Carbon capture is essential for mitigating carbon dioxide emissions. Compared to conventional chemical scrubbing, electrochemically mediated carbon capture utilizing redox-active sorbents such as quinones is emerging as a more versatile and economical alternative. However, the practicality of such systems is hindered by the requirement of toxic, flammable organic electrolytes or often costly ionic liquids. Herein, we demonstrate that rationally designed aqueous electrolytes with high salt concentration can effectively resolve the incompatibility between aqueous environments and quinone electrochemistry for carbon capture, eliminating the safety, toxicity, and at least partially the cost concerns in previous studies. Salt-concentrated aqueous media also offer distinct advantages including extended electrochemical window, high carbon dioxide activity, significantly reduced evaporative loss and material dissolution, and importantly, greatly suppressed competing reactions including under simulated flue gas. Correspondingly, we achieve continuous carbon capture-release operations with outstanding capacity, stability, efficiency and electrokinetics, advancing electrochemical carbon separation further towards practical applications.
Anthropogenic carbon dioxide (CO₂) emissions present a serious challenge to our society. One of the foremost mitigation strategies involves carbon capture, particularly from large stationary emission sources, followed by sequestration or utilization. The incumbent technology for carbon capture is wet chemical scrubbing, which utilizes aqueous amines to chemically absorb CO₂ and subsequently release it by desorption via a temperature-swing process. However, this process is challenged by the substantial energy demand associated with the regeneration step, sorbent degradation, corrosion, environmental concerns, and large footprint. To this end, emerging materials such as water-lean solvents and porous solid sorbents have also been garnering widespread attention, with many showing promising capture properties. Nevertheless, most of these CO₂ storing materials are still reliant on high pressure and/or temperature-swing conditions during their operation cycles. Electrochemically mediated separations offer a low-temperature, ambient-pressure alternative for carbon capture, representing a promising yet largely unchartered research area. While it may be nearly doubles with respect to that under N₂ at the first reduction potential (Fig. 1b, CO₂), indicating the interaction between reduced quinones and CO₂. The complexation of CO₂ with quinone is believed to proceed via an ECEC mechanism (E, electron transfer; C, chemical reaction). Namely, the reduction of AQ affords AQ⁺, which can react chemically with CO₂ through nucleophilic addition; the resulting carbonate renders the quinone aromatic ring relatively neutral such that the second electron transfer can occur at nearly the same potential, generating one additional nucleophilic oxygen for CO₂ attachment (Supplementary Fig. 2).

An aqueous electrolyte is the ideal medium for large-scale electrochemical systems, offering distinct advantages in cost, safety, environmental benignity, and ease of device construction. Carbon capture via redox-active sorbents in aqueous electrolytes is, however, confronted with serious challenges. One critical limitation originates from the narrow electrochemical stability window of water, which is insufficient to support the reductive addition of CO₂ to quinone molecules without incurring parasitic reactions (hydrogen evolution). Compared to aprotic solvents, aqueous media are complicated by the speciation of CO₂ upon dissolution, which can undermine the activity of dissolved molecular CO₂ given inappropriate solution pH. Importantly, various competing reactions with reduced quinone molecules are expected in aqueous solutions, particularly protonation and ion association, hindering severely the kinetics and efficiency of carbon capture. Finally, rapid evaporative loss is a long-standing concern when interfacing aqueous electrolytes with continuous gas flow, which can seriously compromise the service life of the electrochemical systems.

Herein, by drawing inspiration from the recent breakthrough in salt-concentrated electrolytes for battery applications, we develop an aqueous electrolyte formulation that is fully compatible with quinone-mediated electrochemical carbon capture. Salt-concentrated aqueous electrolytes, also known as “water-in-salt” electrolytes, are defined as those in which dissolved salt outnumbers water by both mass and volume. In these solutions, the number of water molecules is insufficient to fully solvate the cations so that no free-state water exists, which can effectively suppress the activity of water molecules to extend the electrochemical stability window. By selecting appropriate salt species, desirable solution pH can be realized so that the total dissolved inorganic carbon is composed primarily of molecular CO₂. It is worthwhile highlighting that, compared to their dilute counterparts, reduced quinone diions are thermodynamically more reactive towards CO₂ and less prone to competing reactions in salt-concentrated electrolytes, due to the modified dielectric environment. This results in enhanced carbon capture kinetics and capacity, as verified by both experiments and theoretical calculations. Additional technical superiorities, including high water retention under continuous gas sparging, and diminished dissolution of electrode-immobilized quinone molecules compared to organic electrolytes, are also observed when using our salt-concentrated electrolyte system. These multifold advantages enable the demonstration of quinone-mediated CO₂ capture-release with excellent capacity, kinetics, reversibility and stability in aqueous media, advancing electrochemical carbon separation further towards practical applications.

Results

Electrochemistry of quinone-mediated CO₂ capture. Figure 1a shows the overall reaction of quinone-mediated CO₂ capture and release, using anthraquinone (AQ) as an example. In aprotic media absent of electrophiles, quinones typically undergo two successive one-electron reductions (Fig. 1b, N₂, and further illustrated in Supplementary Fig. 1), corresponding to the reversible formation of semiquinone radical anion (AQ⁻⁺) and quinone dianion (AQ⁻²). Interestingly, only a single reduction wave can be observed in the presence of electrophilic CO₂ and the current nearly doubles with respect to that under N₂ at the first reduction potential (Fig. 1b, CO₂), indicating the interaction between reduced quinones and CO₂. The complexation of CO₂ with quinone is believed to proceed via an ECEC mechanism (E, electron transfer; C, chemical reaction). Namely, the reduction of AQ affords AQ⁺, which can react chemically with CO₂ through nucleophilic addition; the resulting carbonate renders the quinone aromatic ring relatively neutral such that the second electron transfer can occur at nearly the same potential, generating one additional nucleophilic oxygen for CO₂ attachment (Supplementary Fig. 2).

Similar electrochemical behavior is also commonly observed in protic electrolytes, where reduced quinones can react with protons to form hydroxyl groups via a two-electron reduction process. Correspondingly, one immediate hurdle when...
attempting to carry out quinone-mediated carbon capture in aqueous solutions is the rapid protonation of the reduced quinone molecules (Fig. 1c). For example, the rate constant for AQ\(^{2-}\) protonation was determined to be twice that of the reaction with CO\(_2\), such that full utilization of the carbon capture capacity is hardly achievable in the presence of proton donors\(^{27}\). Moreover, since both the tendency for quinone protonation and the activity of dissolved molecular CO\(_2\) increase with increasing solution acidity, an obvious dilemma is presented for aqueous electrolyte formulation. In addition to protonation through the formation of covalent bonds, cations in the supporting electrolyte can also stabilize quinone anions via Coulombic ion pairing (Fig. 1c), which may further reduce the driving force for CO\(_2\) complexation\(^{29}\).

Salt-concentrated electrolytes, recently gaining momentum in battery research, are proposed to offer an extended electrochemical stability window by eliminating the presence of free-state water through cation solvation\(^{26,30-32}\). For example, when the concentration of LiTFSI was increased to an ultrahigh level of 20m, the cathodic decomposition potential of the electrolyte was shifted to \(-1.11\) V vs. Ag/AgCl, which is sufficient to envelop the electrochemical capture and release of CO\(_2\) by quinones.

To ensure high solubility, salt species with stable and charge-delocalized anions, such as acetate, TFSI, trifluoromethanesulfonate, and bis(fluorosulfonyl)imide, are preferred when formulating water-in-salt electrolytes\(^{33}\). Nevertheless, not all of these salts are compatible with electrochemically mediated carbon capture. Gaseous CO\(_2\) has many possible transformations upon dissolution in water, yet only dissolved molecular CO\(_2\)(aq) is active towards complexation with reduced quinones (Fig. 2b). Therefore, electrolytes for electrochemically mediated carbon capture should have a pH value of less than 6 to ensure the dominance of CO\(_2\)(aq)\(^{15}\). The alkaline nature of acetate anion, being the conjugate base of a weak acid, precludes its use. For this proof of concept study, LiTFSI was selected as it is relatively well studied and satisfies this criterion, having a pH near 4 at both 1m and 20m. The water-in-salt definition applies at concentrations above 5m, and at 20m, the ratio between Li\(^+\) and H\(_2\)O is only \(~1:2.78\).

To verify the feasibility of quinone-mediated CO\(_2\) capture in salt-concentrated aqueous media, CV scans were conducted using poly(1,4-anthraquinone)-carbon nanotube (PAQ-CNT) composite under both N\(_2\) and CO\(_2\) atmospheres. PAQ was selected for the reduced solubility compared to its molecular counterpart. The immobilization of PAQ on CNT through π–π interaction provides stable anchoring as well as fast electrical path

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**Aqueous electrolyte for quinone-mediated CO\(_2\) capture.** The prerequisite for quinone-mediated CO\(_2\) capture-release in aqueous media is a wide electrochemical stability window of the electrolytes. Since only reduced quinones with sufficient electron density can undergo nucleophilic addition with molecular CO\(_2\), the formation of quinone-CO\(_2\) adducts generally occurs at potentials more negative than the reductive decomposition (hydrogen evolution reaction) potential of water\(^{35}\). Figure 2a compares the cyclic voltammogram (CV) of AQ under CO\(_2\) with the stability window of 1m LiTFSI aqueous solution (where m is molality and LiTFSI represents lithium bis(trifluoromethanesulfonyl)imide). The reduction peak of AQ occurred at \(-0.88\) V vs. Ag/AgCl, whereas the electrolyte started to decompose as early as \(-0.81\) V vs. Ag/AgCl (defined as the potential where the current density reaches 0.01 mA cm\(^{-2}\)).
Minimized competing reactions in salt-concentrated electrolyte. To understand the effect of possible competing reactions on the reactivity of PAQ with CO₂, Fig. 3a shows the CV scans of PAQ under N₂ in 1m NH₄NO₃ (pH adjusted to 4 with HNO₃) and LiTFSI electrolytes with different molalities (pH ~ 4). It was previously established that protonation and cation association can stabilize reduced quinone species, inducing anodic shifts in the quinone reduction potential [28]. The reduction potential of PAQ in 1m NH₄NO₃ is significantly more positive than those in LiTFSI-based electrolytes. As NH₄⁺ is generally regarded as a bulky cation with low tendency for anion association, the large anodic shift in the reduction potential in 1m NH₄NO₃ can only be ascribed to the protonation of reduced PAQ [36]. The protonation of reduced PAQ in 1m NH₄NO₃ electrolyte severely undermines its potential carbon capture ability, which can be corroborated by ill-defined adduct oxidation peaks in CV scans under CO₂ (Supplementary Fig. 7).

Protonation is greatly suppressed in LiTFSI-based electrolytes where Li⁺ significantly outnumbers H⁺, rendering ion association the major competing reaction. Notably, though, both the onset and the peak potential of PAQ reduction shift towards more negative values with increasing LiTFSI concentrations (Fig. 3a), indicating weakened ion association. This is likely attributed to the dominance of associated salt species (contact ion pairs, CIP, and aggregates) in concentrated solutions as a result of incomplete cation solvation by water molecules [26,31]. To confirm this mechanistic interpretation, the free energy change (ΔG) for LiTFSI dissociation was calculated by DFT, where the increase in salt concentration can be modeled as a decrease in the dielectric constant of the electrolyte (Fig. 3b, Supplementary Fig. 8) [37]. The DFT results confirmed that the dissociation of LiTFSI becomes nonspontaneous (ΔG > 0) above a certain concentration threshold, such that the kinetics of Li⁺-involved processes are necessarily slowed down in a salt-concentrated environment. More importantly, the thermodynamic driving force for quinone-CO₂ adduct formation, from both AQ²⁻ and AQ₂⁻ complexed...
with Li\(^+\), also increases with increasing electrolyte concentrations based on the DFT calculations (Fig. 3c).

This conclusion of enhanced reactivity of CO\(_2\) with quinones in salt-concentrated media can be seen experimentally by CV measurements under different scan rates (Fig. 3d–f). Interestingly, besides the oxidation of quinone-CO\(_2\) adduct, an additional oxidation peak at less positive potential can be seen during fast scans (shaded area in Fig. 3d–f), corresponding to the oxidation of quinone dianions that are unreacted with CO\(_2\). Such a phenomenon is caused by the much more rapid electrochemical regeneration of reduced quinones compared to the relatively slow CO\(_2\) transport and chemical reaction with reduced quinones, which inevitably becomes more pronounced with increasing scan rates. Quantitatively, while 100% of the electrochemically generated quinone dianions reacted with CO\(_2\) at a scan rate of 100 mV s\(^{-1}\) in LiTFSI-based electrolytes, the value was only 77.5% when using 1m NH\(_4\)NO\(_3\) likely due to the rapid protonation; and compared to 1m LiTFSI, a higher percentage of adduct formation was observed in 20m LiTFSI under all scan rates, thanks to the suppressed Li\(^+\) association (Supplementary Fig. 9).

Therefore, it is clear from the above discussion that the reductive addition of CO\(_2\) to quinones is much less susceptible to competing reactions when carried out in salt-concentrated aqueous media, making them particularly attractive for quinone-mediated carbon capture.

Physicochemical advantages of salt-concentrated electrolyte. In addition to being fully compatible with quinone electrochemistry and minimizing competing reactions, concentrated LiTFSI aqueous electrolytes also possess favorable physicochemical properties for the practical applications of electrochemically mediated carbon capture.

Firstly, salt-concentrated aqueous media demonstrate significantly enhanced resistance to evaporative loss compared to their dilute counterparts. Figure 4a plots the weight evolution of LiTFSI-based electrolytes under continuous, high-flow-rate gas sparging, where a stark contrast in water retention capability exists between the dilute (pure H\(_2\)O, 1m LiTFSI) and the concentrated (10 and 20m LiTFSI) electrolytes. While pure water suffered from >56% weight loss (with respect to the initial electrolyte weight) after 168 h of gas purging, the value was merely 1.3% for 20m LiTFSI. This unique feature effectively dispels the traditional concern of electrolyte dry-out when interfacing aqueous solutions with gas flows, rendering the potential quinone-mediated carbon capture system with flexibility in form factor and durability.

Secondly, salt-concentrated aqueous media containing no free water inhibit the dissolution of electroactive compounds immobilized on heterogeneous substrates, which is desirable for capacity retention during electrochemical cycling of many systems. Quinones, especially in their reduced forms, generally show high dissolution tendency in polar electrolytes\(^{24,35,38}\). Correspondingly,
hydroquinone was selected as a probe molecule, and its solubility in various electrolytes was determined via UV–Vis spectrometry (Fig. 4b, Supplementary Figs. 10 and 11). A high hydroquinone solubility of 463 mg mL⁻¹ was observed in dimethyl sulfoxide (DMSO) which is a representative organic solvent, resulting in visible PAQ dissolution and rapid capacity decay during electrochemical cycling (Fig. 4c and Supplementary Fig. 12). The solubility was decreased by fourfold when the organic solvent was replaced with water, yet the value remained significantly lower than in the previously reported quinone-mediated carbon capture systems. Moreover, the value is on par with, if not higher than, the thermal release in efficiency, the carbon capture capacity of PAQ is ~8.9 mmol g⁻¹, which is close to its theoretical value of 9.7 mmol g⁻¹.

Figure 5a shows the electrochemical CO₂ capture-release voltage profiles at different cycling currents in 20m LiTFSI electrolyte, where the PAQ electrode demonstrated excellent charge transfer kinetics. When cycled at a high current of 0.5 A g⁻¹ (capture/release in ~30 min), a carbon capture capacity of ~8.4 mmol g⁻¹ can be achieved (~86% of the theoretical value), which is higher than in the previously reported quinone-mediated carbon capture systems. Moreover, the value is on par with, if not higher than, the established amine-based processes (~8 mmol per gram of amine, but ~1–2 mmol per gram of the total scrubbing absorbent)²⁴,²⁵. Importantly, the capacity was only affected slightly with further increase in the cycling currents. For example, a working capacity of ~6.6 mmol g⁻¹ could still be obtained when the capture/release process was completed in only ~6 min (2 A g⁻¹). The quinone-mediated CO₂ capture-release also demonstrated outstanding cycling stability and reversibility in salt-concentrated aqueous electrolytes (Fig. 5b). Minimal capacity loss can be observed after 150 cycles at a high current of 2 A g⁻¹, with an average Coulombic efficiency (release capacity/capture capacity) of near unity.

To gain deeper insights into the electrochemical CO₂ capture-release performance in a practical scenario, a prototypical membrane-based flow cell was constructed as shown schematically in Fig. 5c. LiFePO₄ was chosen as the counter electrode due to its well-established Li⁺ intercalation–deintercalation electrochemistry.
in aqueous environments (~0.42 V vs. Ag/AgCl in 20m LiTFSI) with high reversibility and excellent electrokinetics39,40. The capacity of the LiFePO4 electrode was at least four times in excess of that of the PAQ-CNT electrode to minimize the polarization from the counter electrode. The shortened ion-transport distance between the electrodes and the continuous delivery of gas-phase CO2 enabled high-current carbon capture-release with increased active material mass loading41. At a current of 2 A g−1 and a mass loading of 1 mg cm−2, the membrane flow cell demonstrated over 100 cycles of continuous carbon capture and release with good capacity retention (Fig. 5d), highlighting the outstanding capacity, stability, and reversibility of quinone-mediated carbon capture in salt-concentrated aqueous media.

**CO2 response during electrochemical quinone cycling.** Finally, we constructed a gas device to directly monitor the CO2 response in situ during quinone electrochemical cycling. The experimental setup is shown in Fig. 6a. Briefly, the PAQ-LiFePO4 electrochemical cell was mounted on a gasket with CO2 flowing on one side. Gaseous CO2 can dissolve into the electrolyte, transport through the electrochemical cell via diffusion and outgas into the N2 sweep stream on the other side. The permeated CO2 was then carried towards an in-line gas analyzer to determine the concentration. The detailed configuration of the gas cell is provided as Supplementary Fig. 15.

Figure 6b shows the CO2 concentration response of the gas device during constant voltage capture and release (each capture/release step was 10 min, followed by a 10-min rest), where a cyclic fluctuation of the permeated CO2 concentration can indeed be clearly observed. During capture, CO2 dissolved in the electrolyte was being actively consumed by the PAQ-CNT electrode, resulting in a continuous decrease in CO2 permeation; and the subsequent electrochemical oxidation of the PAQ-CO2 adduct generated CO2 rapidly, leading to a sharp concentration spike until all the captured CO2 was released and the permeation restored to its background value (Fig. 6c). Note that the discrepancy between the onset of the electrochemical capture/release and the CO2 signal response was due to system dispersion and the headspace volume of the flow chamber. Importantly, the consistent CO2 response over repeated electrochemical cycling further confirmed the stability of quinone chemistry for carbon capture in salt-concentrated aqueous media.

**System testing in the presence of oxygen.** The potential parasitic reaction (peroxide formation) between reduced quinones and molecular oxygen (O2) present in the feed stream is generally regarded as a source of inefficiency in our described CO2 separation scheme13,17. Therefore, the electrochemical behavior of quinones in salt-concentrated aqueous electrolytes was further evaluated under mixed CO2/O2 conditions.

It is worth emphasizing that electrochemically mediated carbon capture operates in a mass transfer-limited regime. Therefore, we can take advantage of the substantial solubility difference between O2 and CO2 to reduce the efficiency loss...
aqueous solutions at 25 °C, CO₂ has ~26 times higher solubility than O₂ under the same partial pressure. While application for dilute scenarios such as direct air capture requires further investigation, we believe redox-active sorbents are competitive for carbon capture from stationary sources with relatively high CO₂ partial pressure, such as carbon capture from power plants and large industrial processes (steel, cement and fertilizer plants, etc.), as well as in nonoxidizing environments (natural gas and biogas upgrading, etc.), which account for ~60% of the total global greenhouse gas emissions. Flue gas from coal-fired power plants typically contains 12–15% CO₂ and 3–5% O₂; in other industrial plants, the CO₂ concentration in exhaust streams can be even higher (e.g. cement production 15–30%, iron and steel 20–44%). Therefore, the limiting current involving CO₂ can be two orders of magnitude higher than that involving O₂ in our targeted carbon capture scenarios.

The potential of the O₂/O₂^2− redox couple in 1 and 20m LiTFSI was studied using a glassy carbon electrode, which is known to catalyze the two-electron oxygen reduction reaction (ORR, Fig. 7a). The ORR onset potential was relatively negative for both cases, which can be attributed to the limited proton availability. The reaction product is likely to be Li₂O₂, which has been observed to be stable in salt-concentrated aqueous media due to significantly reduced H₂O activity, making it a poor oxidant. Importantly, compared to 1m LiTFSI, the onset potential was 150 mV more negative in 20m LiTFSI together with reduced limiting current, further lowering the driving force for parasitic ORR. No obvious differences in CV can be observed for PAQ in 20m LiTFSI under N₂ and O₂ (Supplementary Fig. 16a). The ratio between the anodic and cathodic peak currents (Ipa/Ipc) is 0.98 and 0.97 under N₂ and O₂ respectively, confirming the relatively low sensitivity of reduced PAQ towards O₂ in 20m LiTFSI. Redox-active sorbents with higher reduction potentials (e.g. 1,4-naphthoquinone, NQ, Supplementary Fig. 16b) can also be chosen to minimize side reactions.

Importantly, the quinone-CO₂ adducts are much more stable against oxidation compared to the corresponding quinone dianions due to their increased oxidation potential and, as described previously, the oxidation potential also increases with increasing LiTFSI concentration (Fig. 2c, d and Supplementary Fig. 6). As shown in Fig. 7b, the onset potential for PAQ-CO₂ oxidation (ca. −0.26 V vs. Ag/AgCl) can become more positive than the onset potential for ORR (ca. −0.43 V vs. Ag/AgCl) in 20m LiTFSI, such that the adduct will be much less prone to react with O₂ in salt-concentrated aqueous media. This is not the case at low salt concentration. Adopting quinones with higher redox potentials can further increase the potential difference for even more stable adducts (Supplementary Fig. 17).

To further investigate the stability of the quinone-CO₂ adduct, we quantified the degree of PAQ-CO₂ oxidation in the presence of O₂. Specifically, the PAQ electrodes were first fully reduced under CO₂ and were then allowed to rest in the presence of different concentrations of O₂, after which the electrodes were reduced again to quantify the capacity loss (Fig. 7c and Supplementary Fig. 18). Merely 9.4% of the adducts were oxidized after resting in air for 15 min, which is the typical capture-release time scale for
our electro-swing CO2 separation; and even when rested in pure O2, only 23.2% of the capacity was lost after 15 min. Moreover, in the presence of 3% O2, which is the typical concentration in coal-fired power plant flue gas, 90% of the capacity was maintained after 1 h. While further improvements will be required for full practical implementation, these results demonstrate a promising pathway to minimizing the effects of oxygen on quinone capture processes.

Figure 7d shows the CV scans of PAQ under pure CO2 and simulated flue gas (15% CO2 with 3% O2). To compare the efficiency under the two scenarios, we calculated the ratio between the total anodic and cathodic capacities (Qa/Qc). At a scan rate of 10 mV s\(^{-1}\), the value was 87.3% under pure CO2 and decreased slightly to 80.7% under simulated flue gas. At an even slower scan rate of 5 mV s\(^{-1}\) to ensure the complete reaction between PAQ dianions and CO2, a high Qa/Qc of 83.6% can be achieved (Supplementary Fig. 19a). Replacing PAQ with NQ afforded further improved efficiency (Supplementary Fig. 19b).

When constant current CO2 capture-release cycling was conducted under simulated flue gas in the membrane flow cell, the capture-release capacity of PAQ during continuous cycling. The cycling was carried out in the membrane flow cell configuration at a current density of 0.5 mA cm\(^{-2}\) and a PAQ mass loading of 0.5 mg cm\(^{-2}\) (immobilized on carbon felt). CC capture and CC/CV release protocol was adopted with the cut-off voltage being −1.35 and −0.1 V, respectively. Each capture/release step has a time scale of ~15 min. Source data are provided as a Source Data file.

**Figure 7** Electrochemical CO2 capture-release in the presence of O2. a Linear sweep voltammograms of 1 and 20m LiTFSI saturated with O2 measured with a glassy carbon electrode at a rate of 10 mV s\(^{-1}\). b Comparison between the oxidation potential of PAQ-CO2 (with onset potential noted) and the ORR potential (from a) in 1 and 20m LiTFSI. c Reduction profiles of PAQ under CO2 in 20m LiTFSI and of the same electrode after being rested in 100% O2 (15 min), air (15 min) and 3% O2 (1 h), respectively. The reductions were carried out at a current density of 0.5 mA cm\(^{-2}\) and the PAQ was immobilized on carbon felt at a mass loading of 1.2 mg cm\(^{-2}\). d CV scans of PAQ under pure CO2 and simulated flue gas in 20m LiTFSI. The ratio between the total anodic and cathodic capacities was 87.3% and 80.7% for pure CO2 and simulated flue gas, respectively. The additional anodic peak observed under simulated flue gas corresponds to the oxidation of quinone anions that did not react with CO2 due to mass transport limitation. For (b) and (d), the CV scans were conducted at a rate of 10 mV s\(^{-1}\) with 1.5 μg PAQ loaded on the glassy carbon electrode. e The capture-release capacity of PAQ during continuous cycling under simulated flue gas. The cycling was carried out in the membrane flow cell configuration at a current density of 0.5 mA cm\(^{-2}\) and a PAQ mass loading of 0.5 mg cm\(^{-2}\) (immobilized on carbon felt). CC capture and CC/CV release protocol was adopted with the cut-off voltage being −1.35 and −0.1 V, respectively. Each capture/release step has a time scale of ~15 min. Source data are provided as a Source Data file.
configuration, high capacity cycling with good stability was still realized (Fig. 7e) with only moderately compromised efficiency (the average Coulombic efficiency over 75 cycles was 95.5%, as shown in Supplementary Fig. 20). The electrical energy consumption of this capture-release process was calculated to be ~56 kJ mol$^{-1}$ CO$_2$ (Supplementary Note 2), which is on par with other carbon capture systems$^{24}$. A more detailed energetic analysis of quinone-mediated carbon capture in a conventional electrolyte can be found in a recent literature$^{16}$. The process energetics can be further improved with engineering optimization.

With the data above, we believe our proposed system of electrochemically mediated carbon capture in salt-concentrated aqueous media can indeed be of practical significance for carbon capture from point sources.

**Discussion**

In this work, we investigated the feasibility of employing salt-concentrated aqueous electrolytes for electrochemically mediated CO$_2$ separation using quinone chemistry. By rationally designing the electrolyte formulation, salt-concentrated aqueous electrolytes can offer an extended electrochemical stability window, high CO$_2$ activity, as well as significantly reduced active material dissolution and evaporative loss. Importantly, reduced quinone species are much less susceptible to competing reactions such as protonation and ion association in salt-concentrated media compared to their dilute counterparts due to the modified dielectric environment, which is highly desirable for quinone-CO$_2$ adduct formation. As a result, high capacity CO$_2$ capture-release cycling with excellent electrokinetics, efficiency and stability was demonstrated in both conventional single-compartment electrochemical cell and prototypical membrane-based flow cell. When tested using simulated flue gas, our proposed system maintained good performance metrics and competitive system energetics. The proved advantages of salt-concentrated aqueous electrolytes are also transferable to other types of redox-active molecular CO$_2$ sorbents. The material design demonstrated through these results advances the practicality of electrochemically mediated carbon capture with redox-active molecular CO$_2$ sorbents broadly, given that the inherent advantages are not limited to our example quinone chemistry.

**Methods**

**Materials.** The electrolytes were prepared by dissolving the corresponding salts, lithium bis(trifluoromethanesulfonfyl)imide (Solvay), and/or tetrabutylammonium hexafluorophosphate (Sigma-Aldrich) in water and/or DMSO (Sigma-Aldrich). Ferrocene and anthraquinone were purchased from Sigma-Aldrich.

**Synthesis of the PAQ-CNT electrode.** PAQ was synthesized following a previously reported procedure$^{22}$. To obtain the PAQ-CNT composite, PAQ was dissolved in chloroform (1 mg mL$^{-1}$) using a probe sonicator at 5 °C (Cole-Parmer Ultrasonic Processor, pulser mode: on 5 s, off 3 s, 60% amplitude). Subsequently, CNT was added into the solution (1 mg mL$^{-1}$), and the mixture was sonicated for another 20 min to afford a homogeneous dispersion. The PAQ-CNT ink was then cast on either glassy carbon electrode (Ø 3 mm, BASi) or 50 µm carbon felt (Filber Glast Carbon Fiber Veil).

**Characterizations.** SEM images were taken with a Zeiss SUPRA 55-VP scanning electron microscope. XPS analysis was obtained on a Thermo Scientific K-Alpha+ XPS equipped with Al (Ka) source with a spot size of 400 µm. High-resolution spectra were collected with a step size of 0.1 eV and an accumulation of ten scans. Hydroquinone solubility was determined based on UV–Vis absorption spectra (Ocean Optics). The evaporative loss rate of the electrolytes was measured by monitoring their weight change under constant N$_2$ purging. In each experiment, 5 mL electrolyte was sealed in a 20 mL glass vial with septum cap, which was equipped with a 21G needle inlet and a 21G needle outlet. Pure N$_2$ gas was bubbled directly into the electrolyte through the needle inlet at a flow rate of 200 mL min$^{-1}$.

**CO$_2$ capture/release quantification by FTIR spectroscopy.** A Nicolet 8700 FTIR spectrometer (Thermo Scientific) and a 100-mm short-path IR gas cell ( Pike Technologies, septum-sealed) were employed for the quantification of CO$_2$ capture/release by PAQ. The spectrometer chamber was under high-flow-rate N$_2$ purging during the measurements to avoid interference from atmospheric CO$_2$.

The calibration curve was made by injecting (Hamilton 1700 series gastight syringe) defined amounts of pure CO$_2$ into the electrolyte and measuring the integrated area of the two CO$_2$ asymmetric stretching bands in absorbance mode (2335 and 2360 cm$^{-1}$). For the quantification, CO$_2$ was captured electrochemically by PAQ (0.25 mg on carbon felt) via two CV reduction cycles at a slow scan rate of 5 mV s$^{-1}$. The electrode was then removed from the electrochemical cell, thoroughly dried with paper towel, and sealed in a 2 mL N$_2$ flushed glass vial with septum cap (Agilent autosampler vial). Thermal release instead of electrochemical release was chosen to avoid the outgassing of physically dissolved CO$_2$ from the electrolyte in the case of electrochemical release, which can compromise the quantification accuracy. After 30 min thermal release on a hotplate, 250 µL headspace gas was taken and injected into the N$_2$ flushed IR gas cell to obtain the corresponding IR spectrum.

**Electrochemical measurements.** All the electrochemical measurements were conducted using a VersaSTAT4 potentiostat (Princeton Applied Research) in a conventional single-compartment electrolytic cell (BASI), unless otherwise specified. The glassy carbon electrode used in this study has a diameter of 3 mm and was polished before each measurement. The LiFePO$_4$ on Al foil electrode was purchased from MTI Corp. (single side coated, active material loading 12 g cm$^{-2}$). The LiFePO$_4$ on carbon felt electrode was prepared by mixing 95% LFP (MTI Corp.), 2.5% carbon black (Super P, MTI Corp.) and 2.5% polyvinylidene fluoride (Sigma-Aldrich) in N-methylpyrrolidinozone followed by casting on 50 µm carbon felt (mass loading 8 mg cm$^{-2}$). For the construction of the PAQ-LiFePO$_4$ membrane flow cell, PAQ-CNT was immobilized on 50 µm carbon felt at a mass loading of 1 mg cm$^{-2}$ and LiFePO$_4$ on Al foil was used as the counter electrode. Both electrodes were cut to 2 cm × 2 cm in area and were separated by an electrolyte-imbibed glass fiber membrane (MilliporeSigma). A hydrophobic gas diffusion layer (Toray carbon paper, PTFE treated, TGP-H-60) was placed on one side of the PAQ electrode to interface with continuous CO$_2$ flow. The reduction of PAQ was conducted with a cut-off voltage of either −0.95 V vs. Ag/AgCl or −1.4 V vs. LiFePO$_4$ counter electrode to prevent the possible TFSI$^-$ decomposition. The oxidation of PAQ-CO$_2$ was conducted via the CC/VC mode (constant current followed by constant voltage). The cut-off voltage for CC oxidation was either 0.2 V vs. Ag/AgCl or −0.1 V vs LiFePO$_4$. CV oxidation was carried out until the current decayed to 10% of the CC oxidation current.

**In situ gas cell measurements.** Measurements on the CO$_2$ permeation response of a carbon electrode were performed using a device consisting of two gas flow chambers separated by the PAQ-LiFePO$_4$ electrochemical cell, all held together with rubber gaskets and screws (detailed gas cell configuration is shown as Supplementary Fig. 16). Both PAQ-CNT (1 mg cm$^{-2}$) and LiFePO$_4$ (8 mg cm$^{-2}$) were cast on 50 µm carbon felt, and the electrode area was 2 cm × 2 cm. The two electrodes were separated with a 25 µm polypropylene separator (Celgard 3501) infused with 20 ml LiTFSI electrolyte. The PAQ-LiFePO$_4$ electrochemical cell was mounted on a brass gasket holder with a 1 cm$^2$ circular opening, and the edges of the electrochemical cell were sealed to block gas-phase CO$_2$ transport between the two flow chambers. The brass gasket also served as the current collector for the PAQ-CNT electrode, and a titanium mesh was used as the current collector for the LiFePO$_4$ electrode. The testing employed a “zero-gap” design where the electrochemical components were all in close contact, such that CO$_2$ cross-over can only occur via dissolution-diffusion through the electrolyte. During operation, CO$_2$ flowed through one gas chamber and N$_2$ sweep gas flowed through the other. The concentration of CO$_2$ in the N$_2$ stream was measured with an in-line CO$_2$ sensor (ExproIR17-W 20% CO$_2$ sensor). The sensor has a measurement range of 0–20% CO$_2$ and an accuracy of ±70 ppm. The electrochemical cycling followed a constant voltage protocol. For each capture-release cycle, the cell was first held at −1.2 V for 10 min to capture CO$_2$, followed by release at 0 V for 10 min and a 10-min rest.

**Data availability**

The datasets generated during and/or analyzed during the current study are available from the corresponding author on reasonable request. The source data underlying Figs. 2a, c, d, 3a, d–f, 4a, c–e, 5a, b, d, 6b, 7a–e and Supplementary Figs. 17, 19, 20 and 22 are provided as a Source Data file.

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Y.L. and T.A.H. conceived the concept and designed the experiments. Y.L. carried out the experiments and H.-Z.Y. conducted the DFT calculations. T.A.H. and T.V.V. supervised the project. Y.L., K.M.D. and T.A.H. prepared the manuscript with input from all the other coauthors.

Competing interests
The authors declare no competing interests.

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