ROYAL SOCIETY OF CHEMISTRY

COMMUNICATION

Check for updates

Interfacial charge transfer driven by surface termination-controlled Ti$_2$C MXene for enhanced hydrogen storage in magnesium†

Min Gyu Kim,[a] ShinYoung Kang,[b] Brandon C. Wood[b] and Eun Seon Cho [ID] *[a]

Two-dimensional transition metal carbides and nitrides (MXenes) with layered structure and high conductivity have been effectively utilized in various energy materials, including as a catalytic support of MgH$_2$ for hydrogen storage. However, the terminal groups formed on the surface during the etching step tend to deactivate reactive Mg metal and add dead mass, thereby deteriorating the catalytic role of MXene in hydrogen sorption of Mg. We exploited a molten-salt derived MXene with easily modifiable −Cl terminations, compared to conventional −O, −OH and −F groups, and synthesized a composite of Mg and delaminated Ti$_2$CCl$_x$ MXene to improve the hydrogen storage performance of Mg through charge transfer. This strategy enables the formation of an intimate interface between Mg and MXene, facilitating charge transfer and thereby boosting the catalytic effect. The resulting composite demonstrates significantly enhanced hydrogen sorption kinetics by modulating Mg−H bond strength. This novel approach of modifying surface terminations leverages the unique properties of MXene as a superior support for active materials, offering broader applications in energy materials.

1. Introduction

With growing apprehensions on the environmental impact of CO$_2$ emission, substituting traditional fossil fuels with renewable energy sources is imperative.[1,2] Hydrogen stands out as a promising energy carrier due to its high energy density, clean-burning feature and versatility, thereby enabling us to store a tremendous amount of energy generated from renewable sources. The efficient storage of hydrogen is a crucial aspect that impacts its viability as a widespread energy carrier.[3] Solid-state hydrogen storage offers considerable benefits compared to conventional pressurized gas or liquefied hydrogen storage in terms of safety and efficiency.[4] Particularly, magnesium hydride (MgH$_2$) has advantages of high theoretical gravimetric capacity (7.6 wt%), good reversibility and abundant natural resources,[5–7] while it faces major challenges such as high operating temperature requirements and sluggish reaction kinetics due to a strong Mg−H bonding nature.[8,9]

Introducing catalytic additives to MgH$_2$ such as transition metals[10–12] and their corresponding oxides[13,14] and carbides[15,16] is one of the feasible strategies to improve its hydrogen storage properties by providing additional activation sites or reaction pathways for hydrogen.[17] Recently, two-dimensional (2D) transition metal carbides and nitrides (MXenes) have been found to show a great catalytic activity for enhancing hydrogen sorption property of MgH$_2$.[18–21] Compared to typical transition metal-based catalysts which are sporadically doped within MgH$_2$ as clusters,[18] atomically thin 2D structured MXene offers the potential to effectively provide a number of active catalytic sites at the interface with MgH$_2$. Although previous studies have shown the validity of MXene as a catalyst for MgH$_2$, the underlying mechanism behind the catalytic effect is not adequately understood.[22] Particularly, studies on Ti-based MXene have suggested that the metallic Ti acts as a key catalytic component by forming an intermediate titanium hydride phase.[18,21,23] However, as metallic Ti is precipitated out of the carbide structure, the 2D structure and properties of a metal−carbon composite might undergo changes, causing it to lose its identity and ceasing its role as a distinct nanomaterial. Apart from the effect of precipitated Ti, the catalytic role of MXene as a metal-carbide has been investigated through computational simulations, yet corresponding experimental observations are currently lacking.[24–26]

On the other hand, during the MAX phase etching reaction for the synthesis of MXene, terminal groups such as =O, −OH, and −F are inevitably formed on the surface.[27,28] However, upon compositing with Mg for the aforementioned catalytic effect, Mg is susceptible to oxidation with these groups, resulting in a possible creation of passivation layers which hinder the reaction kinetics and reduce the active mass.[21] To avoid such

[a]Department of Chemical and Biomolecular Engineering, Korea Advanced Institute of Science and Technology (KAIST), 291 Daehak-ro, Yuseong-gu, Daejeon 34141, Republic of Korea. E-mail: escho@kaist.ac.kr; Tel: +82-42-350-3912

[b]Laboratory for Energy Application for the Future (LEAF), Lawrence Livermore National Laboratory, Livermore, California 94550, USA

† Electronic supplementary information (ESI) available: All experimental and theoretical details, and additional characterization experiments. See DOI: https://doi.org/10.1039/d4ta04563g

issues, a post-modification on surface termination of MXene is essential, while the robust Ti–O and Ti–F bonds introduced during the etching step in acid media make it challenging to replace surface groups through a post-synthetic route.[29] A newly developed synthetic method produces MXenes with –Cl and –Br terminations by using a molten salt, in place of oxygen or fluorine.[29–31] The Ti–Cl and Ti–Br bonds are relatively labile and can be substituted or eliminated to create a variety of MXene derivatives.[29,32] Despite such development, studies about elimination of surface termination from MXene remain limited in using LiH as a reducing agent assisted by molten-salt, which involves multiple steps and harsh reaction conditions.[29]

In this work, we present a facile synthesis of MXene with the removal of the –Cl terminal group (denoted as –Cl_x) via a molten-salt etching method for boosting the catalytic effect on hydrogen sorption kinetics of Mg upon the formation of a composite (Mg@DL-Ti_2CCl_x) with delaminated Ti_2CCl_2 MXene (DL-Ti_2CCl_2). The synthesized composite allows the formation of a Ti–Mg interface which has a catalytic effect on the hydrogen storage property of MgH_2. Interestingly, charge redistribution occurs between MgH_2 and DL-Ti_2CCl_x, resulting in a weakening of the strength of the Mg–H bond strength, which in turn improves hydrogen sorption properties. Additionally, DFT calculations also show that the strength of the interaction depends on the amount of –Cl present, with a lower –Cl content resulting in a stronger interaction. This implies that controlling the terminal groups of MXene is crucial for further enhancing its catalytic function as a support for MgH_2.

2. Results and discussion

The synthesis of Mg@DL-Ti_2CCl_x involves several key steps, as illustrated in Fig. 1a. Multilayer-Ti_2CCl_2 (ML-Ti_2CCl_2) was prepared from the MAX phase via a modified molten-salt etching method.[29,30] This was followed by delamination with lithium pyrene (LiPy) to form DL-Ti_2CCl_2, and simultaneous reduction of DL-Ti_2CCl_2 and the Mg^{2+} precursor was performed to form Mg@DL-Ti_2CCl_x (see ESI† for the detailed procedure). The scanning electron microscopy (SEM) image of ML-Ti_2CCl_2 in Fig. 1b shows a distinctive lamellar microstructure formed by the removal of the Al layer from the Ti_2AlC MAX phase and the decoration with –Cl terminal groups from the chloride salt. The SEM image (Fig. S1a†) of the Ti_2AlC MAX phase displays a combined structure of the Ti_2AlC and excess metal (Ti, Al) precursors,[30,33] which are eliminated during the etching step. Despite the absence of the Al layer, the ML-Ti_2CCl_2 MXene synthesized via the molten-salt etching method maintains a laminated structure due to van der Waals forces, posing challenges for good dispersion during solution-based synthesis and full utilization of its large surface area. Thus, a delamination process is employed, exposing the –Cl terminated basal plane on the surface before the composite synthesis. In previous studies, molten-salt etched MXenes were delaminated using n-butyllithium (n-buLi),[29,34] which has been widely used for transition-metal dichalcogenides via Li^+ ion intercalation.[35,36] However, a pyrophoric feature of n-buLi could pose a substantial safety hazard, especially when produced in a large scale for

hydrogen storage systems. In this work, we employed lithium pyrene (LiPy) for a safe and more effective delamination procedure, in which Li^+ ions are intercalated into ML-Ti_2CCl_2 layers owing to a lower redox potential, enabling successful delamination via simple ultrasonication.[37] The two-dimensional structure of the delaminated-Ti_2CCl_2 (DL-Ti_2CCl_2) is shown by transmission electron microscopy (TEM) and SEM images (Fig. 1c and S1b†). The DL-Ti_2CCl_2 forms a stable colloidal solution in a polar solvent like N-methyl formamide (Fig. S1c†), and its hexagonal crystallinity is retained after the delamination process as shown in the selected area electron diffraction (SAED) pattern (Fig. S2†).

To synthesize the Mg@DL-Ti_2CCl_x composite, a strong reducing agent – lithium biphenyl (LiBp) – was utilized in this work. In addition to reducing the Mg^{2+} precursor to Mg nanoparticles,[38,39] LiBp also partially eliminates the –Cl terminations from the DL-Ti_2CCl_2 surface, forming DL-Ti_2CCl_x, which will be discussed in the later section. This removal of –Cl functional groups exposes the titanium carbide surface, facilitating the formation of a direct Ti–Mg catalytic interface. In other words, the synthesis of Mg@DL-Ti_2CCl_x is accomplished through the removal of the –Cl terminal group from the DL-Ti_2CCl_2 layer along with a simultaneous reduction of Mg^{2+} precursors using LiBp. High-angle annular dark-field (HAADF) scanning transmission electron microscopy (STEM) with energy-dispersive X-ray spectroscopy (EDS) and bright-field TEM images (Fig. 1d and e) show the composite of Mg and DL-Ti_2CCl_x, demonstrating that Mg is homogeneously decorated on DL-Ti_2CCl_x over the Mg@DL-Ti_2CCl_x composite with residual –Cl termination (Fig. 1e).

To elucidate the effect of LiBp treatment on the elimination of –Cl termination, the MXenes – ML-Ti_2CCl_2 and DL-Ti_2CCl_2 – were solely treated with LiBp. The SEM EDS spectra show that the amount of elemental Cl over Ti_2CCl_2 layers is significantly reduced with the LiBp treatment to form ML-Ti_2CCl_x (Fig. S3 and Table S1†). In the subsequently obtained X-ray diffraction (XRD) pattern (Fig. S4a†), the ($11\bar{2}0$) peak whose position highly depends on the surface termination is shifted from 57.0° to 60.2°, implying the change of terminal groups.[29] In addition, the substantially reduced (0001) peak and the shifted ($11\bar{2}0$) peak to 60.2° are consistent with the previously reported result on bare-Ti_2C, in which case the terminal groups were found to be completely removed.[29] The reduction of the peak assigned to –Cl termination in the Raman spectra also supports that the LiBp treatment is effective for removal of –Cl termination (Fig. S4b†). The c lattice parameters were acquired from the high-resolution TEM (HRTEM) image for both ML-Ti_2CCl_2 and ML-Ti_2CCl_x, which are c = 8.76 Å and c = 10.46 Å, respectively (Fig. S4c and d†), closely aligned with the reported values in the literature.[29] The c lattice parameter of ML-Ti_2CCl_2 is consistent with the XRD result, as shown in Fig. S4a,† while the c value increases upon the LiBp-treatment because of space group change from $P\bar{3}m1$ to $P6_3/mmc$. The change in the Ti–C bond with the loss of the –Cl terminal group was investigated by the X-ray photoelectron spectroscopy (XPS) analysis in DL-Ti_2CCl_2 which was used during synthesis in delaminated form (Fig. S5†). The peaks for the Ti–C bond are shifted to lower binding energies in both the Ti 2p and C 1s regions as the electronegative element Cl
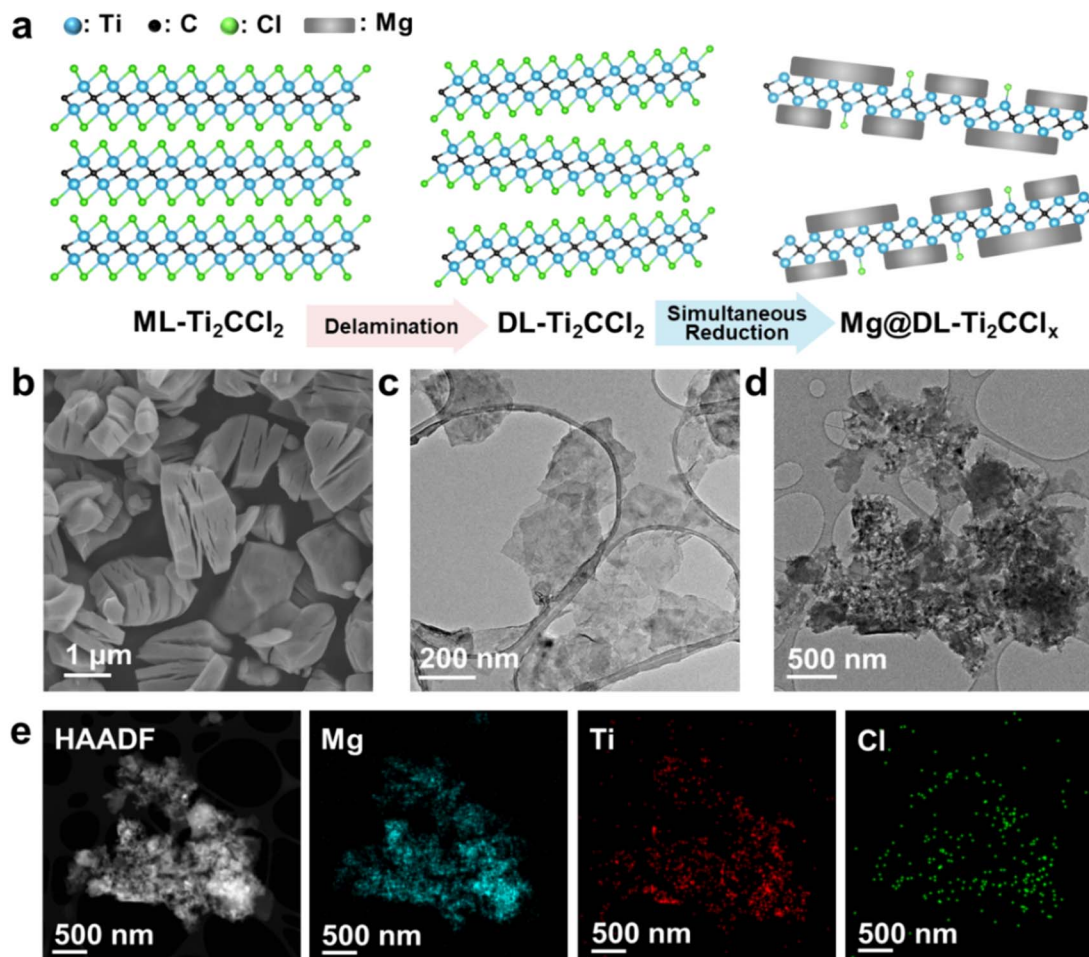
This journal is © The Royal Society of Chemistry 2024

J. Mater. Chem. A, 2024, **12**, 27212–27219 | **27213**

Fig. 1 Scheme and morphology change of the MXene and Mg composite by each synthetic step. (a) Schematic illustration of Mg@DL-Ti$_2$CCl$_x$ composite preparation. (b) Scanning electron microscopy (SEM) image of ML-Ti$_2$CCl$_2$. Transmission electron microscopy (TEM) image of (c) DL-Ti$_2$CCl$_2$ and (d) Mg@DL-Ti$_2$CCl$_x$. (e) High-angle annular dark-field (HAADF) image of Mg@DL-Ti$_2$CCl$_x$ and energy-dispersive X-ray spectroscopy (EDS) mapping of Mg, Ti and Cl representing evenly distributed Mg over DL-Ti$_2$CCl$_x$.

disappears. Additionally, Ti with a higher oxidation state is observed, attributed to the amorphous Ti oxide on the surface, which is reduced by LiBp-treatment. It is noteworthy that the conventional approaches to modifying functional groups over MXene require high temperatures (300–600 °C)[21,40] or molten-salt assisted reactions,[29] whereas LiBp readily removes functional groups under mild conditions. Even though a non-negligible amount of Cl is detected in the SEM-EDS analysis of ML-Ti$_2$CCl$_x$ (Fig. S3 and Table S1†) and TEM-EDS mapping of DL-Ti$_2$CCl$_x$ (Fig. S6†), the –Cl removal is quite evident and is thought to be sufficient to expose a bare Ti surface.

The change of the crystalline structure and terminal group over MXene was further investigated. The powder XRD pattern of ML-Ti$_2$CCl$_2$ shows distinct peaks corresponding to Ti$_2$CCl$_2$, indicating the complete etching of the Al layer from the Ti$_2$AlC MAX phase.[29,34] Prior to the delamination to form DL-Ti$_2$CCl$_2$, an expansion in interlayer spacing is observed in ML-Ti$_2$CCl$_2$ due to Li$^+$ intercalation, shifting the (0001) peak from $c = 8.67$ Å to $c = 8.88$ Å (Fig. S7†).[29] The DL-Ti$_2$CCl$_2$ shows a series of (000l) basal plane peaks since the delaminated sheets are horizontally aligned by vacuum-filtration.[29,41] The Mg@DL-Ti$_2$CCl$_x$ shows

a clear characteristic metallic Mg pattern with the (11$\bar{2}$0) peak of LiBp-treated Ti$_2$CCl$_2$, indicating the simultaneous partial removal of the –Cl terminal group. The formation of the composite was also confirmed by SAED, which shows a clear single-crystalline pattern from DL-Ti$_2$CCl$_x$ and the poly-crystalline ring from decorated Mg particles (Fig. S8†). The change of surface termination on MXene was also identified through Raman spectroscopy (Fig. 2b). The out-of-plane vibration of Ti–C with –Cl termination is observed at 225 cm^{-1} (A$_{1g}$) for ML-Ti$_2$CCl$_2$.[34] DL-Ti$_2$CCl$_2$ retains the A$_{1g}$ peak at 225 cm^{-1} which indicates that –Cl termination is well preserved during the delamination process, while Mg@DL-Ti$_2$CCl$_x$ shows a peak at 341 cm^{-1} (A$_{1g}$) which corresponds to the out-of-plane stretching mode of Ti–C without surface termination,[42] and the A$_{1g}$ peak associated with the –Cl termination at 225 cm^{-1} is significantly diminished. These results suggest that most of the –Cl terminal groups are removed under the highly reductive environment with LiBp, thereby allowing the direct interaction of Mg with Ti atoms in MXene.

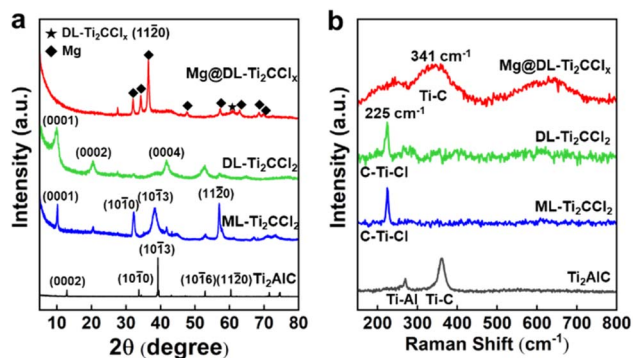In order to examine the hydrogen sorption properties of Mg@DL-Ti$_2$CCl$_x$, isothermal ab/desorption kinetics was

27214 | *J. Mater. Chem. A*, 2024, **12**, 27212–27219

This journal is © The Royal Society of Chemistry 2024

Fig. 2 Change of the crystalline and chemical structures of the Ti_2AlC MAX phase, $ML\text{-}Ti_2CCl_2$, $DL\text{-}Ti_2CCl_2$ and $Mg@DL\text{-}Ti_2CCl_x$ composite based on (a) X-ray diffraction (XRD) patterns and (b) Raman spectra.

measured with Sievert's apparatus (Fig. 3a and b). In the first cycle, 4.1 wt% of H_2 is absorbed within 20 min; however, the kinetics gets sluggish in the subsequent cycle, while its rate is maintained afterwards. On the other hand, the desorption is completed within 20 minutes and maintains a stable rate. To understand these phenomena, we consider the hydrogen sorption process of Mg metal. The H_2 absorption follows several sequential steps: (1) dissociation of hydrogen molecule on the surface, (2) diffusion of hydrogen atom into metal and (3) nucleation and growth of hydride phase. Reversely, desorption from MgH_2 involves: (1) formation of metallic Mg phase, (2) diffusion of hydrogen atom from hydride phase to metallic Mg phase and (3) recombination of hydrogen atoms to form hydrogen molecule.[43,44] It can be assumed that the absorption rate decreases since Mg particles are sintered during the H_2 ab/desorption cycle.[45,46] Since MgH_2 melts at a lower temperature (327 °C) compared to Mg (650 °C), sintering is likely to occur in the hydrogenated state. Increased Mg grain size reduces the surface area for hydrogen dissociation and slows hydrogen atom diffusion to the grain core.[47,48] The grain size growth is

Fig. 3 Hydrogen sorption kinetics of the $Mg@DL\text{-}Ti_2CCl_x$ composite. (a) Isothermal hydrogen absorption at 200 °C under 15 bar of H_2 and (b) hydrogen desorption at 300 °C under 0 bar of H_2. Activation barrier for (c) absorption and (d) desorption as a function of absorbed and desorbed wt% H_2, respectively.

also evident in the XRD result (Fig. S9a†), in which the initially broad Mg peaks for as-synthesized $Mg@DL\text{-}Ti_2CCl_x$ become narrow after H_2 sorption cycles, yet the characteristic $(11\bar{2}0)$ MXene peak is preserved which shows that $DL\text{-}Ti_2CCl_x$ retains its structure (Fig. S9b†). The sintering primarily occurs during the initial cycle, possibly because the $DL\text{-}Ti_2CCl_x$ within the composite acts as a physical barrier, preventing further excessive sintering once it reaches a certain equilibrium state. The structural stability is confirmed by preservation of the hexagonal crystallinity of $DL\text{-}Ti_2CCl_x$ in the composite after the hydrogen sorption cycle, as shown in the SAED pattern (Fig. S10†).

Nevertheless, the desorption rate of $Mg@DL\text{-}Ti_2CCl_x$ is not significantly affected as the hydrogen cycle proceeds, compared to the absorption (Fig. 3b). It is assumed that this unusual discrepancy arises from the different reaction mechanism governing the hydrogen ab/desorption process. Typically, MgH_2 undergoes a localized charge transfer step between Mg^{2+} and H^- during desorption.[49,50] However, due to the lack of free electrons in the conduction band of semi-conductive MgH_2 at the fully hydrogenated state, the nucleation of metallic Mg should be preceded which is commonly referred to as an incubation period.[50–52] Interestingly, the desorption of $Mg@DL\text{-}Ti_2CCl_x$ shows negligible incubation time at the initial desorption stage. We speculate that the high electrical conductivity of $DL\text{-}Ti_2CCl_x$, especially the –Cl removed region,[53] allows it to act as a charge transfer intermediate, facilitating hydrogen recombination without a prior incubation period. For comparison, we additionally synthesized the composite of Mg and conventional Ti_2CT_x (T = –O, –OH and –F) MXene instead of $DL\text{-}Ti_2CCl_2$ ($Mg@Ti_2CT_x$). In the isothermal hydrogen ab/desorption of $Mg@Ti_2CT_x$, it is observed that the rate gradually decreases for both cases, along with a typical incubation period at the initial stage of desorption, as opposed to $Mg@DL\text{-}Ti_2CCl_x$ (Fig. S11†). This suggests that the catalytic activity of Ti_2CT_x deteriorates as the hydrogen cycle progresses, which implies the collapse of the Ti_2CT_x MXene structure. MXene synthesized with an acid solution contains F^- ions (LiF/HCl or HF), and it might become structurally unstable because of the formation of atomic defects.[27,54] Hence, unlike $DL\text{-}Ti_2CCl_x$, Ti_2CT_x might be more vulnerable under a hydrogen cycle environment, potentially leading to the loss of its catalytic function and its role as a physical barrier. Furthermore, it was reported that MXene with a terminal group could cause the formation of a passivation layer at the interface of Mg,[21] suggesting that the elimination of terminal groups is crucial when creating the composite of Mg and MXene. Despite using LiBp as a reducing agent, the removal of terminal groups from conventional Ti_2CT_x is difficult to achieve, supported by SEM-EDS analysis and Raman spectroscopy (Fig. S12 and Table S2†). Unlike $ML\text{-}Ti_2CCl_2$, the terminal groups mostly remain, and both the out-of-plane vibration (A_{1g}) and in-plane vibration of Ti–C (E_g) with terminal groups are observed. Presumably, harsher conditions are required to break the terminal groups attached to Ti_2CT_x. The key factor for creating an efficient catalytic interface lies in the utilization of –Cl decorated MXene and its subsequent removal through LiBp treatment, as demonstrated in this study.
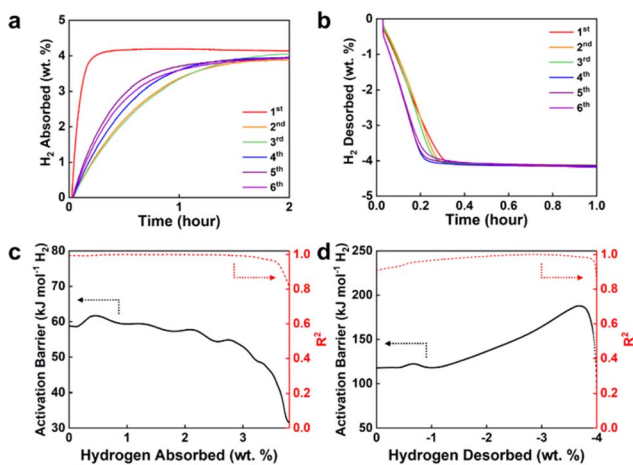
This journal is © The Royal Society of Chemistry 2024

J. Mater. Chem. A, 2024, **12**, 27212–27219 | **27215**

In order to further probe the hydrogen sorption kinetics of Mg@DL-Ti$_2$CCl$_x$, we evaluated activation energies by fitting hydrogen ab/desorption kinetic curves at three different temperatures (Fig. S13†) to the Arrhenius law (Fig. 3c and d) and Johnson–Mehl–Avrami (JMA) equation (Fig. S14 and S15†). The hydrogen ab/desorption kinetic curves and activation barrier plots solely based on the mass of Mg are additionally shown in Fig. S16,† while the overall trend tends to be consistent with the one based on the total composite mass. During absorption, Mg@DL-Ti$_2$CCl$_x$ displays lower activation energy values than the barriers identified in previous studies on hydrogen dissociation on the Mg surface ($E_a \approx 83.9–101.3$ kJ mol^{-1} H$_2$)[55–57] and H atom diffusion through MgH$_2$ ($E_a \approx 95.5$ kJ mol^{-1} H$_2$).[58,59] Thus, we anticipate that the energy barriers for both surface and bulk processes are alleviated in the hydrogen absorption kinetics of Mg@DL-Ti$_2$CCl$_x$. The activation energy acquired from the JMA equation ($E_a \approx 53.5$ kJ mol^{-1} H$_2$) is also aligned with it, while providing additional information regarding the dimensionality of the growth. The best fitting was obtained with an Avrami exponent (n) of $n = 1.0$, indicating that the growth of MgH$_2$ occurs in one-dimensional geometry which can be described by hydrogen diffusion through lattice vacancies followed by thickening of the MgH$_2$ layer in one-dimension as proposed in the literature.[60,61] Furthermore, the activation energy desorption for Mg@DL-Ti$_2$CCl$_x$ determined from both the Arrhenius law (Fig. 3d) and JMA equation (Fig. S15d†) is found to be lower than that for other Mg composites ($E_a \approx 140–219$ kJ mol^{-1} H$_2$)[59,62–64] which are synthesized through a similar Rieke method.[65] This improvement on desorption kinetics is believed to originate from the weakening of the Mg–H bond caused by charge transfer which will be discussed later. A notable characteristic is that the n value of desorption kinetics from the JMA fitting displays a non-integer value ($n = 0.5–0.7$). This suggests that the desorption process may occur *via* diffusion-controlled one-dimensional growth with instantaneous nucleation which aligns with a negligible incubation period as described previously.[66]

To unravel the in-depth catalytic effect of DL-Ti$_2$CCl$_x$ on the hydrogen storage performance of Mg@DL-Ti$_2$CCl$_x$ at the interface of Ti–Mg, the change of the chemical state before and after H$_2$ sorption cycles was investigated through XPS analysis (Fig. 4a and b). As-synthesized Mg@DL-Ti$_2$CCl$_x$ exhibits a comparable binding energy for the Ti–C bond to DL-Ti$_2$CCl$_x$ in both Ti 2p and C 1s spectra (Fig. S5a and b†); however, the relevant peaks are shifted to the lower binding energy after hydrogen sorption cycling, which clearly shows the charge redistribution in the Mg@DL-Ti$_2$CCl$_x$ composite which was induced during the cycling with the existence of Mg. The consistent shift of the Ti–C peak to a lower binding energy in the Ti 2p and C 1s XPS results indicates that charge transfer from Mg to Ti–C has occurred. The retention of the transition metal related peak in the C 1s spectra after the hydrogen sorption process indicates that the Ti–C bonding is well preserved. This aligns with the earlier explanation of structural stability. In addition, the hexagonal crystallinity of DL-Ti$_2$CCl$_x$ in the composite is well maintained after the hydrogen sorption cycle (Fig. S10†). This structural stability is attributed to the
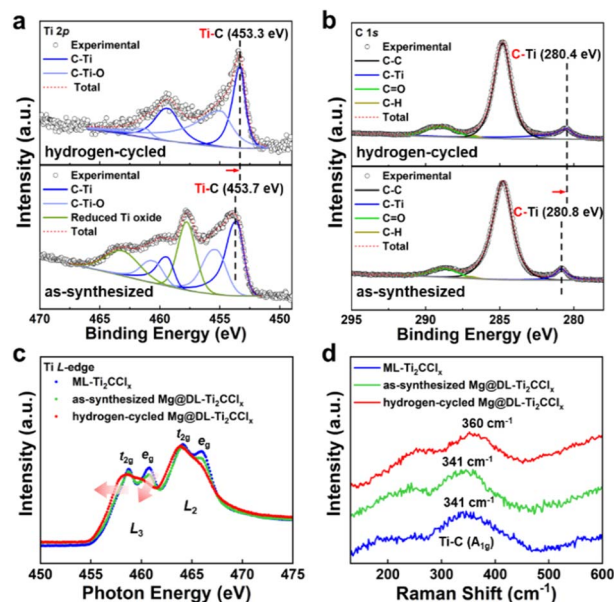

Fig. 4 Charge transfer between DL-Ti$_2$CCl$_x$ and Mg. (a) Ti 2p and (b) C 1s XPS spectra of as-synthesized Mg@DL-Ti$_2$CCl$_x$ (below) and hydrogen-cycled Mg@DL-Ti$_2$CCl$_x$ (upper). (c) Ti L-edge NEXAFS and (d) Raman spectra of ML-Ti$_2$CCl$_x$, as-synthesized and hydrogen-cycled Mg@DL-Ti$_2$CCl$_x$.

molten-salt etching method, wherein hydrofluoric acid (HF) was intentionally excluded from the synthesis process. To the best of our knowledge, our study is the first experimental demonstration of this distinct charge transfer between Mg and Ti–C in the composite of Mg and MXene. This charge transfer effect was further supported by Ti L-edge near edge X-ray absorption fine structure (NEXAFS) and Raman spectroscopy (Fig. 4c and d).[67,68] The Ti L-edge spectrum is related to electron excitation from Ti 2p$_{3/2}$ (L$_3$) and 2p$_{1/2}$ (L$_2$) core levels to 3d states. As the 3d orbital energy level is split by the crystal field effect, each Ti L$_3$ edge and L$_2$ edge is further divided into t$_{2g}$ and e$_g$ subpeaks.[69] In the hydrogen cycled sample, the t$_{2g}$ peaks become more intense at the both L$_2$ and L$_3$ edges, accompanied by the loss of the clear splitting and a shift of the pre-edge towards the lower energy, indicating a lower oxidation state of Ti in MXene.[70–72] The observation of a blue shift (341 cm^{-1} to 360 cm^{-1}) in the A$_{1g}$ peak, associated with the Ti–C stretching mode in the Raman spectra of the hydrogen cycled sample (Fig. 4d), suggests a strengthening of the Ti–C bond due to the charge transfer.[67] These results confirm the charge transfer between DL-Ti$_2$CCl$_x$ and Mg, which suggests that DL-Ti$_2$CCl$_x$ may perturb the Mg–H bonds for the dehydrogenation of MgH$_2$.

To further elucidate the impact of DL-Ti$_2$CCl$_x$ on the MgH$_2$ desorption kinetics, we conducted density functional theory (DFT) calculations,[73,74] in which structures and charge densities were analyzed using the interface models between MgH$_2$ nanoparticle and DL-Ti$_2$CCl$_x$ sheet ($x = 0.0, 0.5, 1.0,$ and 1.5) that are visualized in Fig. S17.† The Bader change analysis calculates the number of electrons belonging to each atom by defining atomic volume within which the charge density is integrated.[75–78] We also computed the Bader charge state of
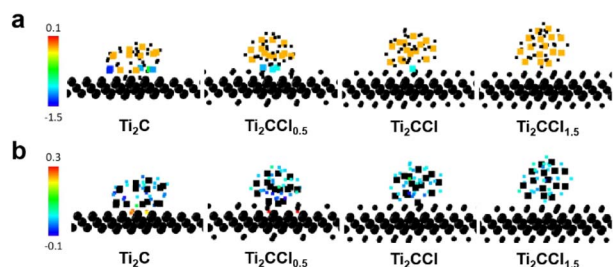
27216 | *J. Mater. Chem. A*, 2024, **12**, 27212–27219

This journal is © The Royal Society of Chemistry 2024

Fig. 5 Changes in the Bader charge of (a) Mg atoms and (b) H atoms in the MgH$_2$@DL-Ti$_2$CCl$_x$ interface models evaluated based on the DFT calculations. Mg and H atoms are denoted by large and small squares, respectively. Circles are Ti, C, and Cl atoms in MXene sheets. Atoms of interest (Mg in (a) and H in (b)) are colored by their valence state change: reddish color indicates electron loss (oxidation), and bluish color indicates electron gain (reduction). Other atoms are colored in black for simplicity.

isolated MgH$_2$ nanoparticles free from MXene as a reference. Fig. 5 displays the Bader charge difference in Mg and H atoms before and after forming the interface, *i.e.*, changes in the number of electrons when forming the MgH$_2$@Ti$_2$CCl$_x$ interface compared to the isolated MgH$_2$ nanoparticles. For Mg, centering from the orange color where the atomic charge is invariant, positive change (reddish color) and negative change (bluish color) indicate oxidation and reduction, respectively. The impact of Ti$_2$CCl$_x$ is most prominent when the –Cl terminal group is completely removed. Many interface Mg atoms obtain additional electrons, which is from Mg^{2+} in MgH$_2$ towards the fully desorbed Mg^0 metal. Similarly, H$^-$ ions in MgH$_2$ oxidize towards H^0 near Ti$_2$C, which can easily recombine into H$_2$ molecule. The charge redistribution phenomenon due to Ti$_2$CCl$_x$ slowly diminishes in the presence of the –Cl terminations, and eventually the Mg and H atoms do not feel the MgH$_2$@Ti$_2$CCl$_x$ interface at $x = 1.5$. Besides the charge redistribution within the MgH$_2$ particle, Ti$_2$CCl$_x$ also participates in the charge transfer (Fig. S18†). However, given that atoms in Ti$_2$CCl$_x$ undergo both oxidation and reduction due to the complexity in the local environment, it is not clear that the role of Ti$_2$CCl$_x$ is monolithic in charge transfer. More systematic investigation exploring various contact area, distance, and local environment at the MgH$_2$@Ti$_2$CCl$_x$ interface is demanded for clarification, which is beyond the scope of this work. Nonetheless, the impact of charge redistribution within MgH$_2$ is clear due to the deformation of its structure. Reduction of Mg and oxidation of H elongate the Mg–H bond from 1.88 Å to 1.91 Å in the MgH$_2$@-Ti$_2$C (Fig. S19†), suggesting that Ti$_2$CCl$_x$ facilitates the initial step of desorption, namely Mg–H bond breakage, and it eventually leads to improved desorption rates as observed in experiments. Our simulation also supports the efficacy of LiBp treatment which helps eliminating the –Cl terminations to maximize the catalytic activity of Ti$_2$CCl$_x$.

3.   Conclusions

In summary, this study demonstrates the role of DL-Ti$_2$CCl$_x$ MXene in enhancing the hydrogen storage property of Mg *via* the formation of an intimate interface with Ti-exposed DL-Ti$_2$CCl$_x$. Such interface allows efficient charge transfer between Mg and Ti, attributed to the removal of surface termination groups over MXene, and it is enabled by introducing relatively weak –Cl terminations to the Ti atoms using the molten-salt etching method, in contrast to the previously reported approaches which require harsh conditions. An identical reduction process was employed for incorporating Mg particles, leading to the one-pot synthesis of the Mg@DL-Ti$_2$CCl$_x$ composite. The composite exhibits improved hydrogen desorption property of MgH$_2$, primarily due to the charge transfer from Mg to DL-Ti$_2$CCl$_x$, which reduces the strength of the Mg–H bond at the interface. Furthermore, due to the excellent electrical conductivity of Ti$_2$CCl$_x$, it potentially serves as an electron mediator, facilitating electron transfer at the initial stage of desorption and thus reducing the incubation period. This work provides a novel approach for synthesizing Mg–MXene complexes for efficient hydrogen storage medium and elucidates the charge transfer at the Ti–Mg interface. This strategy can be applied to other energy materials, such as those used in energy storage and catalysis, to improve their performance through the formation of a well-defined interface and electronic structure control.

Data availability

The authors confirm that the data underlying this article are available within the article and its ESI†.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

Notes and references

1 N. L. Panwar, S. C. Kaushik and S. Kothari, *Renewable Sustainable Energy Rev.*, 2011, **15**, 1513–1524.
2 J. Rockström, O. Gaffney, J. Rogelj, M. Meinshausen, N. Nakicenovic and H. J. Schellnhuber, *Science*, 2017, **355**, 1269–1271.

This journal is © The Royal Society of Chemistry 2024

J. Mater. Chem. A, 2024, **12**, 27212–27219 | **27217**

3 S. Niaz, T. Manzoor and A. H. Pandith, *Renewable Sustainable Energy Rev.*, 2015, **50**, 457–469.

4 S. McQueen, J. Stanford, S. Satyapal, E. Miller, N. Stetson, D. Papageorgopoulos, N. Rustagi, V. Arjona, J. Adams and K. Randolph, *Department of Energy Hydrogen Program Plan*, US Department of Energy (USDOE), Washington DC United States, 2020.

5 Y. Wang and Y. Wang, *Prog. Nat. Sci.: Mater. Int.*, 2017, **27**, 41–49.

6 T. Sadhasivam, H.-T. Kim, S. Jung, S.-H. Roh, J.-H. Park and H.-Y. Jung, *Renewable Sustainable Energy Rev.*, 2017, **72**, 523–534.

7 D. J. Han, K. R. Bang, H. Cho and E. S. Cho, *Korean J. Chem. Eng.*, 2020, **37**, 1306–1316.

8 Y. Sun, C. Shen, Q. Lai, W. Liu, D.-W. Wang and K.-F. Aguey-Zinsou, *Energy Storage Mater.*, 2018, **10**, 168–198.

9 H. Wang, H. Lin, W. Cai, L. Ouyang and M. Zhu, *J. Alloys Compd.*, 2016, **658**, 280–300.

10 G.-x. Liang, J. Huot, S. Boily, A. Van Neste and R. Schulz, *J. Alloys Compd.*, 1999, **292**, 247–252.

11 C. Zhou, J. Zhang, R. C. Bowman Jr and Z. Z. Fang, *Inorganics*, 2021, **9**, 36.

12 J. Cui, J. Liu, H. Wang, L. Ouyang, D. Sun, M. Zhu and X. Yao, *J. Mater. Chem. A*, 2014, **2**, 9645–9655.

13 L. Dan, H. Wang, X. Yang, J. Liu, L. Ouyang and M. Zhu, *ACS Appl. Mater. Interfaces*, 2023, **15**, 30372–30382.

14 M. Song, J.-L. Bobet and B. Darriet, *J. Alloys Compd.*, 2002, **340**, 256–262.

15 Z. Wang, X. Zhang, Z. Ren, Y. Liu, J. Hu, H. Li, M. Gao, H. Pan and Y. Liu, *J. Mater. Chem. A*, 2019, **7**, 14244–14252.

16 M. Tian and C. Shang, *Int. J. Hydrogen Energy*, 2019, **44**, 338–344.

17 X. Zeng, L. Cheng, J. Zou, W. Ding, H. Tian and C. Buckley, *J. Appl. Phys.*, 2012, **111**, 093720.

18 Y. Liu, H. Du, X. Zhang, Y. Yang, M. Gao and H. Pan, *Chem. Commun.*, 2016, **52**, 705–708.

19 W. Zhu, S. Panda, C. Lu, Z. Ma, D. Khan, J. Dong, F. Sun, H. Xu, Q. Zhang and J. Zou, *ACS Appl. Mater. Interfaces*, 2020, **12**, 50333–50343.

20 H. Liu, C. Lu, X. Wang, L. Xu, X. Huang, X. Wang, H. Ning, Z. Lan and J. Guo, *ACS Appl. Mater. Interfaces*, 2021, **13**, 13235–13247.

21 W. Zhu, L. Ren, C. Lu, H. Xu, F. Sun, Z. Ma and J. Zou, *ACS Nano*, 2021, **15**, 18494–18504.

22 J. A. Bolarin, R. Zou, Z. Li, Z. Zhang and H. Cao, *Appl. Mater. Today*, 2022, **29**, 101570.

23 W. Ali, X. Li, Y. Yang, N. Li, B. Huang, C. Wu and S. Ding, *ACS Appl. Mater. Interfaces*, 2023, **15**, 36167–36178.

24 Z. Huang, Y. Wang and M. Zhang, *Int. J. Hydrogen Energy*, 2021, **46**, 33176–33185.

25 C. Lu, H. Liu, L. Xu, H. Luo, S. He, X. Duan, X. Huang, X. Wang, Z. Lan and J. Guo, *J. Magnesium Alloys*, 2022, **10**, 1051–1065.

26 H. Gao, R. Shi, Y. Liu, Y. Zhu, J. Zhang, L. Li and X. Hu, *J. Magnesium Alloys*, 2022, **11**, 3724–3735.

27 K. R. G. Lim, M. Shekhirev, B. C. Wyatt, B. Anasori, Y. Gogotsi and Z. W. Seh, *Nat. Synth.*, 2022, **1**, 601–614.

28 A. VahidMohammadi, J. Rosen and Y. Gogotsi, *Science*, 2021, **372**, eabf1581.

29 V. Kamysbayev, A. S. Filatov, H. Hu, X. Rui, F. Lagunas, D. Wang, R. F. Klie and D. V. Talapin, *Science*, 2020, **369**, 979–983.

30 M. Li, J. Lu, K. Luo, Y. Li, K. Chang, K. Chen, J. Zhou, J. Rosen, L. Hultman and P. Eklund, *J. Am. Chem. Soc.*, 2019, **141**, 4730–4737.

31 Y. Li, H. Shao, Z. Lin, J. Lu, L. Liu, B. Duployer, P. O. Persson, P. Eklund, L. Hultman and M. Li, *Nat. Mater.*, 2020, **19**, 894–899.

32 C. Zhou, D. Wang, F. Lagunas, B. Atterberry, M. Lei, H. Hu, Z. Zhou, A. S. Filatov, D.-e. Jiang, A. J. Rossini, R. F. Klie and D. V. Talapin, *Nat. Chem.*, 2023, **15**, 1722–1729.

33 W. Zhou, B. Mei, J. Zhu and X. Hong, *Mater. Lett.*, 2005, **59**, 131–134.

34 D. Wang, C. Zhou, A. S. Filatov, W. Cho, F. Lagunas, M. Wang, S. Vaikuntanathan, C. Liu, R. F. Klie and D. V. Talapin, *Science*, 2023, **379**, 1242–1247.

35 P. Joensen, R. Frindt and S. R. Morrison, *Mater. Res. Bull.*, 1986, **21**, 457–461.

36 A. Ambrosi, Z. Sofer and M. Pumera, *Small*, 2015, **11**, 605–612.

37 X. Zhu, Z. Su, C. Wu, H. Cong, X. Ai, H. Yang and J. Qian, *Nano Lett.*, 2022, **22**, 2956–2963.

38 N. S. Norberg, T. S. Arthur, S. J. Fredrick and A. L. Prieto, *J. Am. Chem. Soc.*, 2011, **133**, 10679–10681.

39 V. Lomonosov, E. R. Hopper and E. Ringe, *Chem. Commun.*, 2023, **59**, 5603–5606.

40 I. Persson, J. Halim, H. Lind, T. W. Hansen, J. B. Wagner, L. Å. Näslund, V. Darakchieva, J. Palisaitis, J. Rosen and P. O. Persson, *Adv. Mater.*, 2019, **31**, 1805472.

41 M. Alhabeb, K. Maleski, B. Anasori, P. Lelyukh, L. Clark, S. Sin and Y. Gogotsi, *Chem. Mater.*, 2017, **29**, 7633–7644.

42 T. Hu, M. Hu, Z. Li, H. Zhang, C. Zhang, J. Wang and X. Wang, *J. Phys. Chem. A*, 2015, **119**, 12977–12984.

43 J. Fernandez and C. Sanchez, *J. Alloys Compd.*, 2002, **340**, 189–198.

44 C. Zhou, Z. Z. Fang, R. C. Bowman Jr, Y. Xia, J. Lu, X. Luo and Y. Ren, *J. Phys. Chem. C*, 2015, **119**, 22272–22280.

45 P. E. De Jongh, *Nat. Mater.*, 2011, **10**, 265–266.

46 M. Dornheim, N. Eigen, G. Barkhordarian, T. Klassen and R. Bormann, *Adv. Eng. Mater.*, 2006, **8**, 377–385.

47 X. Yao, Z. Zhu, H. Cheng and G. Lu, *J. Mater. Res.*, 2008, **23**, 336–340.

48 C. Zhou, Z. Z. Fang and R. C. Bowman Jr, *J. Phys. Chem. C*, 2015, **119**, 22261–22271.

49 S. Kang, T. Ogitsu, S. A. Bonev, T. W. Heo, M. D. Allendorf and B. C. Wood, *ECS Trans.*, 2017, **77**, 81.

50 I. Gabis, M. Dobrotvorskiy, E. Evard and A. Voyt, *J. Alloys Compd.*, 2011, **509**, S671–S674.

51 N. Takeichi, Y. Sakaida, T. Kiyobayashi and H. T. Takeshita, *Mater. Trans.*, 2014, **55**, 1161–1167.

52 A. S. Gangrade, A. A. Varma, N. K. Gor, S. Shriniwasan and S. S. V. Tatiparti, *Phys. Chem. Chem. Phys.*, 2017, **19**, 6677–6687.

53 M. Khazaei, A. Ranjbar, M. Arai, T. Sasaki and S. Yunoki, *J. Mater. Chem. C*, 2017, **5**, 2488–2503.

54 X. Sang, Y. Xie, M.-W. Lin, M. Alhabeb, K. L. Van Aken, Y. Gogotsi, P. R. Kent, K. Xiao and R. R. Unocic, *ACS Nano*, 2016, **10**, 9193–9200.

55 M. Pozzo and D. Alfe, *Int. J. Hydrogen Energy*, 2009, **34**, 1922–1930.

56 T. Vegge, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2004, **70**, 035412.

57 A. J. Du, S. C. Smith, X. D. Yao and G. Q. Lu, *J. Phys. Chem. B*, 2006, **110**, 21747–21750.

58 J. Čermák and L. Král, *Acta Mater.*, 2008, **56**, 2677–2686.

59 L. F. Wan, E. S. Cho, T. Marangoni, P. Shea, S. Kang, C. Rogers, E. Zaia, R. R. Cloke, B. C. Wood and F. R. Fischer, *Chem. Mater.*, 2019, **31**, 2960–2970.

60 K.-J. Jeon, H. R. Moon, A. M. Ruminski, B. Jiang, C. Kisielowski, R. Bardhan and J. J. Urban, *Nat. Mater.*, 2011, **10**, 286–290.

61 E. S. Cho, A. M. Ruminski, S. Aloni, Y.-S. Liu, J. Guo and J. J. Urban, *Nat. Commun.*, 2016, **7**, 10804.

62 D. J. Han, S. Kim and E. S. Cho, *J. Mater. Chem. A*, 2021, **9**, 9875–9881.

63 H. Cho, S. Hyeon, H. Park, J. Kim and E. S. Cho, *ACS Appl. Energy Mater.*, 2020, **3**, 8143–8149.

64 Y. Cho, S. Kang, B. C. Wood and E. S. Cho, *ACS Appl. Mater. Interfaces*, 2022, **14**, 20823–20834.

65 R. D. Rieke, *Science*, 1989, **246**, 1260–1264.

66 K. Shirzad and C. Viney, *J. R. Soc., Interface*, 2023, **20**, 20230242.

67 Q.-Q. Yan, P. Yin and H.-W. Liang, *ACS Mater. Lett.*, 2021, **3**, 1197–1212.

68 I. C. Gerber and P. Serp, *Chem. Rev.*, 2019, **120**, 1250–1349.

69 F. De Groot, J. Fuggle, B. Thole and G. Sawatzky, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1990, **41**, 928.

70 A. Al-Temimy, K. Prenger, R. Golnak, M. Lounasvuori, M. Naguib and T. Petit, *ACS Appl. Mater. Interfaces*, 2020, **12**, 15087–15094.

71 K. Prenger, Y. Sun, K. Ganeshan, A. Al-Temimy, K. Liang, C. Dun, J. J. Urban, J. Xiao, T. Petit and A. C. Van Duin, *ACS Appl. Energy Mater.*, 2022, **5**, 9373–9382.

72 A. Al-Temimy, B. Anasori, K. A. Mazzio, F. Kronast, M. Seredych, N. Kurra, M.-A. Mawass, S. Raoux, Y. Gogotsi and T. Petit, *J. Phys. Chem. C*, 2020, **124**, 5079–5086.

73 P. Hohenberg and W. Kohn, *Phys. Rev.*, 1964, **136**, B864.

74 W. Kohn and L. J. Sham, *Phys. Rev.*, 1965, **140**, A1133.

75 W. Tang, E. Sanville and G. Henkelman, *J. Phys.: Condens. Matter*, 2009, **21**, 084204.

76 E. Sanville, S. D. Kenny, R. Smith and G. Henkelman, *J. Comput. Chem.*, 2007, **28**, 899–908.

77 G. Henkelman, A. Arnaldsson and H. Jónsson, *Comput. Mater. Sci.*, 2006, **36**, 354–360.

78 M. Yu and D. R. Trinkle, *J. Chem. Phys.*, 2011, **134**, 064111.

This journal is © The Royal Society of Chemistry 2024

J. Mater. Chem. A, 2024, **12**, 27212–27219 | **27219**