

A metal-free organic–inorganic aqueous flow battery

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As the fraction of electricity generation from intermittent renewable sources—such as solar or wind—grows, the ability to store large amounts of electrical energy is of increasing importance. Solid-electrode batteries maintain discharge at peak power for far too short a time to fully regulate wind or solar power output^{1,2}. In contrast, flow batteries can independently scale the power (electrode area) and energy (arbitrarily large storage volume) components of the system by maintaining all of the electro-active species in fluid form^{3–5}. Wide-scale utilization of flow batteries is, however, limited by the abundance and cost of these materials, particularly those using redox-active metals and precious-metal electrocatalysts^{6,7}. Here we describe a class of energy storage materials that exploits the favourable chemical and electrochemical properties of a family of molecules known as quinones. The example we demonstrate is a metal-free flow battery based on the redox chemistry of 9,10-anthraquinone-2,7-disulphonic acid (AQDS). AQDS undergoes extremely rapid and reversible two-electron two-proton reduction on a glassy carbon electrode in sulphuric acid. An aqueous flow battery with inexpensive carbon electrodes, combining the quinone/hydroquinone couple with the Br_2/Br^- redox couple, yields a peak galvanic power density exceeding 0.6 W cm^{-2} at 1.3 A cm^{-2} . Cycling of this quinone–bromide flow battery showed >99 per cent storage capacity retention per cycle. The organic anthraquinone species can be synthesized from inexpensive commodity chemicals⁸. This organic approach permits tuning of important properties such as the reduction potential and solubility by adding functional groups: for example, we demonstrate that the addition of two hydroxy groups to AQDS increases the open circuit potential of the cell by 11% and we describe a pathway for further increases in cell voltage. The use of π -aromatic redox-active organic molecules instead of redox-active metals represents a new and promising direction for realizing massive electrical energy storage at greatly reduced cost.

Solutions of AQDS in sulphuric acid (negative side) and Br_2 in HBr (positive side) were pumped through a flow cell as shown schematically in Fig. 1a. The quinone–bromide flow battery (QBFB) was constructed using a Nafion 212 membrane sandwiched between Toray carbon paper electrodes (six stacked on each side) with no catalysts; it is similar to a cell described elsewhere (see figure 2 in ref. 7). We report the potential–current response (Fig. 1b) and the potential–power relationship (Fig. 1c and d) for various states of charge (SOCs; measured with respect to the quinone side of the cell). As the SOC increased from 10% to 90%, the open-circuit potential increased linearly from 0.69 V to 0.92 V. In the galvanic direction, peak power densities were 0.246 W cm^{-2} and 0.600 W cm^{-2} at these same SOCs, respectively (Fig. 1c). To avoid significant water splitting in the electrolytic direction, we used a cut-off voltage of 1.5 V, at which point the current densities observed at 10% and 90% SOCs were -2.25 A cm^{-2} and -0.95 A cm^{-2} , respectively, with corresponding power densities of -3.342 W cm^{-2} and -1.414 W cm^{-2} .

In Fig. 2 we report the results of initial cycling studies for this battery, to test for consistent performance over longer timescales. Figure 2a shows cycling data at $\pm 0.2 \text{ A cm}^{-2}$ using 50% of the total capacity of

the battery. The cycles are highly reproducible and indicate that current efficiencies for the battery are around 95%. Figure 2b shows constant-current cycling data, collected at $\pm 0.5 \text{ A cm}^{-2}$, using voltage cut-offs of 0 V and 1.5 V. These tests were done using the identical solutions used in the battery for Fig. 1b–d. The galvanic discharge capacity retention (that is, the number of coulombs extracted in one cycle divided by the number of coulombs extracted in the previous cycle) is above 99%, indicating the battery is capable of operating with minimal capacity fade and suggesting that current efficiencies are actually closer to 99%. Full characterization of the current efficiency will require slower cycling experiments and chemical characterization of the electrolyte solutions after extended cycling.

To gain a better understanding of the quinone half-reaction on carbon, AQDS was subjected to half-cell electrochemical measurements. Cyclic voltammetry of a 1 mM solution of AQDS in 1 M sulphuric acid on a glassy carbon disk working electrode shows current peaks corresponding to reduction and oxidation of the anthraquinone species^{9–11} (Fig. 3d, solid trace). The peak separation of 34 mV is close to the value of $59 \text{ mV}/n$, where n is the number of electrons involved, expected for a two-electron process. Rotation of this disk at a variety of rates yields mass-transport-limited currents (Fig. 3a) from which the AQDS diffusion coefficient ($D = 3.8(1) \times 10^{-6} \text{ cm}^2 \text{ s}^{-1}$) can be determined; throughout this paper, the numbers reported in parentheses indicate the standard deviation in the last reported digit. Koutecký–Levich analysis at low overpotentials (Fig. 3b) can be extrapolated to infinite rotation rate and fitted to the Butler–Volmer equation (Extended Data Fig. 3a) to give the kinetic reduction rate constant $k_0 = 7.2(5) \times 10^{-3} \text{ cm s}^{-1}$. This rate constant is greater than that found for other species used in flow batteries such as $\text{V}^{3+}/\text{V}^{2+}$, Br_2/Br^- and $\text{S}_4^{2-}/\text{S}_2^{2-}$ (see table 2 in ref. 3). It implies that the voltage loss due to the rate of surface electrochemical reactions is negligible. The high rate is apparently due to an outer-sphere two-electron reduction into the aromatic π system requiring little reorganizational energy. The electrochemical reversibility of the two-electron redox reaction was confirmed by fitting the slope to the Butler–Volmer equation (Extended Data Fig. 3a), giving the transfer coefficient $\alpha = 0.474(2)$, which is close to the value of 0.5 expected for an ideally reversible reaction. The Pourbaix diagram (Extended Data Fig. 4) confirms that a two-electron, two-proton reduction occurs in acidic solution, and yields approximate $\text{p}K_a$ values of 7 and 11 for the reduced AQDS species¹¹.

Functionalization of the anthraquinone backbone with electron-donating groups such as hydroxy can be expected lower the reduction potential of AQDS (E^0), thereby raising the cell voltage¹². Hydroxy-substituted anthraquinones are synthesized through oxidation reactions that may be performed at minimal cost. They are also natural products that have been extracted for millennia from common sources such as rhubarb and could even provide a renewable source for future anthraquinone-based electrolyte solutions.

Quantum chemical calculations of un-substituted and hydroxy-substituted AQDS were performed to predict how substitution patterns would change both E^0 of the quinone/hydroquinone couples (Fig. 3c)

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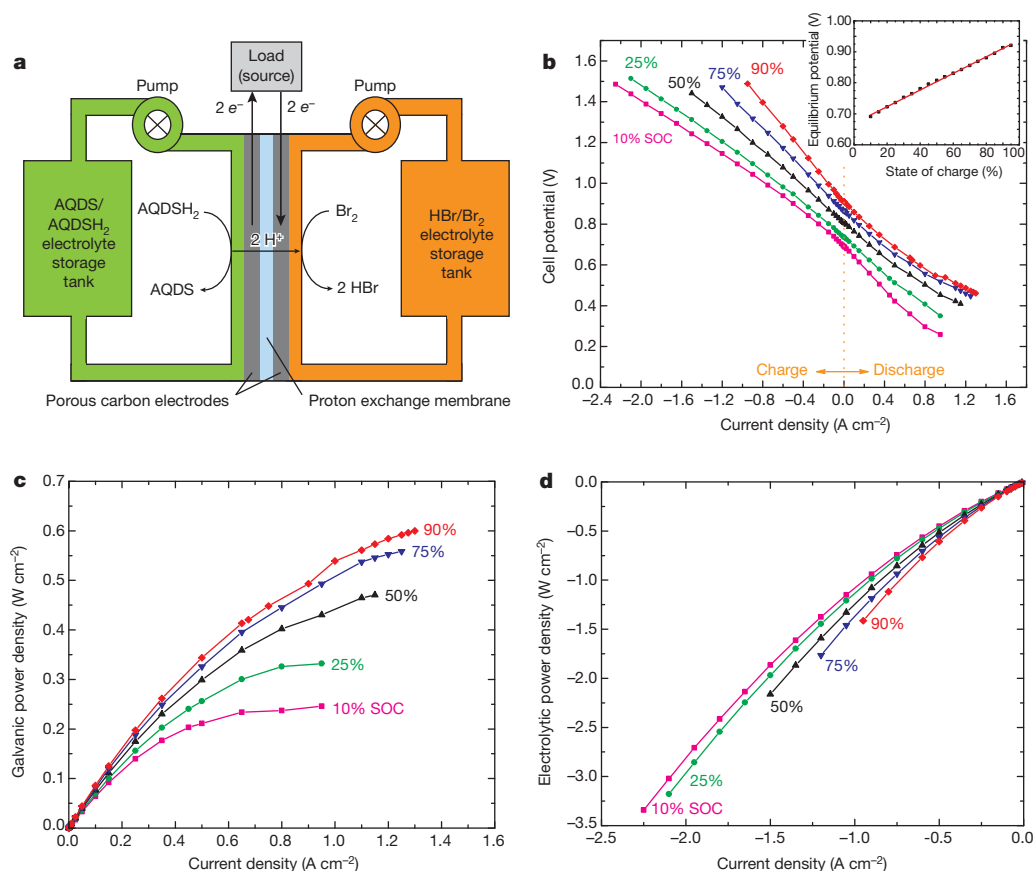


Figure 1 | Cell schematic and cell performance in galvanic and electrolytic modes. **a**, Cell schematic. Discharge mode is shown; the arrows are reversed for electrolytic/charge mode. AQDSH₂ refers to the reduced form of AQDS. **b**, Cell potential versus current density at five different states of charge (SOCs; average of three runs); inset shows the cell open circuit potential

versus SOC with best-fit line superimposed ($E_{eq} = (0.00268 \times \text{SOC}) + 0.670$; $R^2 = 0.998$). **c**, Galvanic power density versus current density for the same SOC levels. **d**, Electrolytic power density versus current density. All data here were collected at 40 °C using a 3 M HBr + 0.5 M Br₂ solution on the positive side and a 1 M AQDS + 1 M H₂SO₄ solution on the negative side.

and the solvation free energy (G_{solv}^0) in aqueous solution (Extended Data Table 1). The addition of $-\text{OH}$ groups is calculated to lower the E^0 by an average of -50 mV per $-\text{OH}$ and provide a wide window for tuning E^0 by almost 0.6 V. In addition, increasing numbers of hydroxy substituents are expected to raise the aqueous solubility due to hydrogen bonding.

In confirmation of the theory, the experimental reduction potential of 1,8-dihydroxy-9,10-anthraquinone-2,7-disulphonic acid (DHAQDS) was found to be 118 mV (versus the standard hydrogen electrode), which is very close to the 101 mV calculated for this species (Fig. 3c and d). The experimental E^0 of DHAQDS was 95 mV lower than AQDS, and would result in an 11% increase in QBFB cell potential.

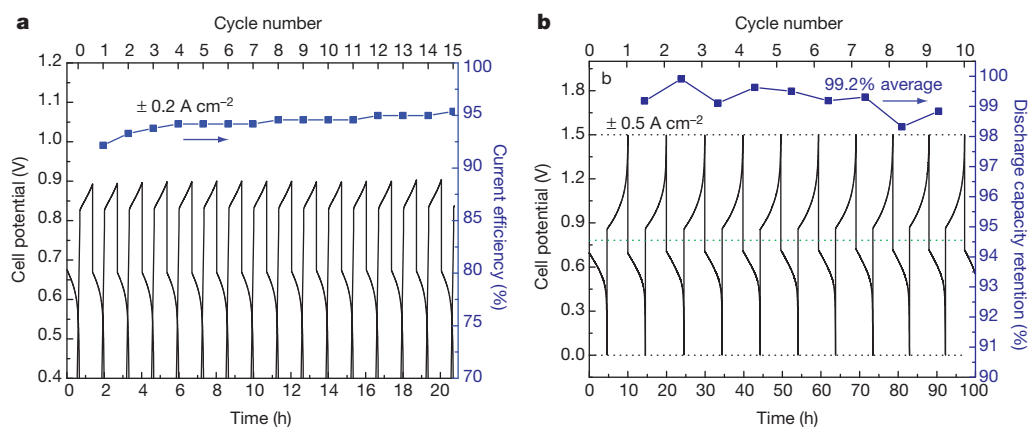


Figure 2 | Cell cycling behaviour. **a**, Constant-current cycling at 0.2 A cm^{-2} at 40 °C using a 2 M HBr + 0.5 M Br₂ solution on the positive side and a 0.1 M AQDS + 2 M H₂SO₄ solution on the negative side; current efficiency is indicated for each complete cycle. **b**, Constant-current cycling at 0.5 A cm^{-2}

at 40 °C using a 3 M HBr + 0.5 M Br₂ solution on the positive side and a 1 M AQDS + 1 M H₂SO₄ solution on the negative side (same solution used in Fig. 1); discharge capacity retention is indicated for each cycle.

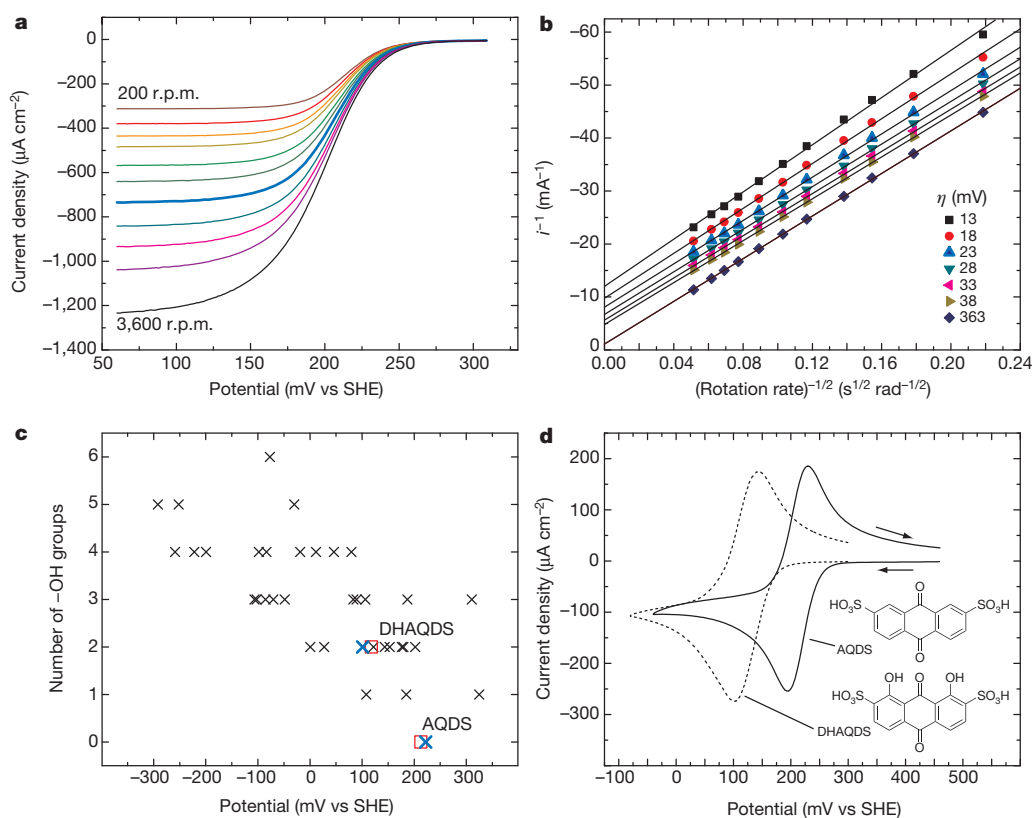


Figure 3 | Half-cell measurements and theory calculations. **a**, Rotating disk electrode (RDE) measurements of AQDS using a glassy carbon electrode in 1 M H_2SO_4 at 11 rotation rates ranging from 200 r.p.m. (red) to 3,600 r.p.m. (black). **b**, Koutecký–Levich plot (current $^{-1}$ versus rotation rate $^{-1/2}$) derived from **a** at seven different AQDS reduction overpotentials, η . **c**, Calculated

reduction potentials of AQDS substituted with -OH groups (black), calculated AQDS and DHAQDS values (blue), and experimental values for AQDS and DHAQDS (red squares). **d**, Cyclic voltammogram of AQDS and DHAQDS (1 mM) in 1 M H_2SO_4 on a glassy carbon electrode (scan rate = 25 mV s^{-1}).

DHAQDS was also found to have faster reduction kinetics ($k_0 = 1.56(5) \times 10^{-2} \text{ cm s}^{-1}$), possibly due to intramolecular hydrogen bonding of the -OH to the ketone (Extended Data Fig. 3b).

The organic approach liberates battery redox chemistry from the constraints of the limited number of elemental redox couples of the periodic table. Although quinones have been used previously in batteries using redox-active solids^{13–15}, their incorporation into all-liquid flow batteries offers the following advantages over current flow-battery technologies. First, scalability: AQDS contains only the Earth-abundant atoms carbon, sulphur, hydrogen and oxygen, and can be inexpensively manufactured on large scales. Because some hydroxy-anthraquinones are natural products, there is also the possibility that the electrolyte material can be renewably sourced. Second, kinetics: quinones undergo extremely rapid two-electron redox on simple, inexpensive carbon electrodes and do not require a costly precious-metal catalyst. Furthermore, this electrode permits higher charging voltages by suppressing the parasitic water-splitting reactions. Third, stability: quinones should exhibit minimal membrane crossover owing to their relatively large size and charge in aqueous solution as a sulphonate anion. Furthermore, although bromine crossover is a known issue in zinc-bromine, vanadium-bromine and hydrogen-bromine cells, AQDS is stable to prolonged heating in concentrated Br_2/HBr mixtures (Extended Data Figs 5 and 6), and the QBFB can be cycled in HBr electrolyte solutions (Extended Data Fig. 9). Fourth, solubility: AQDS has an aqueous solubility greater than 1 M at pH 0, and the quinone solution can thus be stored at relatively high energy density—volumetric and gravimetric energy densities exceed 50 Wh l^{-1} and 50 Wh kg^{-1} , respectively. Last, tunability: the reduction potential and solubility of AQDS can be further optimized by introduction of functional groups such as -OH. Use of DHAQDS is expected to lead to an increase in cell potential, performance and energy density.

These features lower the capital cost of storage chemicals per kilowatt hour, which sets a floor on the ultimate system cost per kilowatt hour at any scale. The precursor molecule anthracene is abundant in crude petroleum and is already oxidized on large scale to anthraquinone. Sulphonated anthraquinones are used on an industrial scale in wood pulp processing for paper¹⁶, and they can be readily synthesized from the commodity chemicals anthraquinone and oleum⁸. In fact, a cyclic voltammogram of the crude sulphonation product of these two reagents is virtually identical to that of pure AQDS (Extended Data Fig. 8). Based on this simple electrolyte preparation that requires no further product separation, we estimate chemical costs of \$21 per kilowatt hour for AQDS and \$6 per kilowatt hour for bromine¹⁷ (see Methods for information on cost calculations). The QBFB offers major cost improvements over vanadium flow batteries with redox-active materials that cost \$81 per kilowatt hour (ref. 18). Optimization of engineering and operating parameters such as the flow field geometry, electrode design, membrane separator and temperature—which have not yet even begun—should lead to significant performance improvements in the future, as it has for vanadium flow batteries, which took many years to reach the power densities we report here⁶. The use of redox processes in π -aromatic organic molecules represents a new and promising direction for cost-effective, large-scale energy storage.

METHODS SUMMARY

The QBFB comprised a mixture of commercially available and custom-made components. Pretreated 2 cm^2 , stacked ($\times 6$) Toray carbon paper electrodes (each of which is about $7.5 \mu\text{m}$ uncompressed) were used on both sides of the cell. Nafion 212 ($50 \mu\text{m}$ thick) was used as a proton-exchange membrane, and PTFE gasketing was used to seal the cell assembly. On the positive side of the cell, 120 ml of 3 M HBr and 0.5 M Br_2 were used as the electrolyte solution in the fully discharged state;

on the negative side, 1 M 2,7-AQDS in 1 M H₂SO₄ was used. AQDS disodium salt was flushed twice through a column containing Amberlyst 15H ion-exchange resin to remove the sodium ions. Half-cell measurements were conducted using a Ag/AgCl aqueous reference electrode (3 M KCl filling solution), a Pt wire counter electrode and a 3-mm-diameter glassy carbon disk electrode. For theoretical calculations, the total free energies of molecules were obtained from first-principles quantum chemical calculations within density functional theory at the level of the generalized gradient approximation (GGA) using the PBE functional. Three-dimensional conformer structures for each quinone/hydroquinone molecule were generated using the ChemAxon suite with up to 25 generated conformers per molecule using the Dreiding force field. Generated conformers were used as input structures for the density functional theory geometry optimization employed for determining the formation energy, which in turn is used to evaluate the reduction potential. In the QBBF cost calculation, a price of \$4.74 per kilogram (eBioChem) was used for anthraquinone. To get the sulphonated form actually used here, anthraquinone must be reacted with oleum (H₂SO₄/SO₃), which adds a negligible cost at scale; this cost is not included here. The price of bromine was \$1.76 per kilogram, based on estimates from the US Geological Survey¹⁷. The cell voltage used to calculate costs here was 0.858 V.

Online Content Any additional Methods, Extended Data display items and Source Data are available in the online version of the paper; references unique to these sections appear only in the online paper.

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Author Contributions B.H. and M.P.M. contributed equally to this work. B.H. and M.P.M. designed and tested the battery, with direction from M.J.A. Both M.P.M. and M.R.G. conducted electrochemistry experiments, with direction from M.J.A. M.P.M. and C.J.G. synthesized chemicals with direction from R.G.G. Theoretical calculations were done by C.S. and S.E., with input from M.P.M. and R.G.G. and direction from A.A.-G. X.C. contributed NMR results. B.H., M.P.M., C.S., M.R.G., S.E., A.A.G., R.G.G. and M.J.A. all contributed to writing the manuscript.

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METHODS

Full cell measurements. The QBFB comprised a mixture of commercially available and custom-made components. Circular endplates were machined out of solid aluminium. Current collectors were 3 inch \times 3 inch pyrolytic graphite blocks with interdigitated flow channels (channel width = 0.0625 inch, channel depth = 0.08 inch, landing between channels = 0.031 inch, Fuel Cell Technologies). Pretreated 2 cm², stacked ($\times 6$) Toray carbon paper electrodes (each of which is about 7.5 μ m uncompressed) were used on both sides of the cell. Pretreatment consisted of a 10 min sonication in isopropyl alcohol followed by a five hour soak in a hot (50 °C) mixture of undiluted sulphuric and nitric acids in a 3:1 volumetric ratio. Nafion 212 (50 μ m thick) was used as a proton-exchange membrane (PEM, Alfa Aesar), and PTFE gasketing was used to seal the cell assembly. Membrane pretreatment was done according to previously published protocols⁷. Six bolts (3/8", 16 threads per inch) torqued to 10.2 N m completed the cell assembly, and PTFE tubing was used to transport reactants and products into and out of the cell. The cell was kept on a hot plate and wrapped in a proportional integral derivative (PID)-controlled heating element for temperature control. On the positive side of the cell, 120 ml of 3 M HBr and 0.5 M Br₂ were used as the electrolyte solution in the fully discharged state; on the negative side, 1 M AQDS in 1 M H₂SO₄ was used. HBr was used on the negative side instead of H₂SO₄ for stability testing results displayed in Extended Data Fig. 9. State-of-charge calculations are based on the composition of the quinone side of the cell. 2,7-Anthraquinone disulphonate disodium salt 98% (TCI) was flushed twice through a column containing Amberlyst 15H ion-exchange resin to remove the sodium ions. Measurements shown here were done at 40 °C. March centrifugal pumps were used to circulate the fluids at a rate of approximately 200 ml min⁻¹. For characterization, several instruments were used: a CH Instruments 1100C potentiostat (which can be used up to ± 2 A), a DC electronic load (Circuit Specialists) for galvanic discharge, a DC regulated power supply (Circuit Specialists) for electrolytic characterization, and a standard multimeter for independent voltage measurements. The cell was charged at 1.5 V until a fixed amount of charge ran through the cell. During this process, the electrolyte colours changed from orange to dark green (AQDS to AQDSH₂) and from colourless to red (Br⁻ to Br₂). Periodically, the open circuit potential was measured, providing the data inset in Fig. 1b. Also, at various SOC, potential–current behaviour was characterized using the equipment described above: a fixed current was drawn from the cell, and the voltage, once stabilized, was recorded (Fig. 1b). For the cycling data in Fig. 2b, the potentiostat was used for constant-current (± 0.5 A cm⁻²) measurements with cut-off voltages of 0 V and 1.5 V. For the cycling data in Fig. 2a, a more dilute quinone solution (0.1 M as opposed to 1 M) was used. Here, the half-cycle lengths were programmed to run at constant current for a fixed amount of time, provided the voltage cut-offs were not reached, so that half of the capacity of the battery was used in each cycle. The voltage cut-offs were never reached during charging, but were reached during discharge. Current efficiencies are evaluated by dividing the discharge time by the charge time of the previous half-cycle.

As shown in Fig. 2, current efficiency starts at about 92% and climbs to about 95% over ~ 15 standard cycles. Note that these measurements are done near viable operating current densities for a battery of this kind. Because of this, we believe this number places an upper bound on the irreversible losses in the cell. In any case, 95% is comparable to values seen for other battery systems. For example, ref. 19 reports vanadium bromide batteries with current efficiencies of 50–90%, with large changes in current efficiency observed for varying membrane conditions. Our system will probably be less dependent on membrane conditions because we are storing energy in anions and neutral species as opposed to cations, which Nafion can conduct reasonably well.

In Fig. 2b we illustrate the capacity retention of the battery (that is, the number of coulombs available for discharge at the n th cycle divided by that available for discharge at the $(n - 1)$ th cycle) to be 99.2% on average, which is quite high and provides direct evidence that our irreversible losses are below 1%. If we attribute all of this loss (the 0.78% capacity fade per cycle) to some loss of redox-active quinone, it would be equivalent to losing 0.0006634 moles of quinone per cycle. If we attribute all of the loss to bromine crossover (which would react with the hydroquinone and oxidize it back to quinone), this corresponds to a crossover current density of 1.785 mA cm⁻², which is within the range of the widely varying crossover values reported in the literature²⁰. Note that these crossover numbers can be very sensitive to membrane pretreatment conditions. It is also important to mention that, to determine very accurate current efficiencies, detailed chemical analyses of the electrolyte are necessary.

Half-cell measurements. These were conducted using a BASi Epsilon EC potentiostat, a BASi Ag/AgCl aqueous reference electrode (RE-5B, 3 M KCl filling solution) and a Pt wire counter electrode. Rotating disk electrode (RDE) measurements were conducted using a BASi RDE (RDE-2) and a 3 mm diameter glassy carbon disk electrode. Electrode potentials were converted to the standard hydrogen electrode (SHE) scale using $E(\text{SHE}) = E(\text{Ag}/\text{AgCl}) + 0.210$ V, where $E(\text{SHE})$ is the

potential versus SHE and $E(\text{Ag}/\text{AgCl})$ is the measured potential versus Ag/AgCl. 2,7-Anthraquinone disulphonate disodium salt 98% was purchased from TCI and used as received. 1,8-Dihydroxy-anthraquinone-2,7-disulphonic acid was made according to the literature procedure²¹. The electrolyte solution was sulphuric acid (ACS, Sigma) in deionized H₂O (18.2 M Ω cm, Millipore). The Pourbaix diagram (plot of E^0 versus pH) shown in Extended Data Fig. 4 was constructed using aqueous 1 mM solutions of AQDS in pH buffers using the following chemicals: sulphuric acid (1 M, pH 0), HSO₄⁻/SO₄²⁻ (0.1 M, pH 1–2), AcOH/AcO⁻ (0.1 M, pH 2.65–5), H₂PO₄⁻/HPO₄²⁻ (0.1 M, pH 5.3–8), HPO₄²⁻/PO₄³⁻ (0.1 M, pH 9.28–11.52), and KOH (0.1 M, pH 13). The pH of each solution was adjusted with 1 M H₂SO₄ or 0.1 M KOH solutions and measured with an Oakton pH 11 Series pH meter (Eutech Instruments).

RDE studies. All RDE data represent an average of three runs. Error bars in Extended Data Figs 2 and 3 indicate standard deviations. Before each run, the glassy carbon disk working electrode was polished to a mirror shine with 0.05 μ m alumina and rinsed with deionized water until a cyclic voltammogram of a solution of 1 mM AQDS in 1 M H₂SO₄ showed anodic and cathodic peak voltage separation of 34 to 35 mV (39 mV for DHAQDS) at a sweep rate of 25 mV s⁻¹. The electrode was then rotated at 200, 300, 400, 500, 700, 900, 1,200, 1,600, 2,000, 2,500 and 3,600 r.p.m. while the voltage was linearly swept from 310 to 60 mV (250 to –100 for DHAQDS) at 10 mV s⁻¹ (Extended Data Fig. 1). The currents measured at 60 mV (–100 for DHAQDS) (that is, the diffusion-limited current density) versus the square root of the rotation rate (ω) is plotted in Extended Data Fig. 2. The data were fitted with a straight line, with the slope defined by the Levich equation as $0.620nFAC_0D^{2/3}\nu^{-1/6}$, where $n = 2$, Faraday's constant $F = 96,485$ C mol⁻¹, electrode area $A = 0.0707$ cm², AQDS concentration $C_0 = 10^{-6}$ mol cm⁻³, kinematic viscosity $\nu = 0.01$ cm² s⁻¹. This gives D values of $3.8(1) \times 10^{-6}$ cm² s⁻¹ for AQDS and $3.19(7) \times 10^{-6}$ cm² s⁻¹ for DHAQDS. The reciprocal of the current at overpotentials of 13, 18, 23, 28, 33, 38 and 363 mV was plotted versus the reciprocal of the square root of the rotation rate (Fig. 3b and Extended Data Fig. 2). The data for each potential were fitted with a straight line; the intercept gives the reciprocal of i_K , the current in the absence of mass transport limitations (the extrapolation to infinite rotation rate). A plot of $\log_{10}(i_K)$ versus overpotential was linearly fitted with a slope of 62 mV (AQDS) and 68 mV (DHAQDS) defined by the Butler–Volmer equation as $2.3\alpha RT/nF$ (Extended Data Fig. 3), where R is the universal gas constant, T is temperature in kelvin and α is the charge transfer coefficient. This gives $\alpha = 0.474(2)$ for AQDS and $0.43(1)$ for DHAQDS. The x -intercept gives the log of the exchange current i_0 , which is equal to FAC_0k_0 , and gives $k_0 = 7.2(5) \times 10^{-3}$ cm s⁻¹ for AQDS and $1.56(5) \times 10^{-2}$ cm s⁻¹ for DHAQDS.

Stability studies. AQDS (50 mg) was dissolved in 0.4 ml of D₂O, and treated with 100 μ l of Br₂. The ¹H and ¹³C NMR spectra (Extended Data Figs 5a, b and 6a, b) were unchanged from the starting material after standing for 20 h at 25 °C. AQDS (50 mg) was then treated with 1 ml of 2 M HBr and 100 μ l of Br₂. The reaction was heated to 100 °C for 48 h and evaporated to dryness at that temperature. The resulting solid was fully dissolved in D₂O giving unchanged ¹H and ¹³C NMR (Extended Data Figs 5c and 6c); however, the ¹H NMR reference was shifted due to residual acid. These results imply that bromine crossover will not lead to irreversible destruction of AQDS.

Sulphonation of anthraquinone and electrochemical study. 9,10-Anthraquinone was treated with H₂SO₄ (20% SO₃) at 170 °C for 2 h according to a literature procedure⁸. The resulting red solution, containing roughly 37% AQDS, 60% 9,10-anthraquinone-2,6-disulphonic acid and 3% 9,10-anthraquinone-2-sulphonic acid, was diluted and filtered. A portion of this solution was further diluted with 1 M H₂SO₄ to ~ 1 mM total anthraquinone concentration. The cyclic voltammogram (Extended Data Fig. 8) is similar to that of pure AQDS, though the anodic/cathodic peak current density is broadened due to the presence of the multiple sulphonate isomers.

Theory and methods. We used a fast and robust theoretical approach to determine the E^0 of quinone/hydroquinone couples in aqueous solutions. We employed an empirical linear correlation of ΔH_f , the heat of formation of hydroquinone at 0 K from the quinone and hydrogen gas, to the measured E^0 values²². Following the treatment of ref. 22, the linear correlation is described as $\Delta G = -nFE^0$, where ΔG is the difference in total free energy between quinone and hydroquinone, n is the number of electrons involved in the reaction and F is the Faraday constant. The entropy contributions to the total free energies of reaction have been neglected because the entropies of reduction of quinones are found to be very similar^{22,23}, and the E^0 of the oxidation–reduction system is linearly expressed as $(-nF)^{-1}\Delta H_f + b$, where b is a constant. It was also assumed that the reduction of quinones takes place in a single-step reaction involving a two-electron two-proton process^{9,24}. The total free energies of molecules were obtained from first-principles quantum chemical calculations within density functional theory (DFT) at the level of the generalized gradient approximation (GGA) using the PBE functional²⁵. The projector augmented wave (PAW) technique and a plane-wave basis set^{26,27} as implemented

in VASP^{28,29} were employed. The kinetic energy cut-off for the plane-wave basis was set at 500 eV, which was sufficient to converge the total energies on a scale of 1 meV per atom. To obtain the ground-state structures of molecules in the gas phase, we considered multiple initial configurations for each molecule and optimized them in a cubic box of 25 Å using Γ -point sampling. The geometries were optimized without any symmetry constraints using the conjugate gradient (CG) algorithm, and the convergence was assumed to be complete when the total remaining forces on the atoms were less than 0.01 eV Å⁻¹.

The search for conformational preference through theoretical calculations for each hydroxylated quinone is crucial because of the significant effects of intramolecular hydrogen bonds on the total free energies of the molecules³⁰. Three-dimensional conformer structures for each quinone/hydroquinone molecule were generated using the ChemAxon suite (Marvin 6.1.0 by ChemAxon, <http://www.chemaxon.com>) with up to 25 conformers generated per molecule using the Dreiding force field³¹. The conformers generated were used as input structures for the DFT geometry optimization employed for determining ΔH_f , which in turn is used to estimate E^0 and G_{solv}^0 .

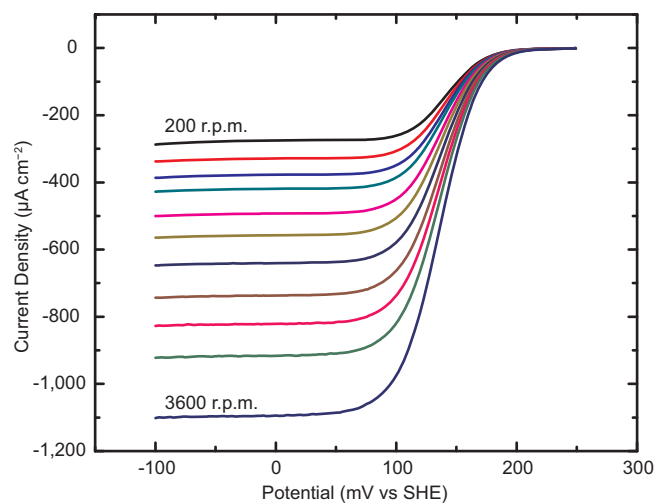
To calculate the E^0 of the hydroxy-substituted AQDS molecules (Fig. 3c), the correlation between ΔH_f and E^0 was calibrated from experimental data on six well-characterized quinones³². Specifically, we used the experimental values of the aqueous E^0 and computed ΔH_f of 1,2-benzoquinone, 1,4-benzoquinone, 1,2-naphthoquinone, 1,4-naphthoquinone, 9,10-anthraquinone and 9,10-phenanthrene³³. The training set ensures that the calibration plot addresses most classes and aspects of quinones, including two quinones each from 1-ring (benzoquinone), 2-ring (naphthoquinone) and 3-ring (anthraquinone and phenanthrene) structures. In addition, the experimental values of E^0 of the training set spanned 0.09 V (9,10-anthraquinone) to 0.83 V (1,2-benzoquinone), providing a wide range for E^0 (Extended Data Fig. 7). The linear calibration plot for E^0 yields an $R^2 = 0.97$ between the calculated ΔH_f and E^0 (Extended Data Fig. 7).

The G_{solv}^0 values of the quinones in water were calculated using the Jaguar 8.0 program in the Schrödinger suite 2012 (Jaguar, version 8.0, Schrödinger). The standard Poisson–Boltzmann solver was employed^{34,35}. In this model, a layer of charges on the molecular surface represents the solvent. G_{solv}^0 was calculated as the difference between the total energy of the solvated structure and the total energy of the molecule in vacuum. A more negative value for G_{solv}^0 corresponds to a quinone that is likely to have a higher aqueous solubility. An absolute prediction of the solubility is not readily available, as the accurate prediction of the most stable forms of molecular crystal structures with DFT remains an open problem³⁶.

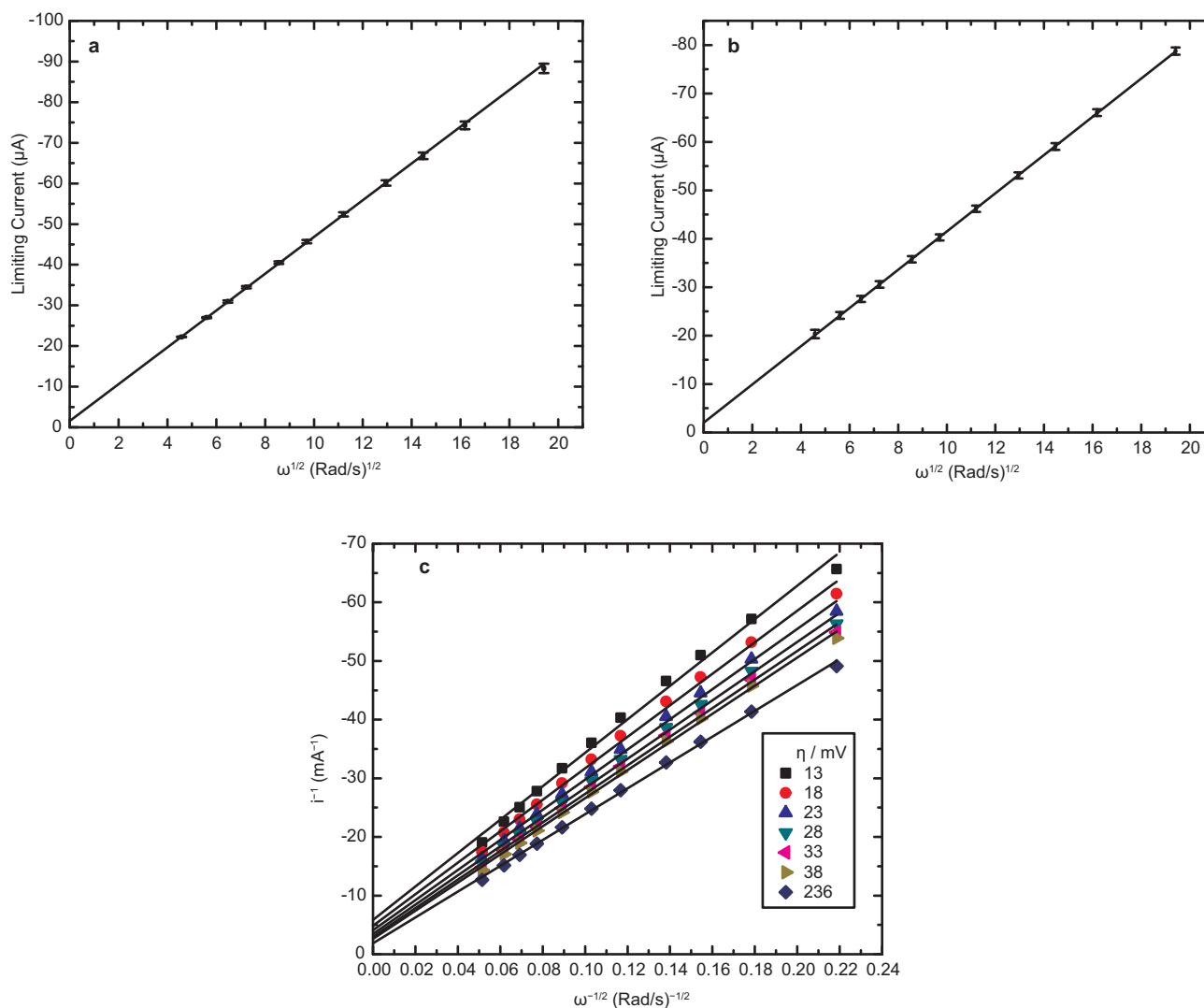
Cost calculations. These were done using the following formula: $C = (3.6 \times 10^3) \times (PM)/(nFE)$, where C is the cost in US dollars of the compound per kilowatt hour, P is the cost in US dollars per kilogram, M is the molecular mass of the compound, F is Faraday's constant (96,485 C mol⁻¹), n is the number of moles of electrons transferred per mole of storage compound (two for the QBFB) and E is the open-circuit voltage (V) of the storage device. In calculating the price for the anthraquinone-bromine battery, a price of \$4.74 per kilogram (eBioChem) was used for anthraquinone

(note that, to get the sulphonated form actually used here, anthraquinone must be reacted with oleum (H₂SO₄/SO₃), which adds a negligible cost at scale; this cost is not included here). The price of bromine was \$1.76 per kilogram, based on estimates from the US Geological Survey¹⁷. The cell voltage used to calculate costs here was 0.858 V. For vanadium, costs were calculated from USGS prices from 2011¹⁸ of vanadium pentoxide at \$14.37 per kilogram, and the cell voltage used was 1.2 V. Balance-of-system costs have not been estimated because the technology is too immature.

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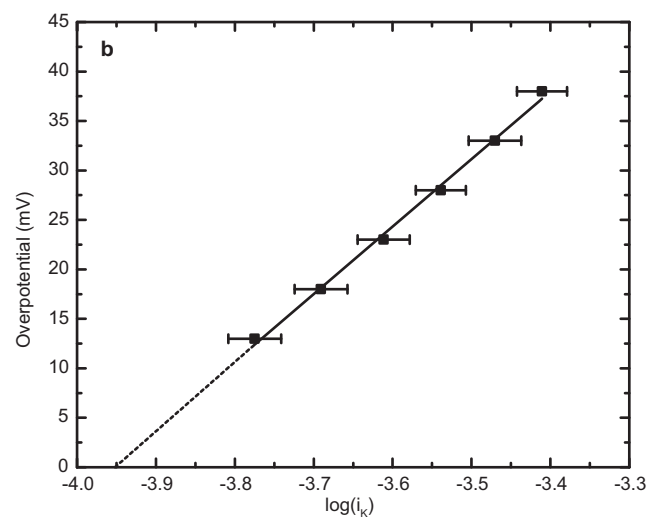
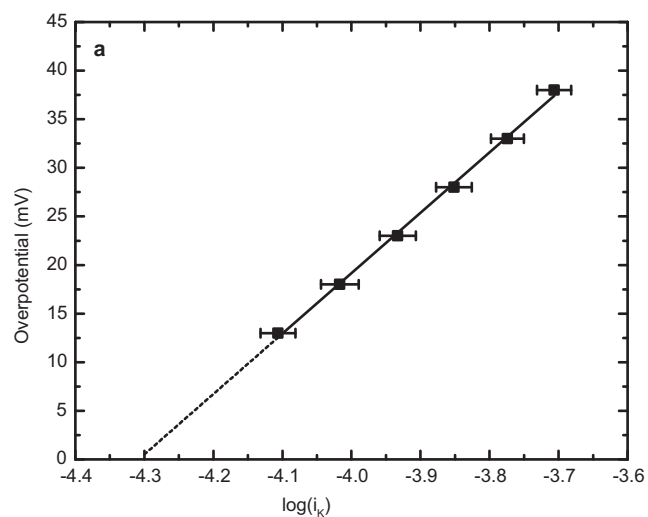


Extended Data Figure 1 | Plot of potential versus current density at different rotation rates of the RDE. The solution is 1 mM DHAQDS (1 mM in 1 M H_2SO_4), using a rotating disk electrode (RDE) of glassy carbon. Rotation rates are 200, 300, 400, 500, 700, 900, 1,200, 1,600, 2,000, 2,500 and 3,600 r.p.m.



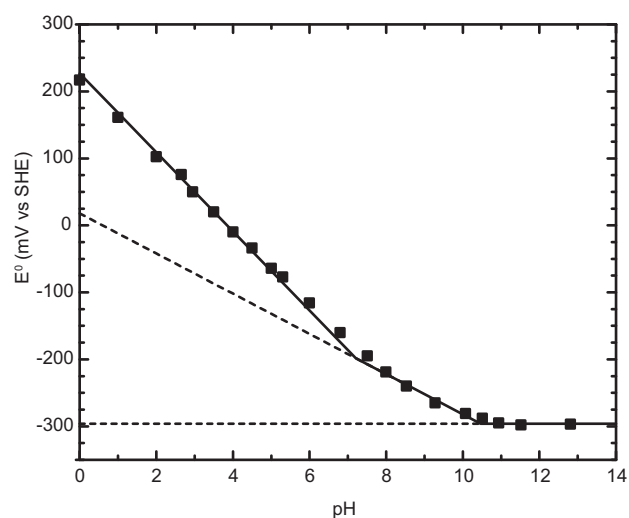
Extended Data Figure 2 | Levich and Koutecký–Levich plots obtained using the RDE. **a**, Levich plot (limiting current versus square root of rotation rate ω) of 1 mM AQDS in 1 M H₂SO₄ (the fitted line has a slope of $4.53(2) \mu\text{A s}^{1/2} \text{rad}^{-1/2}$, giving $D = 3.8(1) \times 10^{-6} \text{cm}^2 \text{s}^{-1}$). Data are an average of three runs; error bars indicate the standard deviation. **b**, As **a** but for

DHAQDS in 1 M H₂SO₄ (slope of $3.94(6) \mu\text{A s}^{1/2} \text{rad}^{-1/2}$ gives $D = 3.19(7) \times 10^{-6} \text{cm}^2 \text{s}^{-1}$). **c**, Koutecký–Levich plot (i^{-1} versus $\omega^{-1/2}$) of 1 mM DHAQDS in 1 M H₂SO₄. The current response, i , is shown for seven different AQDS reduction overpotentials η .

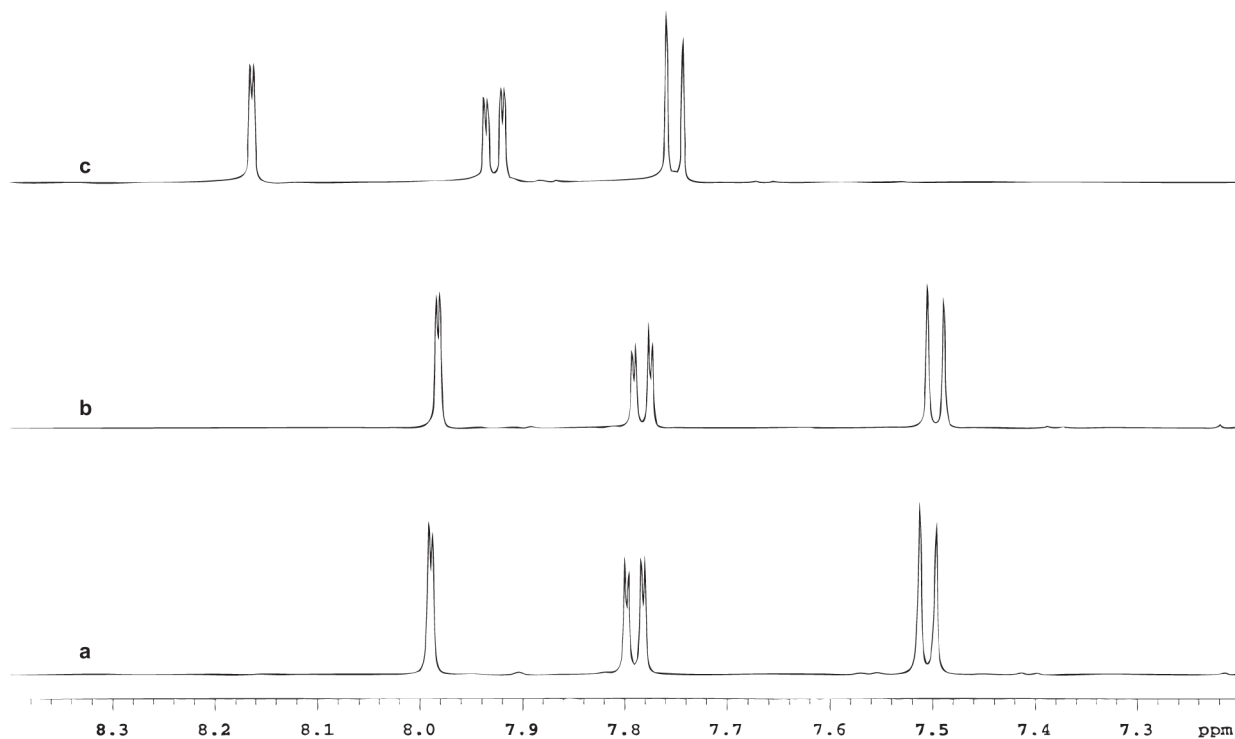


Extended Data Figure 3 | Fit of Butler–Volmer equation. Constructed using the current response in the absence of mass transport at low AQDS reduction overpotentials; i_k is the current extrapolated from the zero-intercept of Fig. 3b and Extended Data Fig. 2c (infinite rotation rate). Data are an average

of three runs; error bars indicate the standard deviation. **a**, AQDS: best-fit line has the equation $y = 62(x + 4.32)$. This yields $\alpha = 0.474(2)$ and $k_0 = 7.2(5) \times 10^{-3} \text{ cm s}^{-1}$. **b**, DHAQDS: best-fit line is the function $y = 68(x + 3.95)$. This yields $\alpha = 0.43(1)$ and $k_0 = 1.56(5) \times 10^{-2} \text{ cm s}^{-1}$.

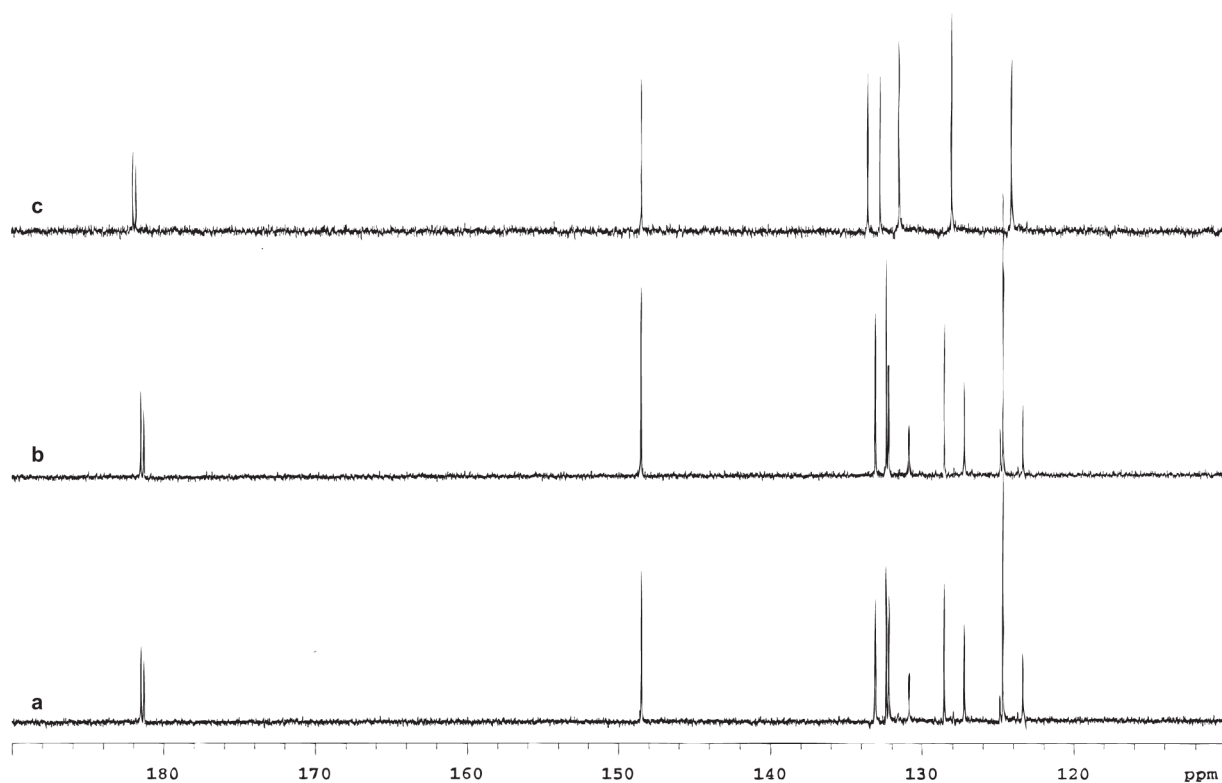


Extended Data Figure 4 | Pourbaix diagram (E^0 vs pH) of AQDS. Data are fitted to three solid lines indicating slopes of -59 mV pH^{-1} , -30 mV pH^{-1} and 0 mV pH^{-1} , corresponding to two-, one- and zero-proton processes, respectively. Dashed lines linearly extrapolate the one- and zero-proton processes to give E^0 values of 18 mV ($2e^-/1H^+$) and -296 mV ($2e^-/0H^+$).



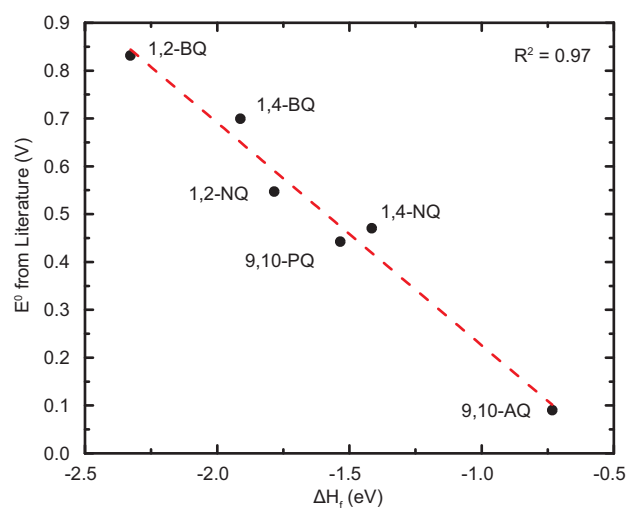
Extended Data Figure 5 | ^1H NMR (500 MHz, D_2O) spectra. **a**, Spectrum of AQDS: chemical shift $\delta = 7.99$ p.p.m. versus tetramethylsilane (TMS) (doublet (d), coupling constant $J = 2$ Hz, 1,8 C–H), 7.79 p.p.m. (doublet of doublets, $J = 2$ and 8 Hz, 4,5 C–H), 7.50 p.p.m. (d, $J = 8$ Hz, 3,6 C–H). **b**, The same

sample, 20 h after addition of Br_2 . **c**, ^1H NMR of AQDS treated with 2 M HBr and Br_2 and heated to 100°C for 48 h. The peaks are shifted due to presence of trace HBr, which shifted the residual solvent peak due to increased acidity. Coupling constants for each peak are identical to **a**.



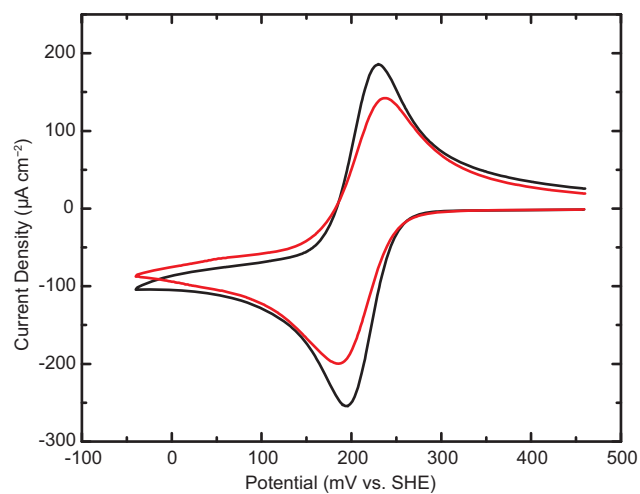
Extended Data Figure 6 | ^{13}C NMR (500 MHz, D_2O) spectra. **a**, AQDS, $\delta = 181.50$ p.p.m. versus TMS (C 9), 181.30 p.p.m. (C 10), 148.51 p.p.m. (C 2,7), 133.16 p.p.m. (C 11), 132.40 p.p.m. (C 12), 130.86 p.p.m. (C 3,6),

128.59 p.p.m. (C 4,5), 124.72 p.p.m. (C 1,8). **b**, The same sample, 24 h after addition of Br_2 . **c**, ^{13}C NMR of AQDS treated with 2 M HBr and Br_2 and heated to 100°C for 48 h.

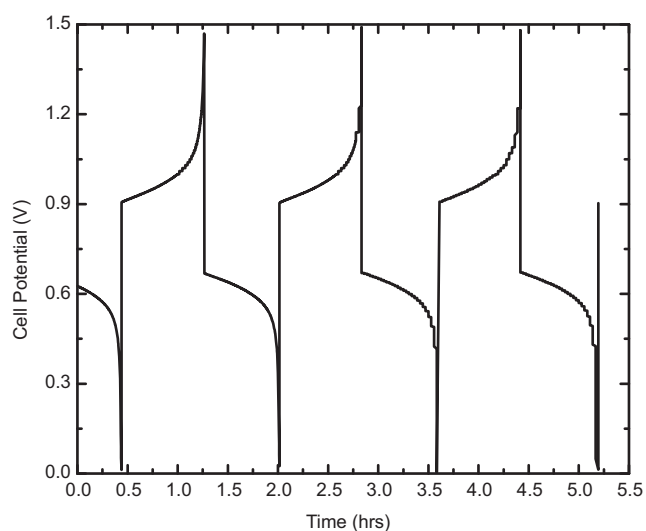


Extended Data Figure 7 | Calibration model for ΔH_f and experimental E^0 .

This shows a linear relationship (red dashed line; $R^2 = 0.97$) between calculated ΔH_f (this work) and experimental E^0 (from the literature) of six quinones in aqueous solutions: BQ, benzoquinone; NQ, naphthoquinone; AQ, anthraquinone; and PQ, phenanthraquinone.

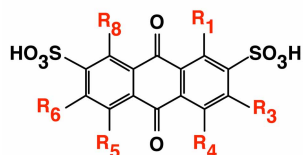


Extended Data Figure 8 | AQDS cyclic voltammograms. Black curve, obtained for a 1 mM solution of AQDS in 1 M H_2SO_4 on a stationary glassy carbon working electrode. Red curve, obtained for a crude anthraquinone sulphonation solution containing a mixture of AQDS, 9,10-anthraquinone-2,6-disulphonic acid and 9,10-anthraquinone-2-sulphonic acid diluted to 1 mM total anthraquinone in 1 M H_2SO_4 .



Extended Data Figure 9 | Flow-battery cycling behaviour with HBr electrolyte on both sides. Data collected by cycling the current at 0.2 A cm^{-2} at 40°C using a $2 \text{ M HBr} + 0.5 \text{ M Br}_2$ solution on the positive side and a $2 \text{ M HBr} + 0.1 \text{ M AQDS}$ solution on the negative side; cell potential versus time performance is comparable to data in Fig. 2.

Extended Data Table 1 | AQDS screened by theoretical calculations



ID	-OH substituted	R ₁	R ₃	R ₄	R ₅	R ₆	R ₈	E ⁰ (V)	G ⁰ _{solv} (kJ mol ⁻¹)
1	Non-	H	H	H	H	H	H	0.222	-81.5
2	Mono-	OH	H	H	H	H	H	0.185	-81.5
3		H	OH	H	H	H	H	0.325	-111.7
4		H	H	OH	H	H	H	0.108	-88.2
5		OH	OH	H	H	H	H	0.176	-110.3
6	Di-	OH	H	OH	H	H	H	0.027	-85.6
7		OH	H	H	OH	H	H	0.122	-96.7
8		OH	H	H	H	OH	H	0.143	-85.7
9		OH	H	H	H	H	OH	0.101	-83.2
10		H	OH	OH	H	H	H	0.153	-105.4
11		H	OH	H	OH	H	H	0.179	-119.1
12		H	OH	H	H	OH	H	0.202	-112.0
13		H	H	OH	OH	H	H	0.000	-95.6
14	Tri-	OH	OH	OH	H	H	H	-0.070	-101.7
15		OH	OH	H	OH	H	H	0.083	-116.2
16		OH	OH	H	H	OH	H	0.187	-114.3
17		OH	OH	H	H	H	OH	0.310	-120.9
18		OH	H	OH	OH	H	H	-0.102	-91.4
19		OH	H	OH	H	OH	H	0.089	-114.0
20		OH	H	OH	H	H	OH	-0.085	-87.1
21		OH	H	H	OH	OH	H	-0.048	-102.8
22		H	OH	OH	OH	H	H	-0.107	-107.8
23		H	OH	OH	H	OH	H	0.106	-136.8
24	Tetra-	OH	OH	OH	OH	H	H	-0.098	-109.0
25		OH	OH	OH	H	OH	H	0.012	-108.4
26		OH	OH	OH	H	H	OH	-0.222	-102.3
27		OH	OH	H	OH	OH	H	-0.019	-132.3
28		OH	OH	H	OH	H	OH	0.046	-114.6
29		OH	OH	H	H	OH	OH	0.080	-111.1
30		OH	H	OH	OH	OH	H	-0.259	-99.0
31		OH	H	OH	OH	H	OH	-0.199	-91.9
32		H	OH	OH	OH	OH	H	-0.083	-120.6
33	Penta-	OH	OH	OH	OH	OH	H	-0.252	-117.1
34		OH	OH	OH	OH	H	OH	-0.292	-108.3
35		OH	OH	OH	H	OH	OH	-0.030	-111.6
36	Hexa-	OH	OH	OH	OH	OH	OH	-0.077	-121.0

The effect of -OH substitution on reduction potential and solvation energy.