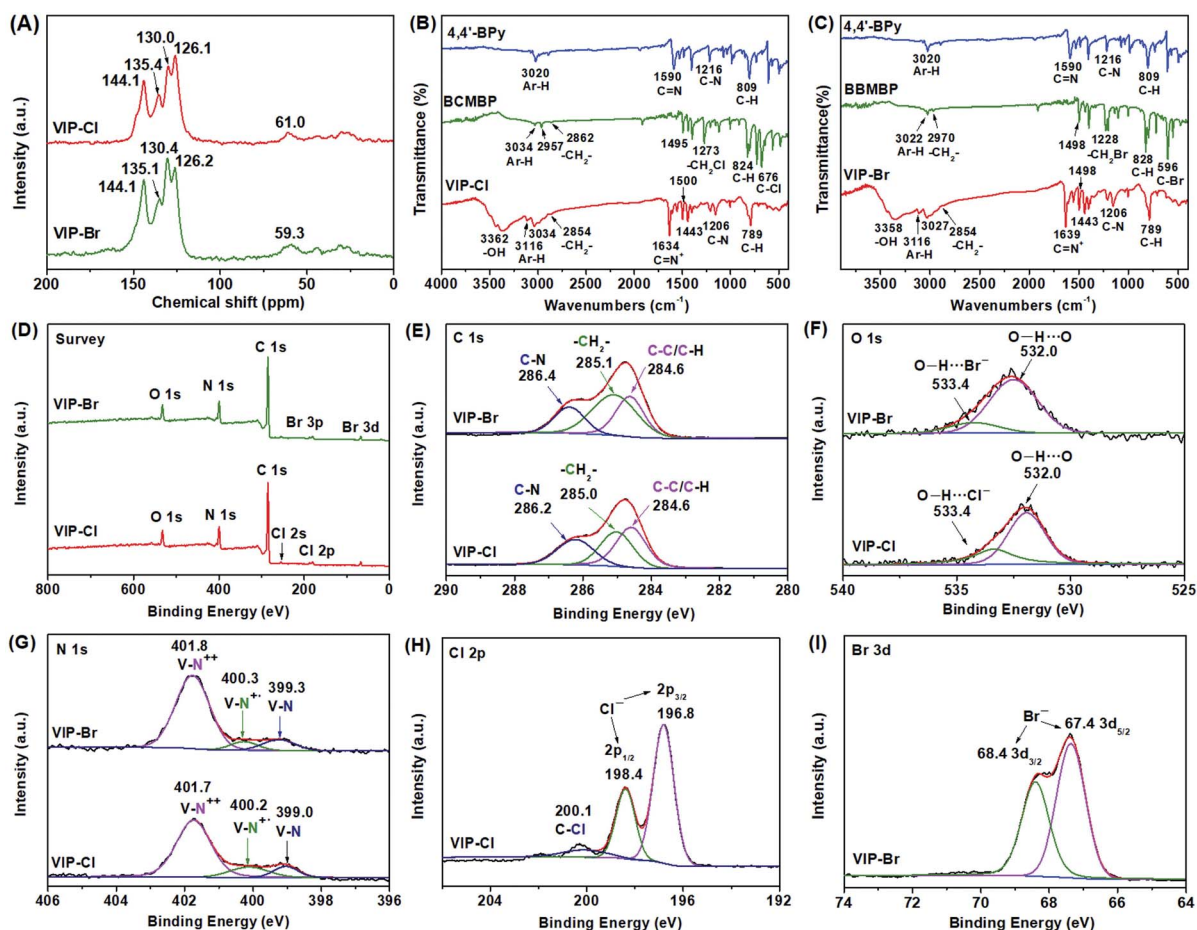


Fig. 4 Characterizations for chemical structures of VIP-Cl and VIP-Br: (A) solid-state <sup>13</sup>C CP/MAS NMR, (B) and (C) FTIR spectra, (D) XPS spectra survey, (E) C 1s, (F) O 1s, (G) N 1s, (H) Cl 2p and (I) Br 3d.



without residual monomers or  $-\text{CH}_2\text{Cl}/-\text{CH}_2\text{Br}$  end groups. Moreover, the broad peaks at 3362 and 3358  $\text{cm}^{-1}$  are assigned to H-bonded water associated with halogen anions in viologen-based ionic polymers, which are different from those of free  $-\text{OH}$  groups in unbounded water.<sup>10</sup> The elemental compositions of VIPs were also analysed by elemental analysis (EA), as listed in Table S2.† The EA results of VIP-X are well consistent with the theoretical values for the targeted closed-loop ionic polymers with repeated viologen units and H-bonded water molecules. Based on the results of EA and TGA (Fig. S4†), the contents of water are approximately 14.6 wt% for the VIP-Cl and 8.5 wt% for the VIP-Br. These desired water molecules are originated from the chemical adsorption of moisture from air under ambient conditions that enhanced by the hydrophilic ionic nature of VIP-X,<sup>11,12</sup> which are beneficial to accelerate  $\text{CO}_2$  conversion by HBD catalysis.<sup>4,14</sup>

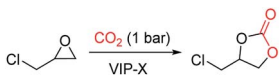
The chemical compositions of VIP-X were further determined by the XPS spectra. The full survey XPS spectra (Fig. 4D) displayed C, N, Cl or Br elements for VIP-Cl and VIP-Br. As shown in Fig. 4E, the C 1s XPS spectra are divided into three distinct peaks at 286.2/286.4, 285.0/285.1 and 284.6 eV, which are assigned to the carbon atoms of  $\text{C}=\text{N}$  bonds in the viologen groups, methylene ( $-\text{CH}_2-$ ) and aromatic rings ( $\text{C}-\text{C}/\text{C}-\text{H}$ ) in the VIP-X.<sup>32</sup> The obvious O 1s peaks (Fig. 4F) appeared at 532.0 and 533.4 eV that assigned to  $\text{O}-\text{H}\cdots\text{O}$  and  $\text{O}-\text{H}\cdots[\text{X}^-]$  ( $\text{X} = \text{Cl}$  or  $\text{Br}$ ), further confirming the presence of H-bonded water in the VIP-X.<sup>55,56</sup> The N 1s spectra for VIP-X (Fig. 4G) could be fitted to three distinct peaks. For VIP-Cl and VIP-Br, the main peaks appeared at 401.7/401.8 eV with very high fitting area ratios (76.1% and 81.4%) are originated from N atoms of dicationic viologen moieties ( $\text{V}-\text{N}^{2+}$ ),<sup>32,35,40</sup> demonstrating that VIP-X have high contents of viologen dicationic sites. Besides, very small peaks appeared at 400.2/400.3 eV with small fitting area ratios can be assigned to some reduced radical cationic viologen species ( $\text{V}-\text{N}^{\cdot+}$ ) that formed during the reaction.<sup>32,35,40</sup> However, these small contents of radical species can't be detected by the solid-state EPR spectroscopy (Fig. S5†). Meanwhile, the small peaks at 399.3/399.0 eV may be assigned to N atoms of fully reduced neutral viologens or unreacted bipyridines.<sup>22</sup> In the XPS Cl 2p spectrum (Fig. 4H) of VIP-Cl, the peaks appeared at 196.8 eV ( $2\text{p}_{3/2}$ ) and 198.4 eV ( $2\text{p}_{1/2}$ ) are assigned to free  $\text{Cl}^-$  anion.<sup>35</sup> Besides, the peak for C-Cl bond at 200.1 eV was very weak, indicating the monomer BCMBP was almost totally consumed. For the VIP-Br, the Br 3d spectrum (Fig. 4I) displays two strong peaks at 68.4 eV ( $3\text{d}_{3/2}$ ) and 67.4 eV ( $3\text{d}_{5/2}$ ), while no signal for C-Br can be observed,<sup>17,32,56</sup> suggesting that all the Br atoms are in the state of  $\text{Br}^-$  anions within the VIP-Br. These results are consistent with FTIR spectra. In other words, these formed VIP-X are highly polymerized linear polymers, not viologen-based oligomers or ionic salts, which possess high contents of ionic moieties paired with halogen anions. Further, the energy-dispersive X-ray absorption spectroscopy (EDS) elemental mapping images (Fig. S6†) describe homogeneous dispersions of the targeted C, N, Cl or Br elements within the VIP-X materials. As a result, the above full characterizations demonstrated that two viologen-based ionic polymers were successfully prepared with high degrees of polymerization and high density

of halogen anions and available H-bonded water *via* one-step Menshutkin reaction.

### Catalytic performance in $\text{CO}_2$ conversion

Owing to the enriched halogen ionic sites and desired bonded water as well as the insolubility in the reaction system, the obtained VIP-X were investigated as heterogeneous catalysts in the cycloaddition of  $\text{CO}_2$  with epoxides without adding any co-catalysts or solvents. To explore the suitable conditions for the synthesis of value-added cyclic carbonates, epichlorohydrin (ECH) was chosen to the model epoxide for this reaction (Table 1). First, the reaction was directly carried out at a relative low temperature of 60 °C using a  $\text{CO}_2$  balloon (1 bar), the catalyst VIP-Cl gives a moderate yield of 75% using 48 h but affords a desired yield of 98% using a longer time of 72 h (Table 1, entries 1 and 2), while the control catalysts VIP-Cl (dioxane), VIP-Cl (NMP) and VIP-Cl (DMF) with low surface areas give a bit lower yields of 95–96%, due to the differences of their surface areas. Another control catalyst VIP-Br achieves a high yield of 99% with an ideal selectivity only using 48 h (Table 1, entry 3). Obviously, the catalyst VIP-Br exhibits a better catalytic activity with a high TOF value of 0.453  $\text{h}^{-1}$  than those of VIP-Cl (TOF values: 0.303 and 0.264  $\text{h}^{-1}$ ) at 60 °C. Stimulated by the good catalytic behaves, the catalysts VIP-Br and VIP-Cl were further performed at near room temperature (40 °C) and room temperature (25 °C). Remarkably, the catalyst VIP-Br achieves the almost full conversion (99%) of ECH only spending 72 h and gives a high TOF of 0.302  $\text{h}^{-1}$  (Table 1, entry 4). In comparison, the catalyst VIP-Cl gives a moderate yield of 60% using 72 h (Table 1, entry 5) and can offer a considerable yield of 90% and a TOF value of 0.146  $\text{h}^{-1}$  by spending a longer time of 120 h (Table 1, entry 6). When the reactions were carried out at 25 °C, VIP-Br can give a moderate yield of 61% but VIP-Cl only shows

Table 1 Cycloaddition of  $\text{CO}_2$  with ECH catalyzed by the VIP-X under different conditions<sup>a</sup>



Entry	Catalyst	$T$ (°C)	$t$ (h)	$Y^b$ (%)	$S^b$ (%)	TOF <sup>c</sup> ( $\text{h}^{-1}$ )
1	VIP-Cl	60	48	75	99	0.303
2	VIP-Cl	60	72	98	99	0.264
3	VIP-Br	60	48	99	99	0.453
4	VIP-Br	40	72	99	99	0.302
5	VIP-Cl	40	72	60	99	0.162
6	VIP-Cl	40	120	90	99	0.146
7	VIP-Br	25	120	61	99	0.112
8	VIP-Cl	25	120	33	99	0.053

<sup>a</sup> Reaction conditions: ECH (2 mmol),  $\text{CO}_2$  balloon (1 bar), the catalyst VIP-X (0.05 g), temperature ( $T = 25-60$  °C), time ( $t = 48-120$  h).

<sup>b</sup> Yield ( $Y$ ) and selectivity ( $S$ ) of the cyclic carbonate were determined by GC and  $^1\text{H}$  NMR. <sup>c</sup> Turnover frequency (TOF) =  $[\text{mmol}(\text{product})]/[\text{mmol}(\text{V ionic content in the catalyst}) \times \text{reaction time (h)}]$ . Viologen ( $V$ ) ionic contents were calculated from EA results (see ESI Table S2).



a low yield of 33% (Table 1, entries 7 and 8). As a result, the catalyst VIP-Br can provide higher catalytic activities that reflected in yields and TOF values than those of VIP-Cl under the same conditions, which can be attributed to the higher nucleophilicity and better leaving ability of the Br<sup>-</sup> anion to Cl<sup>-</sup> anion.<sup>39,57</sup> By full evaluation of the catalytic activities in TOF values, VIP-Br is considered as the ideal heterogeneous catalyst that can achieve the high yield (99%) and the highest TOF value (0.453 h<sup>-1</sup>) under the optimal condition (*i.e.*, 1 bar CO<sub>2</sub>, 60 °C, 48 h). These satisfactory catalytic results are superior to many reported metal-free ionic polymers (IPs)<sup>15,16,27,28,34,35</sup> and even surpass to some IPs with electrophiles (metal sites)<sup>58–61</sup> or task-specific HBD groups (such as hydroxyl,<sup>28,57,62</sup> silanols,<sup>17,32,39</sup> carboxylic acids,<sup>63</sup> and ureas<sup>18</sup>) (see Table S3† for details).

Besides, it is worthy to note that the HBD cocatalyst H-bonded water molecules in VIP-Br play a vital role in activating and enhancing the CO<sub>2</sub>/ECH coupling by forming H-bonds with the oxygen atom of ECH.<sup>4,14,64,65</sup> Fortunately, the high selectivity (99%) of the product cyclic carbonate were still achieved together with high yields catalyzed by the inherent water-contained VIP-Br, indicating that no byproduct diol was formed *via* a possible hydrolysis process. In this work, no additional water was considered to enhance the conversion of the epoxide, because the selectivity for the product would decrease due to the ring opening of ECH to 3-chloro-1,2-propanediol by hot water.<sup>32,66,67</sup> The suitable amount of inbuilt H-bonded water in the catalyst not only can enhance catalytic activities by synergistic catalysis with nucleophilic Br<sup>-</sup> anions but also has no effect on the high selective synthesis of cyclic carbonates.

The structural stability and reusability of heterogeneous catalysts are two key factors that can determine their prospects in practical applications. As shown in Fig. 5, a five-cycle experiments were conducted to assess the catalytic reusability of VIP-Br in the conversion of CO<sub>2</sub> with ECH at 60 °C for 48 h. No

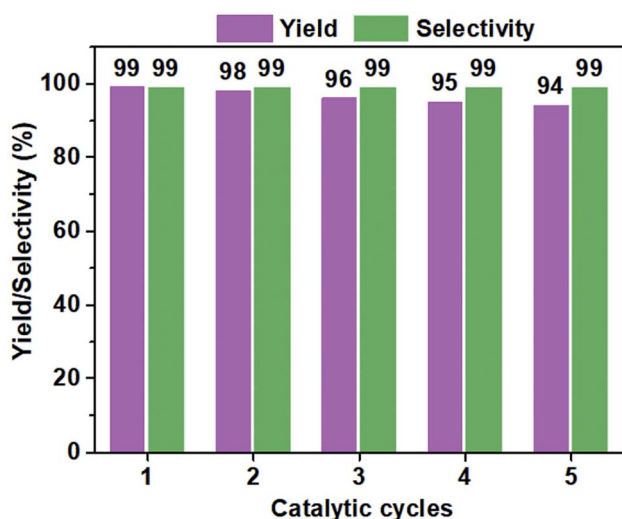


Fig. 5 A five-cycle assessment in the catalytic reusability of VIP-Br for the CO<sub>2</sub> conversion with ECH. Reaction conditions: ECH (2 mmol), CO<sub>2</sub> pressure (1 bar), the catalyst (0.05 g), 60 °C, 48 h.

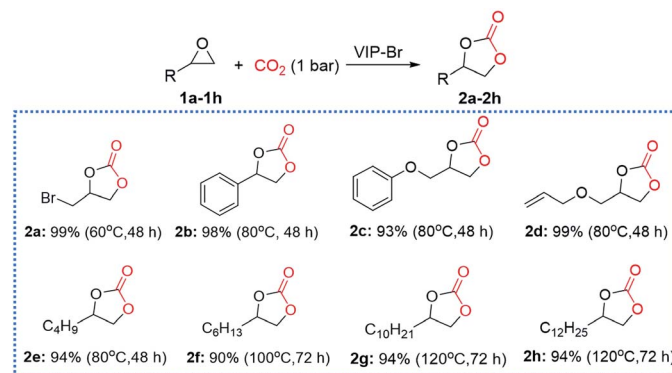


Fig. 6 Substrate compatibility of various epoxides for the CO<sub>2</sub> cycloaddition over the catalyst VIP-Br. Reaction conditions: substrate (2.0 mmol), catalyst (0.05 g), CO<sub>2</sub> (1 bar), 60–120 °C, 48–72 h.

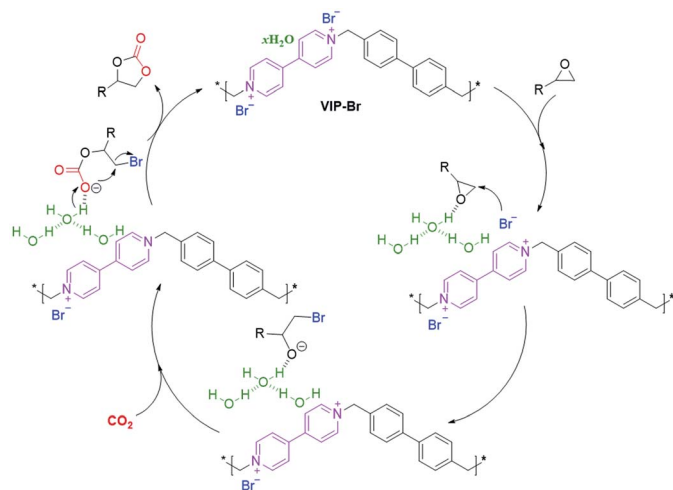
obvious decrease in both the yield and selectivity were observed, owing to its well-preserved chemical structure and composition, and morphology of the recovered catalyst, which were confirmed by its similar results of elemental analysis (C%, 53.1; H%, 4.8; N%, 5.2), FTIR (Fig. S7†) and SEM image (Fig. S8†) of the reused VIP-Br. These above results demonstrate the excellent stability and reusability of the catalyst VIP-Br.

Encouraged by the good catalytic activity and stability of VIP-Br, the substrate compatibility of various epoxides was further explored under atmospheric pressure (1 bar CO<sub>2</sub>). As shown in Fig. 6, many epoxides (1a–1h) could be smoothly converted into the corresponding cyclic carbonates (2a–2h) with high yields that determined by <sup>1</sup>H NMR analyses of the crude products (see Fig. S9–S16†). In detail, the epibromohydrin (1a) could be easily converted into the product 2a with a high yield of 99% under very mild conditions (*i.e.*, 60 °C, 48 h). The conversion of styrene oxide (2b), glycidyl phenyl ether (2c) and allyl glycidyl ether (2d) could be well accomplished with high yields of 93–99% at 80 °C for 48 h, which were required high temperatures (100–140 °C) using various other heterogeneous catalysts.<sup>15,16,24</sup> Remarkably, the extreme inert long carbon-chain alkyl epoxides could be also converted into the corresponding cyclic carbonates (2e–2h) with satisfactory yields (90–94%) under atmospheric pressure using different temperatures and times. We noted that the longer carbon-chain alkyl epoxides (2g and 2h) required a relatively high temperature of 120 °C and longer time (72 h) than the shorter ones (2e and 2f), which ascribed to the increased mass-transfer resistance of these large-sized hydrophobic substrates. In a word, the present heterogeneous catalyst VIP-Br behaves favourable substrate compatibility under mild conditions.

### Insights into the catalytic behavior

Previous works have revealed that the combination of halogen anions and hydrogen bond donors –OH groups can enhance the activation and conversion of CO<sub>2</sub> with epoxides.<sup>14,28,57,68</sup> Obviously, the present VIP-Br has both intrinsic viologen ionic sites with nucleophilic Br<sup>-</sup> anions and H-bonded water molecules, which makes for the good catalytic performance by the synergistic catalysis effect.<sup>64,65</sup> Based on the above experimental





**Scheme 2** A proposed catalytic reaction mechanism for the cycloaddition of CO<sub>2</sub> with epoxides over the catalyst VIP-Br with Br<sup>−</sup> anions and H-bonded water molecules.

results and previous reports, a plausible catalytic reaction mechanism is proposed for the cycloaddition of CO<sub>2</sub> with epoxides over the catalyst VIP-Br (Scheme 2). First, the epoxide substrate was electrophilically activated by the hydrogen-bonding interaction between the oxygen atoms from epoxides and the free –OH groups within H-bonded water, thus inducing the ring opening of the epoxide *via* the C–O bond polarization.<sup>65,69</sup> Second, the nearby nucleophilic Br<sup>−</sup> anion will attack the less hindered β-carbon atom of the activated epoxide, leading to the formation of an oxyanion intermediate that can be stabilized by the available multiple –OH groups from the H-bonded water.<sup>69,70</sup> Subsequently, the insertion of CO<sub>2</sub> into the oxyanion intermediate will create the OH-stabilized alkylcarbonate anion intermediate.<sup>17,32,69</sup> At last, the intramolecular ring-closing of the above alkylcarbonate anion intermediate can afford the cyclic carbonate, along with the release of Br<sup>−</sup> anions and regeneration of the catalyst.

## Conclusions

In summary, a series of viologen-based porous ionic polymers were constructed by a one-pot facile Menshutkin reaction. The obtained VIP-X have special crystalline ordered structures and flower-like porous structures with considerable surface area up to 56 m<sup>2</sup> g<sup>−1</sup>, and abundant halogen anions-paired viologen ionic sites and useful H-bonded water molecules. The typical VIP-Br exhibits promising heterogeneous catalytic activities in the CO<sub>2</sub> conversion with various epoxides into cyclic carbonates under mild conditions, which attributes to the synergistic catalysis of nucleophilic Br<sup>−</sup> anions and HBD co-catalyst H-bonded water. We predict that more task-specific porous ionic polymers will be developed based on this facile and catalyst-free strategy towards diverse targeted applications.

## Conflicts of interest

There are no conflicts to declare.

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