



HAL
open science

Cation desolvation-induced capacitance enhancement in reduced graphene oxide (rGO)

Kangkang Ge, Hui Shao, Encarnacion Raymundo-Piñero, Pierre-Louis Taberna, Patrice Simon

► **To cite this version:**

Kangkang Ge, Hui Shao, Encarnacion Raymundo-Piñero, Pierre-Louis Taberna, Patrice Simon. Cation desolvation-induced capacitance enhancement in reduced graphene oxide (rGO). *Nature Communications*, 2024, 15 (1), pp.1935. 10.1038/s41467-024-46280-1 . hal-04778845

HAL Id: hal-04778845

<https://hal.science/hal-04778845v1>

Submitted on 20 Nov 2024

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Distributed under a Creative Commons Attribution 4.0 International License

Cation desolvation-induced capacitance enhancement in reduced graphene oxide (rGO)

Received: 17 November 2023

Accepted: 15 February 2024

Published online: 02 March 2024

 Check for updates

Kangkang Ge¹, Hui Shao², Encarnacion Raymundo-Piñero^{3,4},
Pierre-Louis Taberna^{1,3}  & Patrice Simon^{1,3} 

Understanding the local electrochemical processes is of key importance for efficient energy storage applications, including electrochemical double layer capacitors. In this work, we studied the charge storage mechanism of a model material - reduced graphene oxide (rGO) - in aqueous electrolyte using the combination of cavity micro-electrode, operando electrochemical quartz crystal microbalance (EQCM) and operando electrochemical dilatometry (ECD) tools. We evidence two regions with different charge storage mechanisms, depending on the cation-carbon interaction. Notably, under high cathodic polarization (region II), we report an important capacitance increase in Zn²⁺ containing electrolyte with minimum volume expansion, which is associated with Zn²⁺ desolvation resulting from strong electrostatic Zn²⁺-rGO interactions. These results highlight the significant role of ion-electrode interaction strength and cation desolvation in modulating the charging mechanisms, offering potential pathways for optimized capacitive energy storage. As a broader perspective, understanding confined electrochemical systems and the coupling between chemical, electrochemical and transport processes in confinement may open tremendous opportunities for energy, catalysis or water treatment applications in the future.

Understanding the charge storage mechanism and, more generally, the processes occurring at the electrode/electrolyte interface, is of key importance for the development of electrochemical energy storage material and devices with improved performance. Unlike batteries, where diffusion limitations in the electrodes are prevalent, charge storage in electrochemical double layer capacitors is governed by a surface-controlled process, thus offering high-rate charge-discharge capability. Historically, the electric double layer (EDL) classical model proposed by Helmholtz and further refined by Gouy–Chapman–Stern (GCS) in the 19th century set the scientific foundations for understanding the charge/discharge mechanisms in capacitive materials¹. Classical EDL-type charge storage mechanism is achieved by ion

adsorption/desorption through electrostatic forces without involving any redox charge transfer. This characteristic ensures fast charging rate, no diffusion limitation and long-term cycling stability². Pseudocapacitive electrodes exhibit similar electrochemical response to EDL-type electrodes like porous carbon and graphene, with a charge changing linearly with the potentials^{3,4}. However, pseudocapacitive charge storage involves charge transfer or partial charge transfer across the electrochemical interface, distinguishing such process from the electrostatic capacitive charge storage observed in EDL-type electrodes⁴, which in turn leads to a lower cycling stability. Both capacitive and pseudocapacitive electrodes exhibit unique confined geometries, such as narrow slit pores for porous carbon or layered

¹Université Paul Sabatier, CIRIMAT UMR CNRS 5085, 118 Route de Narbonne, 31062 Toulouse, France. ²i-Lab, CAS Center for Excellence in Nanoscience, Suzhou Institute of Nano-Tech and Nano-Bionics (SINANO), Chinese Academy of Sciences (CAS), Suzhou 215123, China. ³Réseau sur le Stockage Electrochimique de l'Energie (RS2E), FR CNRS 3459 Amiens, France. ⁴Université Orléans, CNRS, CEMHTI UPR3079 Orléans, France.

 e-mail: pierre-louis.taberna@univ-tlse3.fr; patrice.simon@univ-tlse3.fr

structures for MXenes and graphene-like materials. Confined geometry arises when the traditional planar interfaces approach to each other, leading to an overlap of EDLs⁵. Over the past several decades, significant advancements have been made regarding the characterization of nanoporous carbons for capacitive energy storage^{6,7}. A notable finding was the anomalous increase in capacitance when pore sizes were reduced to below 1 nm⁸. Smaller pore radii permit ions to get closer to the carbon pore walls thanks to partial desolvation, resulting in the creation of image charges in the carbon electrodes, more efficient screening the ionic charges and capacitance increase^{9,10}. As a result, enhanced energy storage capabilities are achieved, suggesting that energy stored at the interface is somehow directly correlated to solvation energy^{11,12}.

Confined geometries such as narrow slit pores are also present in 2D layered materials, including reduced graphene oxide (rGO), MXenes, and others¹³. In particular, water confined between the interlayers of 2D materials is often reported to show improved charge transport kinetics and efficient charge storage in layered metal oxides^{14,15}, including MnO₂^{16,17}, V₂O₅¹⁸, WO₃^{19,20} and so on. Specifically, Augustyn et al. emphasized the impact of confined structural water networks in WO₃·nH₂O, which promotes the structural stability and facilitates a rapid proton intercalation rate¹⁹. Additionally, the promotional effect of structural water has been reported in realizing significant capacity improvements in vanadium oxide nanosheets¹⁸ and MnO₂ birnessite nanolayers, where the intercalation of partial desolvated K⁺ ions resulted in improved capacity by increasing charge transfer to the host¹⁴. These recent studies suggest that capacitive charge storage under confined geometries cannot be simply described as either pure EDL or pseudocapacitance, but instead should be considered more as a continuum based on the solvent-mediated interactions between electrolyte ions and electrode host materials^{13,17}.

For 2D layered materials, the nature of surface terminations has also been identified as a key factor in affecting charge storage capability. For instance, oxygen-containing surface functional groups significantly improve charge storage through efficient surface redox reactions in aqueous electrolytes^{21,22}. Additionally, interlayer confined water is controlled by the presence of -OH functional groups on MXenes, which plays a pivotal role in facilitating the transport of hydrated H⁺ in the interlayer spacing to reach the redox sites²³. Besides, efficient ion transport was observed with nitrogen-terminated MXenes, still in acidic aqueous electrolytes²⁴. However, most of the studies deal with acidic/alkaline electrolytes, basically because of their high (H⁺ and OH⁻) ionic conductivity and chemical reactivity, leaving vast array of neutral electrolytes using metallic salts largely unexplored. One of the challenges is that electrolyte ions cannot simply be treated as point charges. For instance, the adsorption of cations with different valence and solvated ion sizes including Na⁺, Zn²⁺ and Al³⁺, has been studied in highly ordered and compact porous electrodes²⁵. The partially dehydrated divalent Zn²⁺ ion was found to be densely and neatly packed, achieving an unparalleled spatial charge density by balancing the valence and size of charge-carrier ions. Drawing parallels with the findings in porous electrodes, one might wonder whether the ion desolvation is the key for optimizing energy storage in 2D layered conducting materials. To understand the charge storage at confined nanoscale interface, a systematic exploration and deeper comprehension are highly desired.

The present paper provides evidence of the key role of ion partial desolvation on the charge storage mechanism in 2D rGO taken as model material. Thanks to an original combination of several operando techniques including electrochemical quartz crystal microbalance (EQCM) and electrochemical dilatometry (ECD) measurements, we evidenced partial desolvation of cations during cathodic polarization, resulting in enhanced capacitance. In this paper, multilayer rGO particles with conducting graphitic domains and controllable surface oxygen functionalities were used as model materials

to study charge storage mechanism in near-neutral aqueous electrolytes and the effect of ion confinement on the electrochemical behavior. Preliminary studies have already shown a cation-dominated charging mechanism for largely reduced graphene oxide materials²⁶. Ion adsorption kinetics and electrochemical behaviors were also found to be different when changing the type of cation used²⁵⁻²⁷. Key parameters such as charge density, hydration/solvation energies, and specific cation-graphene interactions - like the cation- π interaction - influence interfacial electrochemical activity, making the capacitive charge storage mechanisms of (2D layered) rGO materials more intricate than anticipated.

Here, we aim at clarifying how the ion-carbon interactions and ion desolvation affect the electrochemical behavior of rGO. The electrochemical behavior of rGO electrode was studied in three different electrolytes containing cations including Li⁺, Mg²⁺, and Zn²⁺. Cyclic voltammetry experiments were firstly achieved by using a cavity micro-electrode (CME) setup, which allows for tracking tiny changes in the electrochemical signature, and improves the signal/noise ratio and mitigates ohmic drop. Then, operando EQCM experiments have been used to bring a better understanding about ion fluxes and ion solvation-desolvation occur during the adsorption-desorption process at the rGO surface. Eventually, the operando ECD experiments were made and cross-correlated with EQCM data to better grasp the charge adsorption mechanism in rGO.

Results

Synthesis and physicochemical characterization of rGO

The preparation of the materials is illustrated in Fig. 1a. rGO was synthesized from a graphene oxide (GO) aqueous dispersion at a concentration of 1 mg mL⁻¹ using a mild hydrothermal reaction at 160 °C for 6 h²⁸. Subsequently, to eliminate the residual surface oxygen, a second thermal reduction of the rGO powder was conducted at 900 °C under an Ar atmosphere for 1 h. This product, obtained after a double reduction of GO, is denoted as r²GO. Figure 1b displays the X-ray diffraction (XRD) patterns of both rGO and r²GO powders. The successful synthesis of rGO was validated by the presence of distinct XRD peaks at $2\theta = 24.0^\circ$, corresponding to a d-spacing along the c-axis of 3.7 Å for the (002) diffraction plane and $2\theta = 43.8^\circ$ for the (100) plane. The d-spacing of the (002) plane reflects the interlayer distance between individual graphene layers. The marginally enlarged interlayer distance in our sample may result from the heightened presence of surface functional groups. For r²GO, the interlayer distance decreases to 3.4 Å at $2\theta = 26.2^\circ$, which is attributed to the elimination of various functional groups originally situated between the rGO interlayers²⁹. To identify the oxygen content of rGO, Supplementary Fig. 1 shows the mass loss of rGO during the two-step thermal reduction processes by temperature programmed desorption mass spectrometry (TPD-MS). TPD-MS results indicate that considerable amount of oxygen surface functionalities still exist on the rGO surface after hydrothermal reduction of GO³⁰. Indeed, after quantification of the spilled gas product such as CO and CO₂, the content of surface oxygen is determined as 16.7 wt.% for rGO and 1.3 wt.% for the further reduced sample r²GO.

Due to the residual oxygen-containing surface groups, rGO consistently retains a negative charge in aqueous solutions when the solution's pH exceeds 3³¹. This is confirmed by the zeta potential values of rGO samples in various electrolytes (Fig. 1c), which are negative whatever the electrolyte that has been used. More precisely, we changed the cation type to explore the cation-rGO interactions, given earlier indications that the charge storage mechanism of rGO is predominantly cation-controlled^{26,27}. We maintained the anion as Cl⁻ with electrolyte pH \approx 6 in 0.01 M concentration for zeta potential measurements. The zeta potential for rGO in the presence of LiCl electrolyte was -37 ± 4 mV, consistent with already reported rGO zeta measurement at different pH³¹. Nevertheless, the reported results are focused on the influence of pH without paying attention to the role of cation.

According to our zeta measurements, it turns out that electrolytes containing divalent cations drastically altered the surface charges, as zeta potentials of -11 ± 1 mV and -5 ± 1 mV were measured in MgCl_2 and ZnCl_2 electrolytes, respectively. The important noncovalent cation- π interaction between cations and the delocalized aromatic graphene domains distributed in rGO³² are supposed to be at the origin of the change in the surface charges of rGO³³. In the case of Mg^{2+} , the higher charge density as a divalent cation^{34,35} explains the higher adsorption energy induced by cation- π interaction versus Li^+ ^{35,36}. Therefore, the enrichment of Mg^{2+} onto the rGO surface is expected to effectively screen the surface charge and makes the surface zeta potential less negative than Li^+ . Interestingly, a stronger metal ion- π interaction³⁷⁻³⁹ were observed with transition metal ions - Zn^{2+} in our study - resulting in further decrease of the rGO surface charge. Thus, according to those zeta measurements, the order of surface adsorption at open circuit potential is: $\text{Zn}^{2+} > \text{Mg}^{2+} > \text{Li}^+$.

Cation-dependent electrochemical signature of rGO

In light of the rapid charging kinetics associated with capacitive materials, traditional studies often overlook the tiny differences among electrolytes with similar physiochemical properties; however, we try to address this point in the present work. Figure 2a illustrates a three-electrode setup where a CME is used as the working electrode. The CME consists in a platinum wire enclosed in a silica glass tube. A small amount of active powder material (10^{-7} – 10^{-8} g) is compacted into a cavity of approximately 50 μm in diameter and 20 μm in depth, eliminating the need for conductive agents and binders. Thanks to such electrode design, the measured current is less than few μA , significantly mitigating ohmic drop, which allows for obtaining cyclic voltammograms with better definition. Therefore, we take benefit of this set-up to perform a more accurate electrochemical signature analysis of rGO in three different electrolytes: LiCl , MgCl_2 , and ZnCl_2 (Fig. 2b–d).

As observed in Fig. 2b, in the Li^+ -based electrolyte, there are two current regions: one located between +0.7 V down to +0.4 V vs. Ag/AgCl , with the other between +0.4 V and -0.8 V vs. Ag/AgCl . In both regions, the current plateaus but at different magnitude. According to

some other studies, such current change giving rise to a higher current amplitude at higher overpotentials is highly likely correlated with cation desolvation^{27,40,41}. Interestingly, comparing the divalent Mg^{2+} -based electrolyte to the Li^+ -based electrolyte (Fig. 2c), additional pair of current bumps located at -0.5 V vs. Ag/AgCl is seen in the CV. Meanwhile, a broad current bump emerges in the potential window of -0.5 V to $+0.2$ V vs. Ag/AgCl for the Zn^{2+} -based electrolyte (Fig. 2d)⁴². According to those results, as in each electrolyte anions are the same, these patterns are highly likely correlated with the cation type; moreover, it has also been found out the type of anion does not affect significantly the electrochemical signature (Supplementary Fig. 2). We supposed that the interactions between cations and the rGO seems then to be crucial in controlling the (cation-dominated) electrochemical responses. We further explored the charging kinetics by plotting the logarithm of the bump/peak current against the scan rate. The results suggest a capacitive charging mechanism as current linear increase with scan rate⁴³ (Supplementary Fig. 3). It is worth mentioning, after surface oxygen functionalities have been removed (r^2GO sample), the capacitive current was divided by 2 to 5 (Supplementary Fig. 4), which is attributed to the decrease of surface wettability, and all the CVs display a rectangular shape without any current peaks or bumps. Importantly, the pH of the electrolytes (1 M) shown in Fig. 2 is 6.9 for LiCl , 5.0 for MgCl_2 , and 5.3 for ZnCl_2 . These values are near neutral, despite the slight hydrolysis of multivalent metal cations in water. As a result, cations seem to act as the main charge carrier in our case instead of the hydrated protons, differentiating our study from those examining the faradaic reversible reactions between protons and carbon surface oxygen-containing functionalities.

To further validate the cation- and oxygen-dependent electrochemical behavior, we conducted supplementary Swagelok cell experiments with mass loadings of about 0.7 mg cm^{-2} . CVs shown in Supplementary Fig. 3 mirror a similar trend: while current bumps are present in rGO, they disappear for r^2GO . Furthermore, we calculated the capacitance of both rGO and r^2GO (Supplementary Table 1) across the different electrolytes by integrating the CV with Eq. (1). As expected from previous works⁴⁴, we observed that surface oxygen functionalities considerably enhance capacitance, most possibly attributed

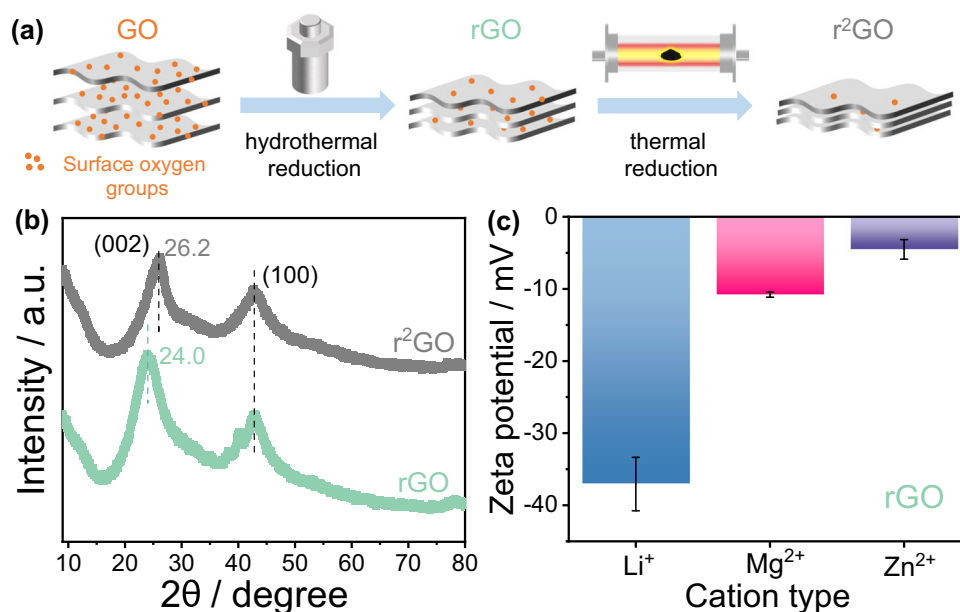


Fig. 1 | Material preparation and physiochemical properties characterization. **a** Experimental procedures of the step reduction of graphene oxide (GO): reduced graphene oxide (rGO) was obtained by first hydrothermal reduction and followed by high-temperature thermal reduction to prepare further reduced material

(marked as r^2GO). **b** X-ray diffraction (XRD) patterns of rGO and r^2GO powder. The 2θ angle corresponding to the 002 crystal plane XRD peaks of GO is shifted from 24.0° (d-spacing 3.7 Å) to 26.2° (d-spacing 3.4 Å) after reduction. **c** Zeta potential of rGO in the presence of aqueous 0.01 M LiCl , MgCl_2 and ZnCl_2 solutions.

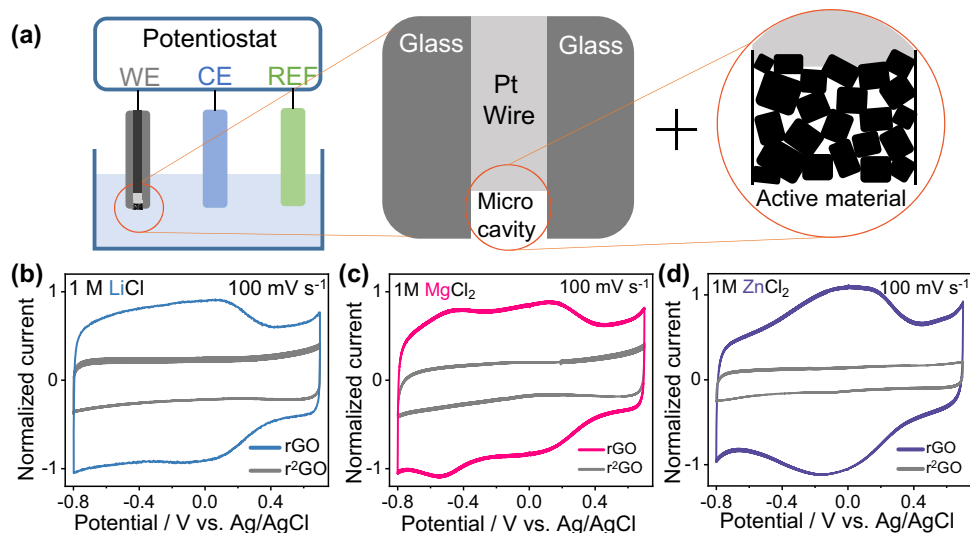


Fig. 2 | Electrochemical response of rGO in three different electrolytes. Experimental set-up and electrochemical behavior of rGO in different neutral aqueous electrolytes in a three-electrode configuration with a cavity micro-electrode (CME) serving as working electrode. **a** illustration of the three-electrode

set-up: the active material (rGO or r²GO) is encapsulated in the cavity (50 μm diameter and 20 μm depth). Cyclic voltammetry curves of rGO and r²GO in the electrolyte of **(b)** 1 M LiCl, **(c)** 1 M MgCl₂, and **(d)** 1 M ZnCl₂ at a scan rate of 100 mV s⁻¹.

to increased exposure of active sites and enhanced ion transport due to improved wetting capabilities.

Two regions of cation solvation-desolvation

EQCM stands as an effective operando technique for monitoring ionic fluxes at polarized electrode interfaces. According to the Sauerbrey equation⁴⁵, the electrode mass change (Δm) arising from the electrochemical ion adsorption/desorption in/from the carbon electrode is negatively correlated to the change of the quartz resonant frequency (Δf) (see Eq. (2)): a decrease in Δf indicates an increase in Δm and vice versa. Figure 3a–c show the CV curves in three electrolytes obtained using the EQCM cell, which are similar to those previously obtained with CMEs. As a whole, in all plots, the frequency change Δf decreases (Δm increases) during cathodic polarization and Δf increases (Δm decreases) during anodic polarization. Globally speaking, the overall trend of Δf keeps the same regardless of cation type.

The next part is focused on the detailed analysis of the Δf curves in conjunction with both the potential and the cation need to be further analyzed. The cathodic polarization span, ranging from 0.5 V to -0.8 V vs. Ag/AgCl, was chosen for a deeper examination of the cation-dependent charge storage mechanism. According to the minimal change in motional resistance (ΔR) shown in Supplementary Fig. 5, the coating onto the quartz can be considered as rigid so that the gravimetric model of the Sauerbrey equation remains valid. Δm was plotted against the accumulated charges (ΔQ) as depicted in Fig. 3d–f. ΔQ was calculated from the integration of the CV curves (current over time) and was normalized to zero at 0.5 V vs. Ag/AgCl, where the potential of zero charge seems to be. Δm increases with the accumulated cathodic charges, basically suggesting a cation adsorption mechanism, which is in a good agreement with other EQCM studies that reported a predominantly cation-driven charging mechanism for alkali metal ions and rGO^{26,27}. Two distinct regions can be defined from the Δm - ΔQ plots (Fig. 3d–f): Region I displays a steady growth with a constant slope, while Region II shows a smooth transition and a reduced slope. These regions are particularly pronounced for divalent Zn²⁺. Generally, the steady mass increase at low polarized potentials in Region I, driven by cation adsorption, is shared by the three cations, but with different slopes. Interestingly, the charge storage in Region II is more efficient considering less mass is required to balance equivalent ΔQ at large polarizations when compared to Region I. Regarding the cations,

especially multivalent ones, they are known to be highly hydrated in bulk electrolyte due to their high charge density and solvation free energy. So, a possible explanation of the slope decrease of the Δm - ΔQ plots at high charge (large polarizations) points towards cation dehydration.

To have a deeper look into the ion dehydration process during cathodic polarization (cation adsorption) and clarify the relationship between ion dehydration and the cation-dependent current responses, we used multi-linear fitting of the Δm - ΔQ plots. Each calculated slope within the linear fitting scale is interpreted as a molecular weight according to Faraday's law (Eq. (3)), as illustrated in Fig. 3g–i. The white dashed lines show the multi-linear fitting results, with the cut-off potentials for each scale indicated by vertical black dashed lines. Finally, we plotted both the current and the calculated molecular weight versus the electrode potential to establish the correlation between the observed current bumps and ion dehydration.

In Region I, all three cations exhibited a similar trend: an increasing current paralleled by cation adsorption with a stable molecular weight. The molecular weight of these hydrated cations reveals varying hydration levels among the cations. For instance, the average solvation number for monovalent Li⁺ ion is 4.6 (Fig. 3g), while Mg²⁺ (Fig. 3h) and Zn²⁺ (Fig. 3i) average 18.4 and 12.7 H₂O molecules, respectively. Interestingly, the idea of each Li⁺ carrying 4–5 H₂O molecules at mild polarization was also proposed from electrogravimetric impedance studies²⁷. Due to higher hydration free energy for multivalent cations, the average solvation number for Mg²⁺ is found to be as high as 18.4, which well aligns with previous reports with a value of 18, counting both the primary coordination number (6) and the secondary solvation shell number (12)^{46,47}. In the case of ZnCl₂ electrolyte, 75% of Zn(H₂O)_n²⁺ likely serves as the primary charge carrier (25% of Zn(H₂O)_nCl⁺) in 1 M ZnCl₂ solutions, while the potential formation of zinc clusters usually occurs in highly concentrated ZnCl₂ solutions^{48,49}. Besides, the electrochemical response shows no anion dependence when altering the anion including NO₃⁻ and SO₄²⁻ (Supplementary Fig. 2), which also evidences that the electrochemical response are related to cation Zn²⁺ but not the zinc clusters. Given the comparable hydration free energy of Zn²⁺ (-1995 kJ mol⁻¹) to Mg²⁺ (-1830 kJ mol⁻¹)²⁵, an average solvation number of 12.7 possibly include the primary hydration shell (6) and extended hydration shells⁵⁰.

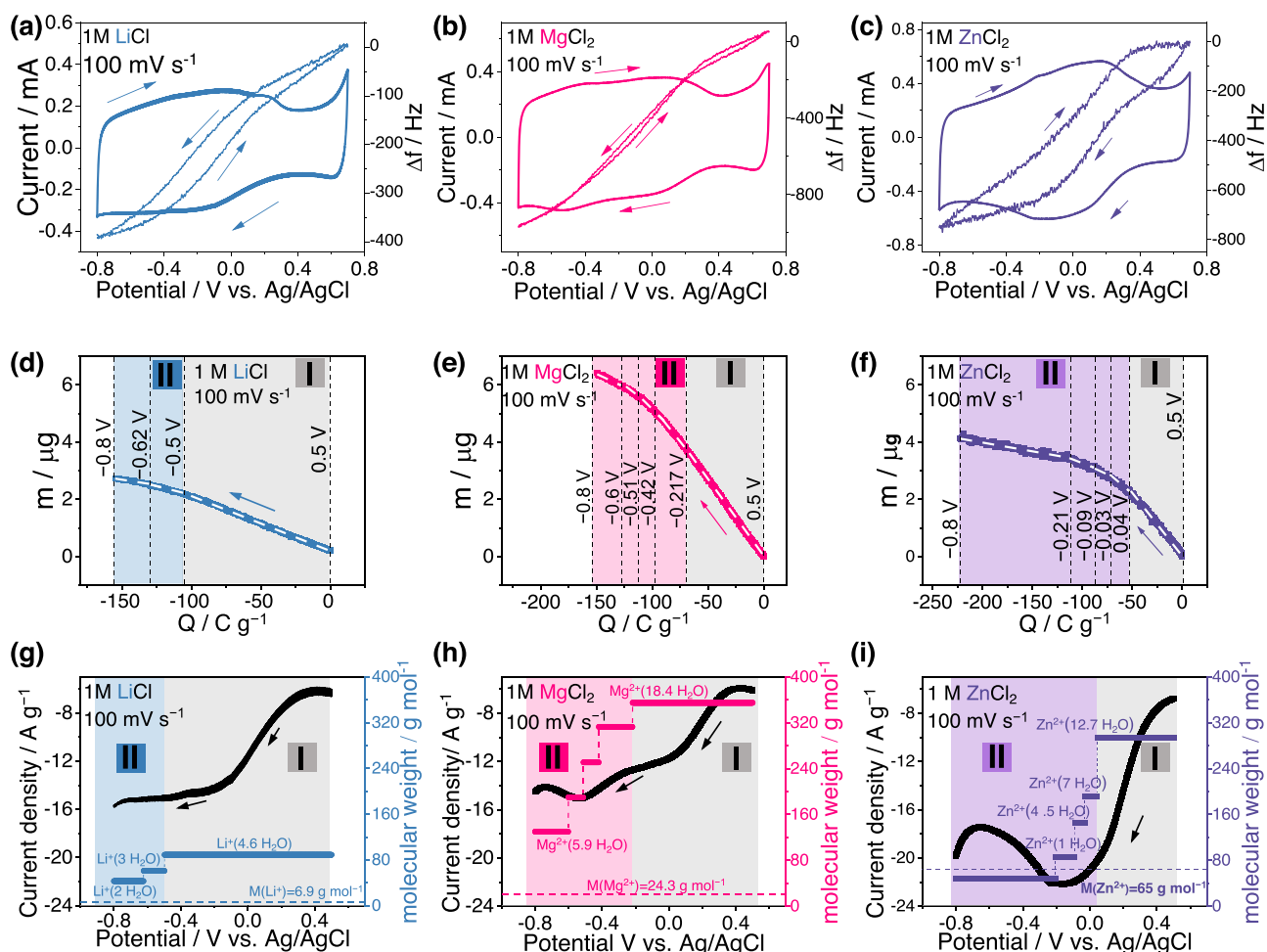


Fig. 3 | Electrochemical quartz crystal microbalance (EQCM) measurement of rGO in three different electrolytes. The electrolytes are 1M LiCl (**a, d, g**), 1M MgCl₂ (**b, e, h**) and 1M ZnCl₂ (**c, f, i**), the potential scan rate is 100 mV s⁻¹. **a, b, c** Cyclic voltammograms with corresponding frequency change (Δf), indicated by arrows representing the polarization direction and Δf change direction. **d, e, f** Electrode mass change (Δm) is plotted against accumulated charges (ΔQ), with two distinct regions shaded in different colors. The white dashed lines

show the multi-linear fitting results, with cut-off potentials of each scale separated in vertical black dashed lines. Region I is highlighted in light gray and region II is delineated in blue (Li⁺), pink (Mg²⁺), and light purple (Zn²⁺). **g, h, i** Current and equivalent molecular weight obtained from the slope of Δm - ΔQ plots by Faraday's equation is plotted vs. potential, presented in the same two regions as depicted in (**d-f**). Dashed lines in (**g-i**) represent the theoretical molecular weight of the naked cations.

When moving to Region II, the calculation of the molar weight from the Δm - ΔQ plots highlighted the existence of a dehydration process at larger cathodic potentials for the highly solvated cations (Mg²⁺ and Zn²⁺), while minor dehydration was observed for Li⁺. This progressive dehydration seems to be associated with the presence of a current broad peak (bump). The hydrated Mg²⁺ discards a part of its solvation shell, transitioning from Mg²⁺ (18.4 H₂O) to Mg²⁺ (5.9 H₂O), which well aligns with the current bump situated at -0.5 V (Fig. 3h). Similarly, the Zn²⁺ current peak located at -0.3–0 V corresponds to a substantial removal of its solvation shell (Fig. 3i). Meanwhile, the current for Li⁺ remained steady in Region II, as a result of minor structural change in the Li⁺ solvation shell. To summarize, the pronounced dehydration observed for Mg²⁺ and Zn²⁺ during high cathodic polarization (Region II) is assumed to be at the origin of the observed current bumps. This might be due to the reduced effective ion size after removing solvation shells, allowing for a more efficient screening of the electrode charge by the adsorbed ionic species, as desolvated cations are expected to get closer to the carbon surface favoring specific interactions^{9,10,51}.

The potential-driven step dehydration process provides intriguing insights into the dynamics of cation-electrode interactions and their impact on the electrochemical behavior of the system. The

current bump for Mg²⁺ being located at a more negative potential (-0.5 V) versus that observed for Zn²⁺ (-0.1 V), it suggests a more substantial energy barrier for the electrochemical adsorption of Mg²⁺, resulting in easier Zn²⁺ dehydration and subsequent adsorption into the carbon electrode. This observation is consistent with the zeta potential measurements, suggesting a more robust interaction of Zn²⁺ with rGO. To sum up, the cation-specific electrochemical behaviors studied via both CME and EQCM techniques were found to be closely tied to the interactions between the cations and the rGO electrode. In the case of the Mg²⁺-based electrolyte, the current bump (-0.5 V) arises from the strong Mg²⁺-rGO interaction, often referred as cation- π interaction. This cation- π interaction is even further increased after dehydration, as the discard of thick solvation shell strengthens the Mg²⁺- π bond. This interaction is notably weaker in the Li⁺-based electrolyte, thus explain the absence of current peaks / bumps for Li⁺ adsorption in both region I and II. In contrast, Zn²⁺ exhibits a distinct behavior. As a transition metal cation, Zn²⁺ maintains a (+II) oxidation state, mirroring that of Mg²⁺ and sharing comparable hydration free energies and effective ion sizes²⁵. Nevertheless, the inherent characteristics of Zn²⁺, especially its filled 3d shell and an available 4s state, strengthen its interaction with the conjugated π systems of

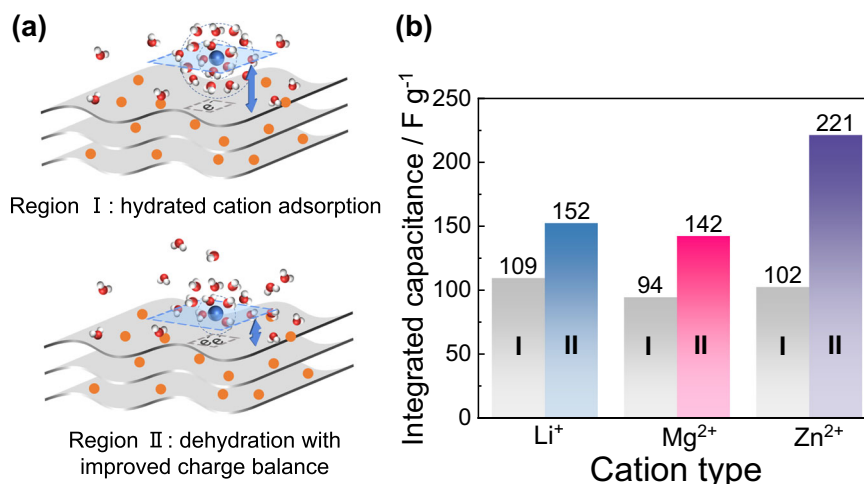


Fig. 4 | Two regions of charging process and capacitance enhancement. **a** Schematic illustration of cation-dependent two regions charge storage process of rGO when applying aqueous near neutral electrolyte. **b** Integrated capacitance of two regions when applying different electrolytes.

rGO^{38,39}. This specific interaction is often termed as transition metal- π interaction. Consequently, the stronger Zn²⁺-rGO interaction facilitate Zn²⁺ dehydration and shift the current bump potential to less negative region, causing a confluence of regions I and II, resulting in a broadened current bump in the CV profile.

Cation desolvation-induced capacitance enhancement

Figure 4a proposes a schematic illustration of the charge storage processes across the two regions. During cathodic polarization, in region I, solvated or hydrated cations predominantly balance the electronic charges on the rGO surface. This is typical of the electric double layer capacitor (EDLC) behavior, where charge storage happens at the electrode/electrolyte interface without faradaic (redox) processes. Conversely, in region II, cations after losing a part of their solvation shell can approach closer to the electrode surface, leading to more efficient charge screening and storage. This in line with a molecular dynamic study reporting that the more counter-ion interact with the surface, the more the effective counter-charge¹⁰. These two charging processes were further confirmed by the capacitance analysis (Fig. 4b and Eq. (1)): the nearly constant integrated capacitance ($100 \pm 10 \text{ F g}^{-1}$) calculated in region I for all cations well agrees with the expected EDLC behavior. This region, as suggested, is dominated by the adsorption of solvated cations: the polarizable water molecules of the solvation shell highly screen the ionic charges, making it less effective to balance the electronic charges from electrode side. The dramatic capacitance increase observed in region II can originate from partial desolvation of the cations, that results in a more efficient charge screening thanks to the shortened carbon-cation distance, resulting in higher capacitance.

The difference in capacitance increase observed for the different cations (Li⁺, Mg²⁺, and Zn²⁺) in region II highlights the specific interactions between each cation and the rGO. The pronounced capacitance enhancement for Zn²⁺ suggests stronger cation interaction with rGO after dehydration. This mirrors the discussions around transition metal- π interactions mentioned previously. The magnitude of enhancement follows the order: Zn²⁺ > Mg²⁺ > Li⁺, which well aligns with the expected cation-rGO interaction strength. In essence, the findings reiterate the significance of cation hydration and its dynamic interaction with rGO in driving the electrochemical behavior.

Charge storage active sites and charging process

Due to the remarkable capacitance achieved in Zn²⁺ electrolyte, we tried to get further understanding about the Zn²⁺ charge storage

mechanism. X-ray diffraction (XRD) measurements were conducted to investigate the active sites. It is worth noting that charge storage sites in reduced graphene oxide (rGO) materials can have two origins: (1) the graphitic-like space situated between the graphene interlayers, corresponding to the $2\theta = 24^\circ$ (001) diffraction peak (Fig. 1b); and (2) the gallery domains formed by the stacking of multilayer rGO particles, characterized by the $2\theta = 11\text{--}13^\circ$ diffraction peaks⁵². In our case, rGO particles do not exhibit well defined and organized gallery domains (Supplementary Fig. 6) because of a random stacking of rGO particles. Besides, an operando XRD analysis evidenced the absence of shift of the (002) diffraction peak of rGO during cycling in Zn²⁺-containing electrolyte⁴², evidencing that Zn²⁺ cannot intercalate between the graphene interlayers of rGO. Consequently, the charge storage process primarily takes place at the surface of rGO particles in the gallery domains without intercalation.

To get further insights about the charge storage mechanism, we used ECD technique to track the changes in thickness/displacement under polarization. Dilatometry is a well-established technique utilized to measure the expansion or contraction of macroscopic samples. ECD measures the expansion/contraction of a sample during polarization. Figure 5 presents operando ECD plots of the rGO electrode during polarization in different electrolytes. Initially, the pristine electrode, as measured by SEM, is approximately 4 μm -thick (Supplementary Fig. 7). The thin active material coating and the scan rate of 20 mV s^{-1} have been optimized to offset the ECD cell resistance arising from the thick frit separator. Figure 5a–c show the CVs of the rGO samples in the different electrolytes and the electrode displacement (along the z-axis) recorded during the polarization. The displacement was determined by normalizing the thickness change to the pristine thickness of the rGO electrode. Drawing parallels with EQCM data analysis, we examined the cathodic scan from 0.5 V to -0.8 V . In these plots, both accumulated charges and displacement were normalized to zero at 0.5 V. A direct correlation of the (increase of the) displacement with the accumulated charge can be observed, indicating volume expansion due to ion adsorption. Notably, the thickness variation in all three electrolytes were small (below 1%), which well aligns with a typical EDL-like ion adsorption process^{33,34}. Typically, intercalation processes would result in more pronounced volume change^{55,56}. A linear volume expansion with charge accumulation was found for both Li⁺ (Fig. 5d) and Mg²⁺ (Fig. 5e) adsorption. This electrostatic volume expansion results from the transportation of solvated cations from the electrolyte bulk into the capacitive storage sites, which are the spaces formed by the random stacking of rGO particles (Fig. 5g, h). The constant ion sizes result in a continuous linear change of displacement in region I.

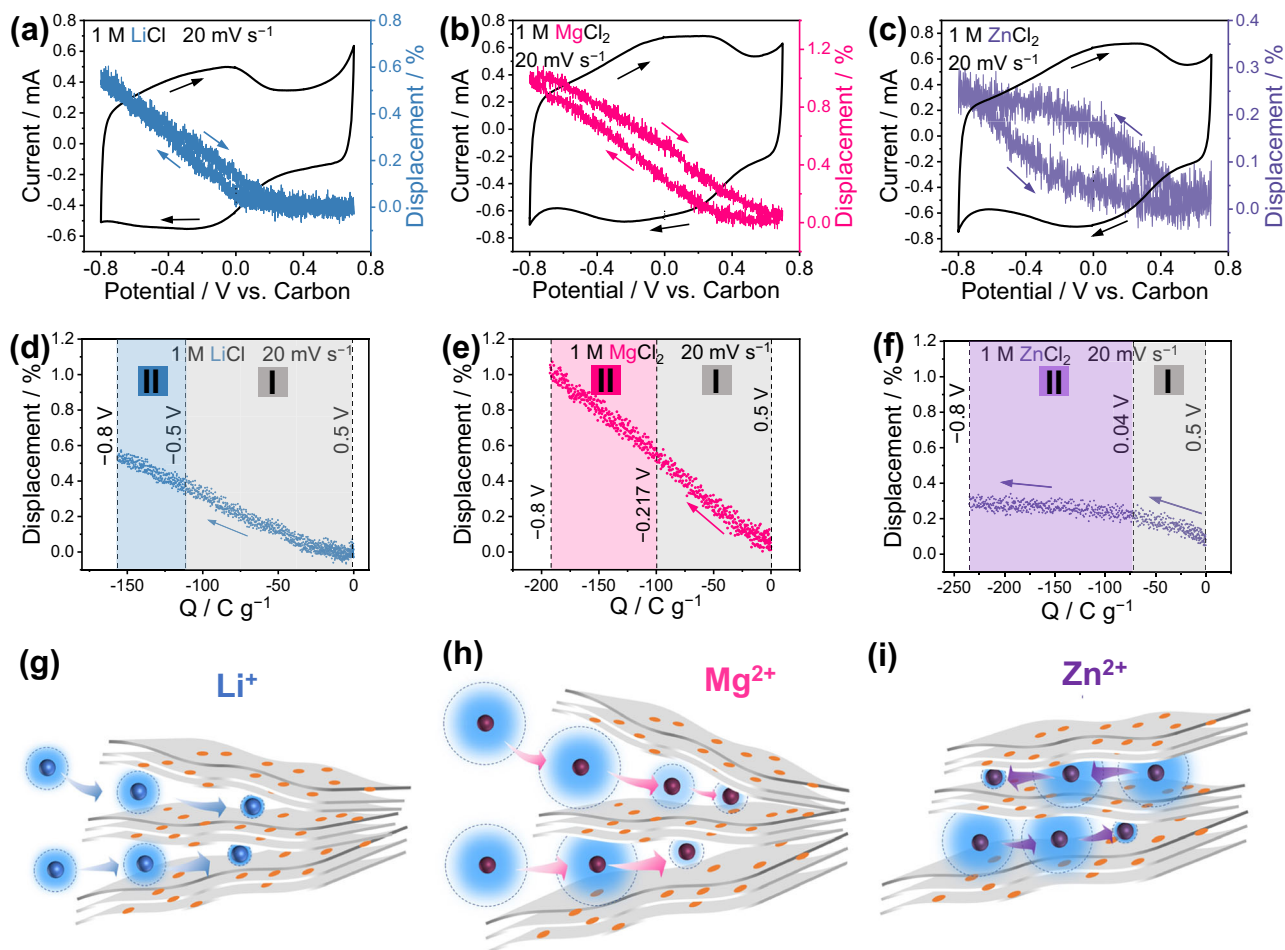


Fig. 5 | Charge storage active sites and charging mechanism related with cation desolvation. Dilatometer measurements of rGO with different electrolytes by using a three-electrode operando electrochemical dilatometry cell. **a–c** Cyclic voltammograms at scan rate of 20 mV s^{-1} with corresponding displacement change, indicated by arrows representing the polarization direction and displacement change direction. **d–f** Displacement change (also named “height change”) of rGO electrodes with different electrolytes at potential scan rate of 20 mV s^{-1} .

The zero displacement was set at 0.5 V vs. carbon. **g–i** Schematic illustration depicting the charge storage sites within rGO particles and the charging process accompanied by cation dehydration. Noting that polarization is not depicted in this Figure. The hydration shell encompassing various cations is delineated in varying shades of light blue, showing hydrated cation size changes. Electrolytes are **(a, d, g)** 1 M LiCl , **(b, e, h)** 1 M MgCl_2 , and **(c, f, i)** 1 M ZnCl_2 .

Interestingly, the electrode expansion rate remains constant, even when desolvation occurs in region II. This suggests that main effective charges in region II could be the solvated ions already present in the electrode (region I). As a whole, the hydrated cation continues to expand the electrodes at shallow adsorption sites near the edges of multilayer particles in region II and the partial dehydration locally occurs by ion and solvent rearrangement, at less accessible surface, that is inside the particles (Fig. 5g, h)^{27,57}. However, the behavior of Zn^{2+} diverges from other cations (Fig. 5f). There is only a minimal volume expansion for Zn^{2+} , which may be attributed to the robust binding affinity between Zn^{2+} and rGO. Zn^{2+} are presumably already present in energy storage sites at open circuit potential (Fig. 5i), and the available Zn^{2+} balance the charge more efficiently by partial cation desolvation, accompanied by a minimal expansion. Overall speaking, the charge storage of Zn^{2+} is more efficient than Li^+ and Mg^{2+} , which also explain the higher average capacitance of Zn^{2+} -based electrolyte.

In conclusion, the presented study offers significant insights into the multifaceted electrochemical behaviors depending on cation-rGO interactions, and we found cation-dependent and oxygen-determined specific electrochemical behavior by using CMEs. The charge storage mechanism was firstly studied by EQCM, that suggested the presence of two distinct regions: one dominated by solvated cation adsorption, and the other characterized by cation dehydration, leading to

enhanced charge compensation on the rGO surface. Notably, different cations exhibited varying degrees of interaction strength with rGO, with the sequence being $\text{Zn}^{2+} > \text{Mg}^{2+} > \text{Li}^+$. The dynamic cation-rGO interaction at different polarizations are responsible for the varying electrochemical behavior and energy storage capacity. Furthermore, operando ECD revealed the presence of shallow adsorption sites and accessible, confined sites for Li^+ and Mg^{2+} adsorption, resulting in constant volume expansion during charge. Differently, the electrode volume change in Zn^{2+} charge storage mechanism was less, about 0.2% , suggesting a reorganization of Zn^{2+} present in the electrode at OCV, by solvent reorganization, resulting in partial desolvation. Such findings underscore the potential for tailored electrolyte and electrode design, aiming to optimize capacitive charge storage for high-performance supercapacitors, batteries, and other energy storage systems.

Methods

Material preparation

Graphene oxide (GO) was firstly prepared by a modified Hummers method⁵⁸. Then 1.0 mg mL^{-1} of GO aqueous dispersion was used as the precursor to prepare the reduced graphene oxide (rGO) by hydrothermal reduction at $160 \text{ }^\circ\text{C}$ for 6 h and followed by freeze drying. Afterwards, the further reduced sample (r^2GO) was made from rGO

powder by an annealing thermal reduction at 900 °C in argon atmosphere for 1 h, with a heating rate of 10 °C min⁻¹.

Electrode preparation

First, we commenced the electrode preparation process by thoroughly blending the active powder material, which could be either GO or rGO, together with the conducting additives - carbon black - in a PVDF/NMP solution. The mass ratio used for this combination was 8:1:1 (active material: carbon black: PVDF), and the mixture was stirred overnight. Subsequently, the resulting homogeneous slurry was applied onto the surface of a titanium foil (as current collector) using the doctor blade method. After this step, the electrodes were considered ready for use after vacuum dried at 80 °C for 24 h. Following the drying process, the electrodes were cut into small disks with a diameter of 12 mm. The prepared electrodes were utilized in both three-electrode Swagelok cells and in-situ dilatometry measurements, with a loading mass of approximately 0.7 mg cm⁻².

Electrochemical tests

The cyclic voltammetry (CV) experiments were conducted using the Biologic VMP-300 electrochemical workstation (France), employing a three-electrode cell setup. The potential range in these CV tests spanned from -0.8 to +0.7 V vs. Ag/AgCl (saturated in a KCl solution). This potential range remained consistent across all our experiments, encompassing CME tests, operando EQCM, and operando ECD measurements.

The choice of scan rate varied based on factors such as the electrochemical cell resistance and the loading mass of the electrodes. For instance, in the CME configuration, where the ohmic drop was negligible, we adopted a high scan rate of 500 mV s⁻¹. This scan rate aimed to enhance current amplification that comes from the active materials and minimize background current. Similarly, in the case of operando EQCM measurements, a slightly reduced scan rate of 100 mV s⁻¹ was employed due to the higher loading mass (in the microgram scale) when compared to the CME test, which featured a loading mass in the nanogram range. For the operando ECD cell, the higher cell resistivity was primarily attributed to the thick frit serving as a separator, making it the most resistive among the different cell configurations, including Swagelok cells, CME setup and EQCM cells. Consequently, a relatively lower scan rate of 20 mV s⁻¹ was selected. It's worth emphasizing that the choice and careful adjustment of these scan rates, tailored to the specific characteristics of each cell configuration, played a crucial role in elucidating the cation-dependent electrochemical behaviors.

The reference electrode for operando ECD tests is the YP-50F activated carbon electrode film, which has similar potential ($\Delta U = 50$ mV) with Ag/AgCl electrode that we used. And the potential of YP50F reference electrode is relatively stable and not polarizable⁵⁹.

Capacitance calculation

The capacitance of rGO was calculated in specific regions of interest. In each region, we integrate of current with respect to time, and subsequently divided it by the potential window of each region, as indicated by Eq. (1):

$$C = \frac{\int_0^t i dt}{Vm} \quad (1)$$

where C is the gravimetric capacitance (F g⁻¹), t is the recorded time range (s), i is the response current (A), V is the potential window (V), and m is the active material mass (g).

Electrochemical quartz crystal microbalance (EQCM) measurements

To prepare the EQCM samples, Au-coated quartz crystals (basic oscillating frequency of 9 MHz, AWSensors, Spain) were

homogeneously coated with a slurry comprising 90 wt.% of active material rGO, 10 wt.% of polyvinylidene fluoride (PVDF, provided by Arkema), and dissolved in N-Methyl-2-pyrrolidone (NMP, Sigma-Aldrich). The slurry concentration is 1 mg mL⁻¹ (taking all solid mass into account). The coating process was conducted using a spray gun, with a distance 10–15 cm away from the quartz surface. 170 μ L of the slurry was added into the spray gun for each quartz, and the coated quartz was vacuum dried overnight at 80 °C before use. The loading mass of the active material coated on the quartz surface is around 20–40 μ g cm⁻². The calibration plot (frequency vs. mass for different weight loading) as shown in Supplementary Fig. 8, indicate that the rGO weight loading selected for EQCM experiments falls within the linear change of frequency vs. mass, evidencing the presence of a rigid rGO coating in air. Subsequently, the 3-electrode EQCM setup was applied with the coated QCM quartz served as working electrode, the platinum wire functioned as counter electrode, and Ag/AgCl employed as reference electrode. These three electrodes were placed within a glassware container and immersed in different aqueous electrolyte that we studied.

All EQCM electrochemical measurements were performed using a Maxtek RQCM system in conjunction with the Biologic potentiostat for simultaneous EQCM and electrochemical measurements. The EQCM data was analyzed according to the Sauerbrey equation⁴⁵ (Eq. (2)):

$$\Delta m = -C_f \Delta f \quad (2)$$

where Δm represents the change in mass of the coating, and C_f denotes the sensitivity factor of the crystal. The sensitivity factor for the coated quartz was determined through a copper deposition experiment conducted in a solution containing 10 mM CuSO₄ mixed with 0.5M H₂SO₄. In the stable CV cycles, C_f was calculated as 6.98 ng·Hz⁻¹ (or 5.43 ng·Hz⁻¹ cm⁻², considering the Au crystal electrode surface area of 1.28 cm²). For consistency in results, a few cycles were run prior to initiating EQCM measurements to ensure that the data began from stable and reproducible electrochemical signatures.

For calculating the molecular weight from Δm - ΔQ plots, we apply the Faraday's law (Eq. (3)):

$$\frac{\Delta m}{\Delta Q} = \frac{M_w}{nF} \quad (3)$$

where Δm represents the change in mass of the coating, and ΔQ was the accumulated charges obtained by integrating current with time, n is the ion valence number and F is the Faraday constant (96 485 C mol⁻¹), then M_w is the molecular weight. In our case, $\Delta m/\Delta Q$ is the slope of Δm - ΔQ plot, we consider the charge carriers are Li⁺, Mg²⁺ and Zn²⁺ in Fig. 3, so the n is 1 for Li⁺ and 2 for Mg²⁺ and Zn²⁺ to calculate the molecular weight.

Other characterizations

XRD data were collected a D4 X-ray diffractometer (Bruker, Germany) equipped with CuK α radiation ($\lambda = 0.154$ nm). Scanning electron microscope (SEM) measurements were conducted on TESCAN VEGA3. Zeta potential was determined by Zetasizer Nano ZS90 with (Malvern Co., UK). Samples were prepared by preparing rGO dispersions in 0.01 M salt solutions containing LiCl, MgCl₂ and ZnCl₂. All the samples present similar pH at around 6 as tested by pH paper. Zeta potential was carried out every 10 measurements for each sample and every 20 times scan for each measurement. The mean value and distribution as indicated by the phase plot are good, then we present the mean value among 10 times measurements with error bars. The in-situ displacement of electrode was measured by an ECD-3-nano electrochemical dilatometer, the dilatometer was placed inside an oven with fixed temperature at 25 °C during the whole measurement. TPD-MS measurement was conducted under the Ar atmosphere 75 mL min⁻¹. rGO

powder was placed in a thermo-balance and heat up to 900 °C at a heating rate of 10 °C min⁻¹. The resulted decomposition products were monitored by online mass spectrometry (Skimmer, Netzsch, Germany).

Reporting summary

Further information on research design is available in the Nature Portfolio Reporting Summary linked to this article.

Data availability

The data that support the findings of this study are available from the corresponding author upon reasonable request.

References

- Bard, A. J., Faulkner, L. R. *Electrochemical methods: fundamentals and applications* (John Wiley & Sons, 2001).
- Wu, J. Z. Understanding the electric double-layer structure, capacitance, and charging dynamics. *Chem. Rev.* **122**, 10821–10859 (2022).
- Fleischmann, S. et al. Pseudocapacitance: from fundamental understanding to high power energy storage materials. *Chem. Rev.* **120**, 6738–6782 (2020).
- Schoetz, T. et al. Disentangling faradaic, pseudocapacitive, and capacitive charge storage: a tutorial for the characterization of batteries, supercapacitors, and hybrid systems. *Electrochim. Acta* **412**, 140072 (2022).
- Jaugstetter, M., Blanc, N., Kratz, M. & Tschulik, K. Electrochemistry under confinement. *Chem. Soc. Rev.* **51**, 2491–2543 (2022).
- Shao, H., Wu, Y. C., Lin, Z., Taberna, P. L. & Simon, P. Nanoporous carbon for electrochemical capacitive energy storage. *Chem. Soc. Rev.* **49**, 3005–3039 (2020).
- Xu, F. et al. Facile synthesis of ultrahigh-surface-area hollow carbon nanospheres for enhanced adsorption and energy storage. *Nat. Commun.* **6**, 7221 (2015).
- Chmiola, J. et al. Anomalous increase in carbon capacitance at pore sizes less than 1 nanometer. *Science* **313**, 1760–1763 (2006).
- Liu, Y. M., Merlet, C. & Smit, B. Carbons with regular pore geometry yield fundamental insights into supercapacitor charge storage. *ACS Cent. Sci.* **5**, 1813–1823 (2019).
- Merlet, C. et al. Highly confined ions store charge more efficiently in supercapacitors. *Nat. Commun.* **4**, 2701 (2013).
- Chmiola, J., Largeot, C., Taberna, P. L., Simon, P. & Gogotsi, Y. Desolvation of ions in subnanometer pores and its effect on capacitance and double-layer theory. *Angew. Chem. Int. Ed.* **47**, 3392–3395 (2008).
- Lin, R. et al. Solvent effect on the ion adsorption from ionic liquid electrolyte into sub-nanometer carbon pores. *Electrochim. Acta* **54**, 7025–7032 (2009).
- Fleischmann, S. et al. Continuous transition from double-layer to faradaic charge storage in confined electrolytes. *Nat. Energy* **7**, 222–228 (2022).
- Augustyn, V., Simon, P. & Dunn, B. Pseudocapacitive oxide materials for high-rate electrochemical energy storage. *Energy Environ. Sci.* **7**, 1597–1614 (2014).
- Simon, P. & Gogotsi, Y. Confined water controls capacitance. *Nat. Mater.* **20**, 1597–1598 (2021).
- Shan, X. et al. Structural water and disordered structure promote aqueous sodium-ion energy storage in Sodium-Birnessite. *Nat. Commun.* **10**, 4975 (2019).
- Boyd, S. et al. Effects of interlayer confinement and hydration on capacitive charge storage in Birnessite. *Nat. Mater.* **20**, 1689–1694 (2021).
- Charles, D. S. et al. Structural water engaged disordered vanadium oxide nanosheets for high capacity aqueous potassium-ion storage. *Nat. Commun.* **8**, 15520 (2017).
- Mitchell, J. B. et al. Confined interlayer water promotes structural stability for high-rate electrochemical proton intercalation in tungsten oxide hydrates. *ACS Energy Lett.* **4**, 2805–2812 (2019).
- Mineo, G., Bruno, E. & Mirabella, S. Advances in WO₃-based supercapacitors: state-of-the-art research and future perspectives. *Nanomater* **13**, 1418 (2023).
- Fan, X. Z., Lu, Y. H., Xu, H. B., Kong, X. F. & Wang, J. Reversible redox reaction on the oxygen-containing functional groups of an electrochemically modified graphite electrode for the pseudo-capacitance. *J. Mater. Chem.* **21**, 18753–18760 (2011).
- He, Y. et al. Capacitive mechanism of oxygen functional groups on carbon surface in supercapacitors. *Electrochim. Acta* **282**, 618–625 (2018).
- Shao, H. et al. Unraveling the charge storage mechanism of Ti₃C₂T_x MXene electrode in acidic electrolyte. *ACS Energy Lett.* **5**, 2873–2880 (2020).
- Liu, L. Y. et al. Tuning the surface chemistry of MXene to improve energy storage: example of nitrification by salt melt. *Adv. Energy Mater.* **13**, 2202709 (2023).
- Ma, H. et al. Maximization of spatial charge density: an approach to ultrahigh energy density of capacitive charge storage. *Angew. Chem. Int. Ed.* **59**, 14541–14549 (2020).
- Gao, W., Debiecme-Chouvy, C., Lahcini, M., Perrot, H. & Sel, O. Tuning charge storage properties of supercapacitive electrodes evidenced by in situ gravimetric and viscoelastic explorations. *Anal. Chem.* **91**, 2885–2893 (2019).
- Goubaa, H. et al. Dynamic resolution of ion transfer in electrochemically reduced graphene oxides revealed by electrogravimetric impedance. *J. Phys. Chem. C* **121**, 9370–9380 (2017).
- Shao, H. et al. Synthesis of MAX phase nanofibers and nanoflakes and the resulting MXenes. *Adv. Sci.* **10**, 2205509 (2023).
- Alam, S. N., Sharma, N. & Kumar, L. Synthesis of Graphene Oxide (GO) by modified hummers method and its thermal reduction to obtain reduced Graphene Oxide (rGO). *Graphene* **06**, 1–18 (2017).
- Acik, M. et al. The role of oxygen during thermal reduction of graphene oxide studied by infrared absorption spectroscopy. *J. Phys. Chem. C* **115**, 19761–19781 (2011).
- Konkena, B. & Vasudevan, S. Understanding aqueous dispersibility of graphene oxide and reduced graphene oxide through pK a measurement. *J. Phys. Chem. Lett.* **3**, 867–872 (2012).
- Kumpf, R. A. & Dougherty, D. A. A mechanism for ion selectivity in potassium channels - computational studies of Cation- π interactions. *Science* **261**, 1708–1710 (1993).
- Jeong, S. Y. et al. Highly concentrated and conductive reduced graphene oxide nanosheets by monovalent Cation- π interaction: toward printed electronics. *Adv. Funct. Mater.* **22**, 3307–3314 (2012).
- Xia, X. M. et al. Unexpectedly efficient ion desorption of graphene-based materials. *Nat. Commun.* **13**, 7247 (2022).
- Yi, R. B. et al. Ultrahigh permeance of a chemical cross-linked graphene oxide nanofiltration membrane enhanced by Cation- π interaction. *RSC Adv.* **9**, 40397–40403 (2019).
- Shi, G. et al. Ion enrichment on the hydrophobic carbon-based surface in aqueous salt solutions due to Cation- π interactions. *Sci. Rep.* **3**, 3436 (2013).
- Demircan, C. A. & Bozkaya, U. Transition metal Cation- π interactions: complexes formed by Fe²⁺, Co²⁺, Ni²⁺, Cu²⁺, and Zn²⁺ binding with benzene molecules. *J. Phys. Chem. A* **121**, 6500–6509 (2017).
- Marshall, M. S., Steele, R. P., Thanthiriwatt, K. S. & Sherrill, C. D. Potential energy curves for Cation- π interactions: off-axis configurations are also attractive. *J. Phys. Chem. A* **113**, 13628–13632 (2009).
- Sinnokrot, M. O. & Sherrill, C. D. High-accuracy quantum mechanical studies of π - π interactions in benzene dimers. *J. Phys. Chem. A* **110**, 10656–10668 (2006).

40. Hu, L. T., Guo, D. Q., Feng, G., Li, H. Q. & Zhai, T. Y. Asymmetric behavior of positive and negative electrodes in carbon/carbon supercapacitors and its underlying mechanism. *J. Phys. Chem. C* **120**, 24675–24681 (2016).
41. Urita, K., Ide, N., Isobe, K., Furukawa, H. & Moriguchi, I. Enhanced electric double-layer capacitance by desolvation of lithium ions in confined nanospaces of microporous carbon. *ACS Nano* **8**, 3614–3619 (2014).
42. Shao, Y. Y. et al. Regulating oxygen substituents with optimized redox activity in chemically reduced graphene oxide for aqueous Zn-ion hybrid capacitor. *Adv. Funct. Mater.* **31**, 2007843 (2021).
43. Wang, J., Polleux, J., Lim, J. & Dunn, B. Pseudocapacitive contributions to electrochemical energy storage in TiO₂ (Anatase) nanoparticles. *J. Phys. Chem. C* **111**, 14925–14931 (2007).
44. Liu, W. & Speranza, G. Tuning the oxygen content of reduced graphene oxide and effects on its properties. *ACS Omega* **6**, 6195–6205 (2021).
45. Hillier, A. C. & Ward, M. D. Scanning electrochemical mass sensitivity mapping of the quartz crystal microbalance in liquid media. *Anal. Chem.* **64**, 2539–2554 (1992).
46. Varma, S. & Remppe, S. B. Coordination numbers of alkali metal ions in aqueous solutions. *Biophys. Chem.* **124**, 192–199 (2006).
47. Rao, J. S., Dinadayalane, T. C., Leszczynski, J. & Sastry, G. N. Comprehensive study on the solvation of mono- and divalent metal cations: Li⁺, Na⁺, K⁺, Be²⁺, Mg²⁺ and Ca²⁺. *J. Phys. Chem. A* **112**, 12944–12953 (2008).
48. Harris, D. J., Brodholt, J. P., Harding, J. H. & Sherman, D. M. Molecular dynamics simulation of aqueous ZnCl₂ solutions. *Mol. Phys.* **99**, 825–833 (2001).
49. Wang, C. et al. Toward flexible zinc-ion hybrid capacitors with superhigh energy density and ultralong cycling life: the pivotal role of ZnCl₂ salt-based electrolytes. *Angew. Chem. Int. Ed.* **60**, 990–997 (2021).
50. Cauet, E. et al. Structure and dynamics of the hydration shells of the Zn²⁺ ion from ab initio molecular dynamics and combined ab initio and classical molecular dynamics simulations. *J. Chem. Phys.* **132**, 194502 (2010).
51. Prehal, C. et al. Tracking the structural arrangement of ions in carbon supercapacitor nanopores using small-angle X-ray scattering. *Energy Environ. Sci.* **8**, 1725–1735 (2015).
52. Banda, H. et al. Ion sieving effects in chemically tuned pillared graphene materials for electrochemical capacitors. *Chem. Mater.* **30**, 3040–3047 (2018).
53. Escher, I., Hahn, M., Guillermo, A. F. & Adelhelm, P. A practical guide for using electrochemical dilatometry as operando tool in battery and supercapacitor research. *Energy Technol.* **10**, 2101120 (2022).
54. Hantel, M. M., Presser, V., Kötz, R. & Gogotsi, Y. In situ electrochemical dilatometry of carbide-derived carbons. *Electrochem. Commun.* **13**, 1221–1224 (2011).
55. Huesker, J. M., Winter, M. & Placke, T. Dilatometric study of the electrochemical intercalation of Bis(trifluoromethanesulfonyl) imide and hexafluorophosphate anions into carbon-based positive electrodes. *ECS Trans.* **69**, 9 (2015).
56. Huesker, J., Froböse, L., Kwade, A., Winter, M. & Placke, T. In situ dilatometric study of the binder influence on the electrochemical intercalation of Bis(trifluoromethanesulfonyl) imide anions into graphite. *Electrochim. Acta* **257**, 423–435 (2017).
57. Levi, M. D. et al. Solving the capacitive paradox of 2D MXene using electrochemical quartz-crystal admittance and in situ electronic conductance measurements. *Adv. Energy Mater.* **5**, 1400815 (2014).
58. Lv, W. et al. Low-temperature exfoliated graphenes: vacuum-promoted exfoliation and electrochemical energy storage. *ACS Nano* **3**, 3730–3736 (2009).
59. Shao, H., Lin, Z., Xu, K., Taberna, P.-L. & Simon, P. Electrochemical study of pseudocapacitive behavior of Ti₃C₂T_x MXene material in aqueous electrolytes. *Energy Storage Mater.* **18**, 456–461 (2019).

Acknowledgements

K.G. was supported by a grant from the China Scholarship Council. H.S. thanks the National Natural Science Foundation of China (Grant no. 22309202). P.S. and P.-L.T. are grateful for support from the European Research Council (ERC) and Réseau sur le Stockage Electrochimique de l’Energie (RS₂E) and the LABEX STOREX. This research was funded by ERC Synergy Grant MoMa-Stor #951513. For the purpose of open access, the author has applied a Creative Commons Attribution (CC BY) licence to any Author Accepted Manuscript version arising.

Author contributions

K.G., H.S., P.-L.T., and P.S. designed the research, P.-L.T., and P.S. supervised the project. K.G., and H.S. performed the material synthesis, electrochemical test, data analysis, and composed the manuscript. E.R.-P. carried out the TPD-MS measurement and performed data analysis. All authors discussed and interpreted the results and contributed to the writing of the manuscript.

Competing interests

The authors declare no competing interests.

Additional information

Supplementary information The online version contains supplementary material available at <https://doi.org/10.1038/s41467-024-46280-1>.

Correspondence and requests for materials should be addressed to Pierre-Louis Taberna or Patrice Simon.

Peer review information *Nature Communications* thanks Guohui Zhang and the other, anonymous, reviewer(s) for their contribution to the peer review of this work. A peer review file is available.

Reprints and permissions information is available at <http://www.nature.com/reprints>

Publisher’s note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if changes were made. The images or other third party material in this article are included in the article’s Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article’s Creative Commons licence and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit <http://creativecommons.org/licenses/by/4.0/>.

© The Author(s) 2024