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A general Lewis acidic etching route for preparing MXenes with enhanced electrochemical performance in non-aqueous electrolyte

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Two-dimensional carbides and nitrides of transition metals, known as MXenes, are a fast-growing family of materials that have attracted attention as energy storage materials. MXenes are mainly prepared from Al-containing MAX phases (where A = Al) by Al dissolution in F-containing solution; most other MAX phases have not been explored. Here a redox-controlled A-site etching of MAX phases in Lewis acidic melts is proposed and validated by the synthesis of various MXenes from unconventional MAX-phase precursors with A elements Si, Zn and Ga. A negative electrode of Ti₃C₂ MXene material obtained through this molten salt synthesis method delivers a Li⁺ storage capacity of up to 738 C g⁻¹ (205 mAh g⁻¹) with high charge-discharge rate and a pseudocapacitive-like electrochemical signature in 1 M LiPF₆ carbonate-based electrolyte. MXenes prepared via this molten salt synthesis route may prove suitable for use as high-rate negative-electrode materials for electrochemical energy storage applications.

Two-dimensional (2D) transition metal carbides or carbonitrides (MXenes) are one of the latest additions to the family of 2D materials. MXenes are prepared by selective etching of the A-layer elements in MAX-phase precursors, where M represents an early transition metal element (Ti, V, Nb and so on), A is an element mainly from groups 13–16 (Al, Si and so on) and X is carbon and/or nitrogen¹. Their general formula can be written as M_{n+1}X_nT_x (n = 1–3), where T_x stands for the surface terminations, generally considered to be -F, -O or -OH. Due to their unique 2D layered structure, hydrophilic surfaces and metallic conductivity (>6,000 S cm⁻¹), MXenes show promise in a broad range of applications, notably electrochemical energy storage^{2,3}.

Following the first report of Ti₃C₂ MXene synthesis in 2011, MXenes are currently mainly prepared by selective etching of the A-layer in MAX phases by aqueous solutions containing fluoride ions, such as aqueous HF (ref. ¹), LiF + HCl mixtures⁴ or ammonium bifluoride ((NH₄)HF₂) (ref. ⁵). To date, the high reactivity of Al with fluoride-based aqueous solutions has limited the synthesis of MXenes mainly to those derived from Al-containing MAX-phase precursors. Although Alhabet et al. reported the synthesis of Ti₃C₂ MXene through oxidant-assisted etching of Si from Ti₃SiC₂ MAX phase⁵, the etching mechanism was still based on the use of hazardous HF solution. The main challenges for MXene synthesis are therefore (1) to find non-hazardous synthesis routes; and (2) to establish a broader range of MAX-phase precursors.

Li et al. recently reported the synthesis of Ti₃ZnC₂ MAX phase from the reaction of Ti₃AlC₂ in ZnCl₂ Lewis acidic molten salt at 550 °C, via a replacement reaction mechanism⁶. Ti₃ZnC₂ could be further transformed into Ti₃C₂Cl₂ MXene by increasing the MAX:ZnCl₂ ratio. However, the formation mechanism of Ti₃C₂Cl₂ MXene was not fully understood in terms of the chemistry; since the molten salt and the MAX phase shared the same element (Zn), a reaction mechanism assuming the existence of a low valence Zn₂²⁺ cation was proposed.

Herein we generalize this synthesis route to include, in addition to Zn, the A-site elements Al, Si and Ga from various MAX-phase precursors, by adjusting the chemistry of the MAX precursor and the Lewis acid melt composition. We propose a generic method to etch MAX phases by direct redox coupling between the A element and the cation of the Lewis acid molten salt, which allows us to predict the reactivity of the MAX in the molten salt and drastically increases the number of MXenes that can be prepared by this method. We also show that this method can be used to obtain MXene from MAX phases with A = Ga. The etching process is illustrated here using Ti₃C₂ prepared from Ti₃SiC₂ immersion in CuCl₂ molten salt. The obtained MXenes exhibit enhanced electrochemical performance with high Li⁺ storage capacity combined with high-rate performance in non-aqueous electrolyte, which makes these materials promising electrode materials for high-rate battery and hybrid devices such as Li-ion capacitor applications^{7,8}. This method enables the production of new 2D materials that are difficult or

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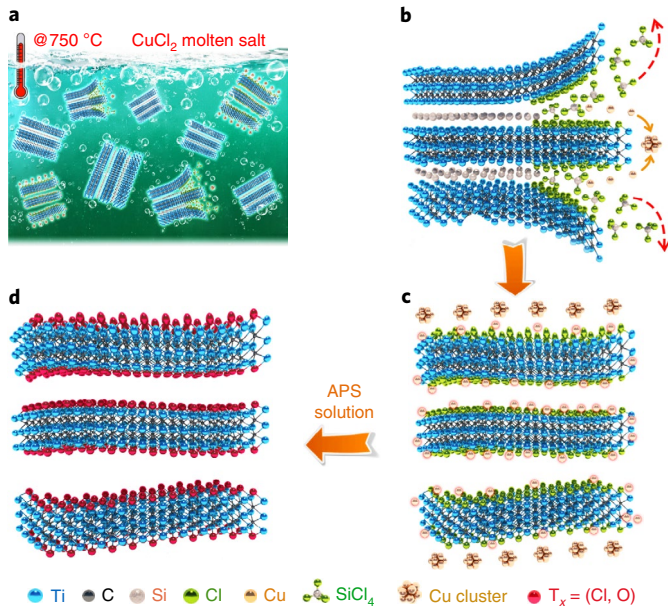
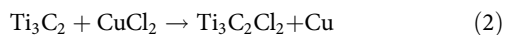
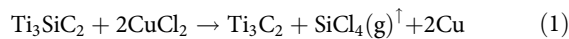


Fig. 1 | Schematic of $\text{Ti}_3\text{C}_2\text{T}_x$ MXene preparation. **a**, Ti_3SiC_2 MAX phase is immersed in CuCl_2 Lewis molten salt at 750°C . **b,c**, The reaction between Ti_3SiC_2 and CuCl_2 results in the formation of $\text{Ti}_3\text{C}_2\text{T}_x$ MXene. **d**, $\text{MS-Ti}_3\text{C}_2\text{T}_x$ MXene is obtained after further washing in ammonium persulfate (APS) solution.

even impossible to prepare via existing synthesis methods, such as HF etching. This synthetic route expands the range of MAX-phase precursors that can be used and offers important opportunities for tuning the surface chemistry and properties of MXenes.

Figure 1 shows a sketch of the $\text{Ti}_3\text{C}_2\text{T}_x$ MXene synthesis from the reaction between Ti_3SiC_2 and CuCl_2 at 750°C ; this synthesis involves the following reactions:



Ti_3SiC_2 MAX precursor is immersed at 750°C in molten CuCl_2 ($T_{\text{melting}} = 498^\circ\text{C}$). The exposed Si atoms weakly bonded to Ti in the Ti_3C_2 sublayers are oxidized into Si^{4+} cations by the Lewis acid Cu^{2+} , resulting in the formation of the volatile SiCl_4 phase ($T_{\text{boiling}} = 57.6^\circ\text{C}$) and concomitant reduction of Cu^{2+} to Cu metal (equation (1)). Similar to what has been recently reported⁶, excess Cu^{2+} partially reacts with the exposed Ti atoms from Ti_3C_2 to form metallic Cu, while charge compensation is ensured by Cl^- anions reacting to form $\text{Ti}_3\text{C}_2\text{Cl}_2$ (equation (2)). The formation mechanism of $\text{Ti}_3\text{C}_2\text{Cl}_2$ from Ti_3SiC_2 is analogous to that of chemical etching of Ti_3AlC_2 in HF solution¹: Cu^{2+} and Cl^- act as H^+ and F^- , respectively. The as-prepared powder of $\text{Ti}_3\text{C}_2\text{Cl}_2$ and Cu metal (Supplementary Fig. 1) were further immersed in ammonium persulfate (APS, $(\text{NH}_4)_2\text{S}_2\text{O}_8$) solution to remove Cu particles from the $\text{Ti}_3\text{C}_2\text{Cl}_2$ MXene surface, which also results in the addition of O-based surface groups (Supplementary Fig. 2). The final material prepared via this molten salt route is named $\text{MS-Ti}_3\text{C}_2\text{T}_x$ MXene, where T_x stands for O and Cl surface groups, and MS refers to molten salt.

X-ray diffraction (XRD) patterns of the pristine Ti_3SiC_2 before (black) and after reaction with CuCl_2 at 750°C for 24 h (noted as $\text{Ti}_3\text{SiC}_2\text{-CuCl}_2$, red), and final product after APS washing ($\text{MS-Ti}_3\text{C}_2\text{T}_x$, purple), are shown in Fig. 2a. Compared with pristine Ti_3SiC_2 , most of the diffraction peaks disappear in the final product,

leaving (00l) peaks as well as several broad and low-intensity peaks in the 2θ range from 5° to 75° ; these features indicate the successful reduction of Ti_3SiC_2 into layered Ti_3C_2 (MXene)⁹. Additionally, the shift of Ti_3C_2 (002) diffraction peaks from 2θ 10.13° to 8.07° ($\pm 0.02^\circ$) indicates an expansion of the interlayer distance from 8.8 to 10.98 Å (± 0.03 Å). The sharp and intense peaks located at approximate 2θ angles of 43.29° , 50.43° and 74.13° can be indexed as metallic Cu (Fig. 2a, red plot), which confirms the proposed etching mechanism in Lewis acid melt (equation (1)). The XRD pattern of the final product (Fig. 2a, purple plot) exhibits only the (00l) MXene peaks, confirming the removal of the Cu. A scanning electron microscopy (SEM) image of the final $\text{MS-Ti}_3\text{C}_2\text{T}_x$ sample is shown in Fig. 2b. After etching in molten salt, the Ti_3SiC_2 particle (Supplementary Fig. 1a) turns into an accordion-like microstructure (Supplementary Fig. 1b), similar to previously reported for MXenes obtained by etching in HF- or F-containing electrolyte¹. The spherical particles observed on the Ti_3C_2 before APS treatment (Supplementary Fig. 1b) are assumed to be metallic Cu produced during the etching process from equations (1) and (2) (Supplementary Fig. 1c), which can be removed by immersion in APS solution (Supplementary Fig. 2).

The lamellar microstructure of the $\text{MS-Ti}_3\text{C}_2\text{T}_x$ MXene is clearly visible in scanning transmission electron microscopy (STEM) images, as shown in Fig. 2c. The SiCl_4 gas molecules formed in situ during the etching reaction (equation (1)) are believed to act as an effective expansive agent to delaminate the MXene, similar to the preparation of expanded graphite through the decomposition of intercalated inorganic acids¹⁰.

$\text{MS-Ti}_3\text{C}_2\text{T}_x$ MXene sample surface was further characterized by X-ray photoelectron spectroscopy (XPS) analysis. Supplementary Fig. 3a shows an overview XPS spectrum for the Ti_3SiC_2 precursor (black) and $\text{MS-Ti}_3\text{C}_2\text{T}_x$ MXene (red), where the signals of Si 2p, C 1s, Ti 2p and O 1s are observed at 102, 285, 459 and 532 eV, respectively¹¹. The disappearance of the Si signal confirms the effectiveness of Si removal by the Lewis acid etching reaction (Supplementary Fig. 3b). Similarly, no notable amounts of Cu or S element were detected (Supplementary Fig. 4a,b). The deconvolution of the Ti 2p spectra (Fig. 2d) in the energy range between 454 and 460 eV was achieved following previous works^{6,12} and details are given in Supplementary Table 1. The Ti 2p spectra show the existence of Ti-O and Ti-Cl chemical bonds, most likely from O and Cl surface groups associated with partial surface oxidation. The observed Ti-C bonds come from the core $[\text{TiC}_6]$ octahedral building blocks of the Ti_3C_2 MXene. The fitting of the O 1s (Supplementary Fig. 4c) and C 1s (Supplementary Fig. 4d) spectra show O-terminated surface functional groups on the $\text{MS-Ti}_3\text{C}_2\text{T}_x$ sample. The XPS signal of the Cl 2p energy level confirms the presence of Ti-Cl bonds (Supplementary Fig. 4e). The Cl groups are expected from equation (2), while O surface functional groups are formed during the oxidation treatment in APS solution and subsequent washing process¹³. Elemental analysis (Supplementary Table 3) revealed an O-termination-group content of about 12 wt% and a Cl-termination-group content of about 14 wt% in the $\text{MS-Ti}_3\text{C}_2\text{T}_x$ MXene, resulting in a composition of $\text{Ti}_3\text{C}_{1.94}\text{Cl}_{0.77}\text{O}_{1.71}$ for the $\text{MS-Ti}_3\text{C}_2\text{T}_x$ MXene.

Temperature-programmed desorption coupled with mass spectroscopy analysis was performed on $\text{MS-Ti}_3\text{C}_2\text{T}_x$ MXene samples and MXene prepared via conventional etching treatment in HF, termed herein $\text{HF-Ti}_3\text{C}_2\text{T}_x$ (Supplementary Fig. 5 and Supplementary Table 4). H_2O release was observed below 400°C for both samples, corresponding to surface-adsorbed and intercalated water resulting from the washing with water after synthesis¹⁴. Unlike $\text{HF-Ti}_3\text{C}_2\text{T}_x$, $\text{MS-Ti}_3\text{C}_2\text{T}_x$ MXene shows substantial CO_2 release below 600°C , which could be ascribed to the partial oxidation of C by the APS. Also noteworthy is the absence of any release of -OH surface groups for $\text{MS-Ti}_3\text{C}_2\text{T}_x$ MXene, decreasing the hydrophilicity of the surface. The Cl group is stable on Ti_3C_2 at 750°C (ref. 15), but a trace of

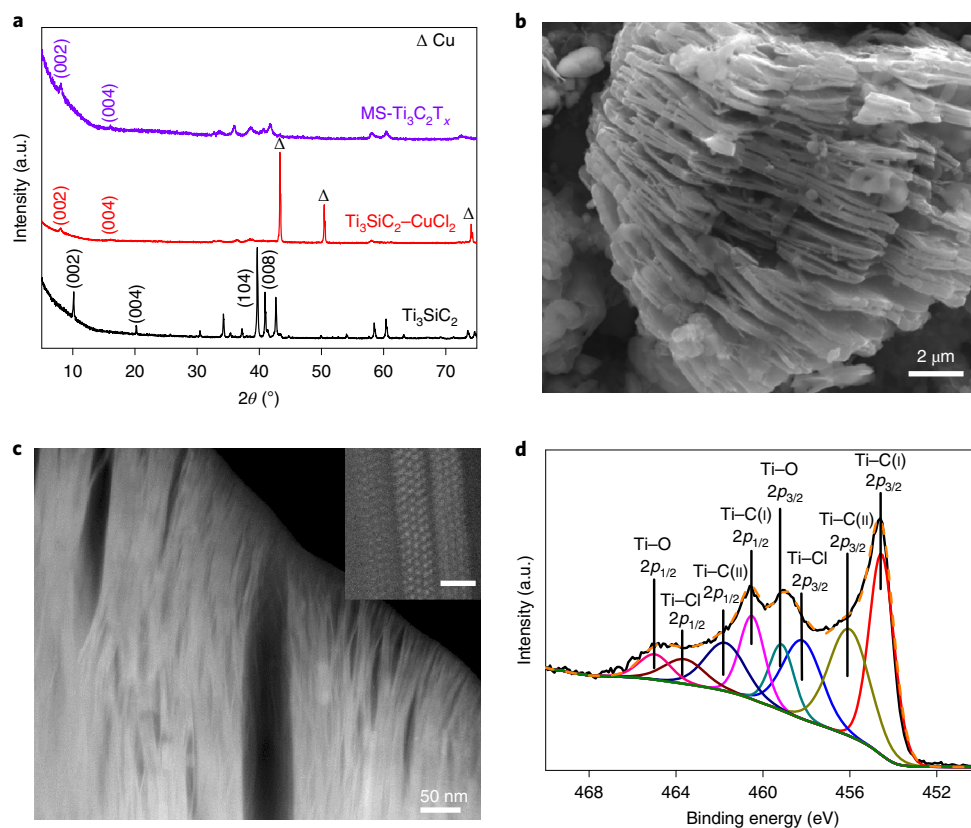
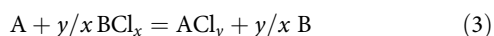


Fig. 2 | Morphological and structural characterizations of MS-Ti₃C₂T_x MXene. **a**, XRD patterns of pristine Ti₃SiC₂ before (black line) and after (red line) reaction with CuCl₂, and final MS-Ti₃C₂T_x MXene obtained after washing in 0.1M (NH₄)₂S₂O₈ solution (purple line). **b, c**, SEM image (**b**) and cross-sectional STEM image (**c**) showing the nanolaminate nature of the material (scale bar in the atomically resolved image inset in **c** is 1 nm). **d**, XPS analysis of the Ti 2p energy level from the MS-Ti₃C₂T_x MXene sample.

released Cl, as well as SO₂, is still detected below 600 °C, the latter coming from the APS treatment.

The ability of Lewis acids to withdraw electrons from A elements in the MAX phase tends to be accurately reflected by their respective electrochemical redox potentials in halide melts. For instance, the Si⁴⁺/Si couple has a redox potential as low as -1.38 V versus Cl₂/Cl⁻ at 700 °C. As a result, CuCl₂ molten salt (redox potential of -0.43 V versus Cl₂/Cl⁻) can easily oxidize Si into Si⁴⁺ (etching/exfoliation of MAX phase into MXene). The present Lewis acid etching process can then be generalized to prepare a broad family of MXene materials. Figure 3a shows a Gibbs free energy mapping prepared from thermodynamic data (see equation (3) and Supplementary Fig. 6) to guide the selection of effective Lewis acids for MAX phases with different A elements. In these calculations, the etching is independent of both the composition of the MX layer and the value of *n* in M_{n+1}AX_n. The colour of each spot/star in Fig. 3a indicates the value of the Gibbs free energy of the reaction between the selected A element in the MAX phase and the Lewis acid Cl melt at 700 °C (equation (3)).



From these thermodynamic calculations, etching of an A element from MAX can be achieved by using a Lewis acid with a higher redox potential. Based on this map, a series of MAX phases—specifically Ti₂AlC, Ti₃AlC₂, Ti₃AlCN, Nb₂AlC, Ta₂AlC, Ti₂ZnC and Ti₃ZnC₂—were successfully exfoliated into corresponding MXenes (Ti₂CT_x, Ti₃C₂T_x, Ti₃CNT_x, Nb₂CT_x, Ta₂CT_x, Ti₂CT_x, Ti₃C₂T_x) using various chloride molten salts (CdCl₂, FeCl₂, CoCl₂, CuCl₂, AgCl, NiCl₂) as

indicated by the stars in Fig. 3a. The SEM images in Fig. 3b–g show the lamellar microstructures of various obtained MXenes. The successful preparation of Ta₂CT_x and Ti₃C₂T_x MXenes from Ta₂AlC and Ti₃SiC₂, which were theoretically predicted to be hard to exfoliate¹⁶, provides evidence of the effectiveness of the Lewis acid molten salt route. Additional information about the as-prepared MXenes can be found in Supplementary Figs. 8–15, including the use of FeCl₃ as alternative etching agent to remove Cu (Supplementary Fig. 16).

The diversity and green chemistry of Lewis acids in inorganic salts offer an unexplored parameter space to optimize such etching methodology. At the same time, this approach broadens the choice of MAX-phase families for MXene fabrication and offers opportunities to tune the surface chemistry of MXene materials by using various molten salts based on other anions (such as Br⁻, I⁻, SO₄²⁻ or NO₃⁻). Supplementary Fig. 18a,c shows examples of the successful preparation of MS-Ti₃C₂ MXene using CuBr₂ and CuI molten salts as Lewis acidic etchants. Energy-dispersive spectroscopy (EDS) analysis (Supplementary Fig. 18b,d) shows the presence of -I and -Br surface groups on MXenes, confirming that the surface functional groups can be tuned by careful selection of the Lewis acid etchant. By successfully extending the etchant from Cl to Br and I, the present Lewis acidic etching route offers many opportunities to tune the surface chemistry of MXene materials.

Layered MS-Ti₃C₂T_x MXene powders derived from Ti₃SiC₂ (Fig. 2b) were further used to prepare electrodes by mixing with carbon-containing conducting additive/binder (see Methods for details). A conducting additive, 15 wt% carbon black, was added to improve the electrical percolation in the electrode film. Figure 4a shows cyclic voltammetry profiles of the MS-Ti₃C₂T_x MXene

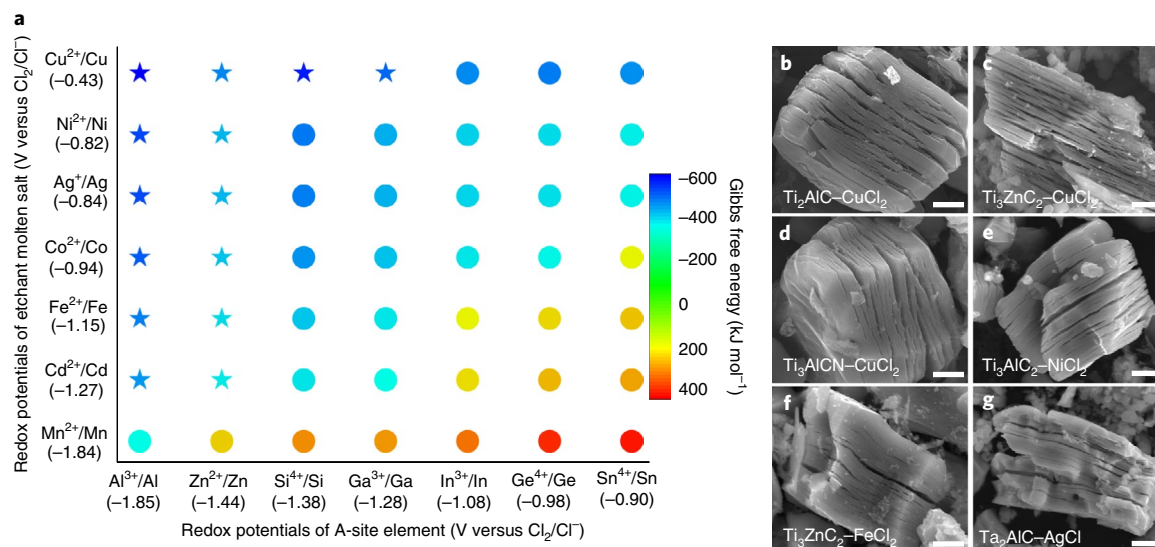


Fig. 3 | Generalization of the Lewis acid etching route to a large family of MAX phases. **a**, Gibbs free energy mapping (700 °C) guiding the selection of Lewis acid Cl salts according to the electrochemical redox potentials of A-site elements in MAX phases (x axis) and molten salt cations (y axis) in Cl melts. Stars mark the corresponding MXenes demonstrated in the current study. All symbols (stars and spots) were calculated using the same approach; the star symbols were experimentally verified in this paper, while spot symbols remain theoretical prediction. **b-g**, SEM images reveal the typical accordion morphology of MXenes from different MAX phases etched by various Lewis acid Cl salts, such as Ti_2AlC by CuCl_2 (**b**), Ti_3ZnC_2 by CuCl_2 (**c**), Ti_3AlCN by CuCl_2 (**d**), Ti_3AlC_2 by NiCl_2 (**e**), Ti_3ZnC_2 by FeCl_2 (**f**) and Ta_2AlC by AgCl (**g**). Scale bars, 2 μm .

electrode in 1 M LiPF_6 (in 1:1 vol/vol ethylene carbonate/dimethyl carbonate) electrolyte recorded at 0.5 mV s^{-1} with different negative cut-off potentials. The electrochemical signature is remarkable as it differs from what is currently known for conventional MXene prepared from etching in HF- or F-containing electrolyte (hereinafter termed HF-MXene) in non-aqueous electrolytes¹⁷⁻¹⁹. The Li^+ intercalation/deintercalation reaction in HF-MXene²⁰⁻²⁴ occurs over a large range of potentials (usually from $\sim 0.05 \text{ V}$ up to 3 V versus Li^+/Li). Such a large operating potential window, resulting in a high average operating potential for a negative electrode, drastically limits the possibilities for using MXene materials in (Li-ion) battery applications. Moreover the Li^+ intercalation/deintercalation reaction in HF- Ti_3C_2 MXene is achieved in a sequential way, resulting in the presence of sets of redox peaks associated with Li^+ intercalation in interlayer slits²¹. However, the charge storage mechanism in MS- $\text{Ti}_3\text{C}_2\text{T}_x$ MXene is achieved with a constant current versus applied potential, similarly to what is observed in a pseudocapacitive material, with an almost constant current during the reduction and oxidation process in a potential range between 0.2 and 2.2 V versus Li^+/Li . The discharge capacity of the MS- $\text{Ti}_3\text{C}_2\text{T}_x$ MXene powder in this non-aqueous Li-ion battery electrolyte reaches 738 C g^{-1} (205 mAh g^{-1}) at 0.5 mV s^{-1} within the full potential window of 2.8 V, which translates into 323 F g^{-1} within 2 V (see Supplementary Fig. 19). These are, to the best of our knowledge, the highest capacitance values reported for Ti_3C_2 MXene in non-aqueous electrolytes^{17,25,26}. This performance makes MXene materials suitable for use as negative electrodes in non-aqueous energy storage devices. It is also important to note that unlike previous works in which electrodes had to be prepared from filtration of delaminated MXene suspensions to achieve high electrochemical performance²⁷, raw, non-delaminated MXene powders (Fig. 2b) were used here to prepare the electrode films. This broadens the range of application of the materials to prepare electrodes for energy-storage devices.

The charge storage mechanism was investigated using in situ XRD during cyclic voltammetry experiments at 0.5 mV s^{-1} . Figure 4b shows the change in the d -spacing calculated from the

(002) peak during anodic and cathodic scans for three different cycles. The initial d -spacing was found to be $\sim 11.02 \text{ \AA}$, and the d -spacing was found to be roughly constant during the polarization with a maximum change of 0.25 \AA . The small value of the d -spacing indicates that MXene layers are separated by $\sim 3 \text{ \AA}$: this supports the intercalation of desolvated Li^+ ions between the MXene layers, such as recently reported¹⁷, blocking the co-intercalation of solvent molecules and resulting in improved electrochemical performance. During the cathodic scan (Fig. 4a), Li^+ ions are intercalated between the MXene layers; this is assumed to be associated with the change in the oxidation state of Ti, such as observed in Li-ion batteries during Li^+ intercalation²². Li^+ deinsertion from the MXene structure occurs during the anodic potential scan, with a remarkable mirror-like cyclic voltammetry profile shape. During the first cycle upon reduction, an irreversible capacity is observed (Supplementary Fig. 20a), as a result of the formation of the solid electrolyte interphase layer²⁸.

Figure 4c shows the change of the $\text{Ti}_3\text{C}_2\text{T}_x$ MXene capacity with discharge time calculated from cyclic voltammetry profiles obtained at various potential scan rates (Supplementary Fig. 20b and Supplementary Table 8). The capacity reaches 738 C g^{-1} (205 mAh g^{-1}) for a discharge time of 1.5 h (C/1.5 rate). This value corresponds to a minimum of 1.22 F exchanged per mole of Ti_3C_2 , which is ~ 0.4 electrons transferred per Ti atom (considering 10 mAh g^{-1} coming from the acetylene black), much higher than previously reported values^{17,29}. Interestingly, a similar number of 0.4 electrons transferred per Ti was found for Ti_2CT_x MXene, as shown in Supplementary Fig. 21, demonstrating that Ti is the redox active centre.

Compared with OH-terminated MXene surfaces, Xie et al. reported density field theory calculations giving a lower adsorption energy of Li onto O-terminated MXenes, resulting in stronger interaction between O and Li and improved Li capacity²². As a result, the absence of OH surface groups and the high oxygen content (Supplementary Fig. 5 and Supplementary Table 3) of the MS- $\text{Ti}_3\text{C}_2\text{T}_x$ MXene in relation to the specific synthesis route in molten salt are also assumed to contribute to the improvement in capacity.

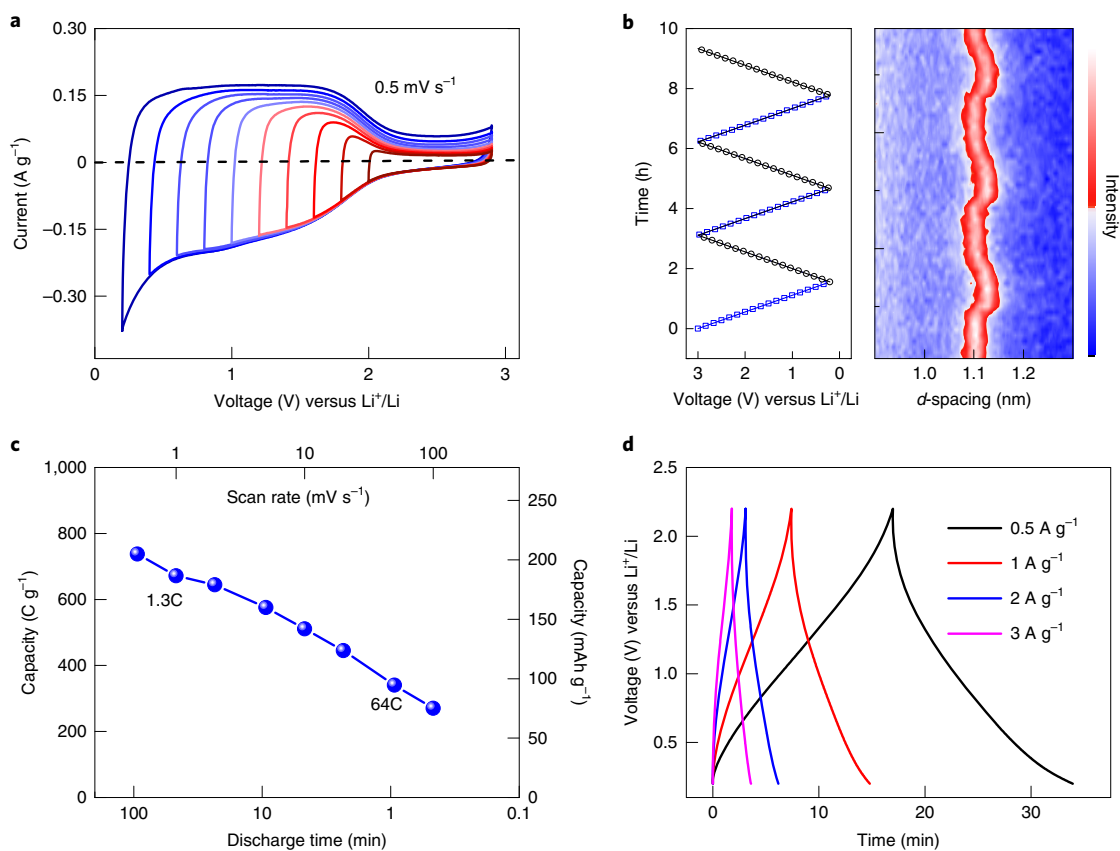


Fig. 4 | Electrochemical characterizations of MS-Ti₃C₂T_x MXene electrode in 1M LiPF₆ (in 1:1 ethylene carbonate/dimethyl carbonate) electrolyte.

a, Cyclic voltammety profiles at a 0.5 mV s⁻¹ potential scan rate with various cut-off negative potentials; the profiles exhibit a mirror-like shape with no redox peaks during the Li⁺ intercalation/deintercalation redox reaction. **b**, In situ XRD maps of the (002) peak derived *d*-spacing during anodic and cathodic scans for three different cycles; the change of *d*-spacing is less than 0.25 Å during cycling. **c**, Change of the MXene electrode capacity versus the discharge time during cyclic voltammety recorded at various potential scan rates from anodic scans; 1C rate corresponds to a full discharge in 1h. The active material weight loading is 1.4 mg cm⁻². **d**, Galvanostatic charge/discharge curves at current densities from 0.5 to 3 A g⁻¹.

The Ti₃C₂T_x electrode nevertheless delivers 142 mAh g⁻¹ for a 280 s discharge time (13C rate) and 75 mAh g⁻¹ for a discharge time of less than 30 s (128C rate). Taken together with the galvanostatic plots achieved at various current densities (Supplementary Fig. 22a), these results highlight the high power performance of the present Ti₃C₂ MXene material as an electrode during the Li⁺ ion intercalation reaction, which occurs at lower average potential versus Li⁺/Li compared with previously reported pseudocapacitive materials^{30,31}. As a result of these high power performances, the electrochemical impedance spectroscopy plots recorded at various bias potentials (Supplementary Fig. 23) show a charge-transfer resistance of ~25 Ω cm² followed by a restricted-diffusion behaviour with a rapid increase in the imaginary part of the impedance at low frequency³².

An increase in the electrode weight loading (4 mg cm⁻²) does not substantially affect the power capability (Supplementary Fig. 20c,d). Galvanostatic charge/discharge measurements (Fig. 4d) confirm the unique electrochemical signature of the electrode in non-aqueous electrolyte with a sloping voltage profile within a potential range of 0.2–2.2 V versus Li⁺/Li, as expected from the cyclic voltammety profiles shown in Fig. 4a. Last, but not least, cycle stability was impressive with 90% capacity retention after 2,400 cycles (Supplementary Fig. 20b). Similar remarkable electrochemical signatures and performances were obtained for other MS-MXenes studied here, such as can be seen more specifically from the cyclic voltammety profiles and power performance of a Ti₃C₂T_x electrode prepared from Ti₃AlC₂ MAX phase (Supplementary Fig. 24).

In conclusion, the combination of a mirror-like electrochemical signature during Li⁺ intercalation/deintercalation in non-aqueous Li⁺-containing electrolyte, together with high capacity, high rate discharge and charge performance (<1 min) and the low operating potential range (0.2–2.2 V versus Li⁺/Li), makes these MXenes prepared via a molten salt route potentially useful as negative electrodes in electrochemical energy storage devices (batteries and Li-ion capacitors). The general Lewis acid etching route proposed here expands the range of MAX-phase precursors that can be used to prepare new MXenes, and offers unprecedented opportunities for tailoring the surface chemistry, and consequently the properties, of MXene materials.

Online content

Any methods, additional references, Nature Research reporting summaries, source data, extended data, supplementary information, acknowledgements, peer review information, details of author contributions and competing interests, and statements of data and code availability are available at <https://doi.org/10.1038/s41563-020-0657-0>.

References

- Naguib, M. et al. Two-dimensional nanocrystals produced by exfoliation of Ti₃AlC₂. *Adv. Mater.* **23**, 4248–4253 (2011).

2. Lukatskaya, M. R. et al. Cation intercalation and high volumetric capacitance of two-dimensional titanium carbide. *Science* **341**, 1502–1505 (2013).
3. Anasori, B., Lukatskaya, M. R. & Gogotsi, Y. 2D metal carbides and nitrides (MXenes) for energy storage. *Nat. Rev. Mater.* **2**, 16098 (2017).
4. Ghidui, M., Lukatskaya, M. R., Zhao, M.-Q., Gogotsi, Y. & Barsoum, M. W. Conductive two-dimensional titanium carbide ‘clay’ with high volumetric capacitance. *Nature* **516**, 78–81 (2014).
5. Feng, A. et al. Fabrication and thermal stability of NH_4HF_2 -etched Ti_3C_2 MXene. *Ceram. Int.* **43**, 6322–6328 (2017).
6. Li, M. et al. Element replacement approach by reaction with Lewis acidic molten salts to synthesize nanolaminated MAX phases and MXenes. *J. Am. Chem. Soc.* **141**, 4730–4737 (2019).
7. Naoi, K. et al. Ultrafast charge-discharge characteristics of a nanosized core-shell structured LiFePO_4 material for hybrid supercapacitor applications. *Energy Environ. Sci.* **9**, 2143–2151 (2016).
8. Lukatskaya, M. R., Dunn, B. & Gogotsi, Y. Multidimensional materials and device architectures for future hybrid energy storage. *Nat. Commun.* **7**, 12647 (2016).
9. Alhabeib, M. et al. Selective etching of silicon from Ti_3SiC_2 (MAX) to obtain 2D titanium carbide (MXene). *Angew. Chem. Int. Ed.* **130**, 5542–5546 (2018).
10. Yang, S. et al. Ultrafast delamination of graphite into high-quality graphene using alternating currents. *Angew. Chem. Int. Ed.* **56**, 6669–6675 (2017).
11. Kisi, E., Crossley, J., Myhra, S. & Barsoum, M. Structure and crystal chemistry of Ti_3SiC_2 . *J. Phys. Chem. Solids* **59**, 1437–1443 (1998).
12. Halim, J. et al. X-ray photoelectron spectroscopy of select multi-layered transition metal carbides (MXenes). *Appl. Surf. Sci.* **362**, 406–417 (2016).
13. Çakır, O. Review of etchants for copper and its alloys in wet etching processes. *Key Eng. Mater.* **364–366**, 460–465 (2008).
14. Shpigel, N. et al. Direct assessment of nanoconfined water in 2D Ti_3C_2 electrode interspaces by a surface acoustic technique. *J. Am. Chem. Soc.* **140**, 8910–8917 (2018).
15. Lu, J. et al. $\text{Ti}_{n+1}\text{C}_n$ MXenes with fully saturated and thermally stable Cl terminations. *Nanoscale Adv.* **9**, 3680–3685 (2019).
16. Khazaei, M. et al. Insights into exfoliation possibility of MAX phases to MXenes. *Phys. Chem. Chem. Phys.* **20**, 8579–8592 (2018).
17. Wang, X. et al. Influences from solvents on charge storage in titanium carbide MXenes. *Nat. Energy* **4**, 241–248 (2019).
18. Wang, X. et al. Pseudocapacitance of MXene nanosheets for high-power sodium-ion hybrid capacitors. *Nat. Commun.* **6**, 6544 (2015).
19. Come, J. et al. A non-aqueous asymmetric cell with a Ti_2C -based two-dimensional negative electrode. *J. Electrochem. Soc.* **159**, A1368–A1373 (2012).
20. Ren, C. E. et al. Porous two-dimensional transition metal carbide (MXene) flakes for high-performance Li-Ion storage. *ChemElectroChem* **3**, 689–693 (2016).
21. Cheng, R. et al. Understanding the lithium storage mechanism of $\text{Ti}_3\text{C}_2\text{T}_x$ MXene. *J. Phys. Chem. C* **123**, 1099–1109 (2019).
22. Xie, Y. et al. Role of surface structure on Li-ion energy storage capacity of two-dimensional transition-metal carbides. *J. Am. Chem. Soc.* **136**, 6385–6394 (2014).
23. Kajiyama, S. et al. Enhanced Li-ion accessibility in MXene titanium carbide by steric chloride termination. *Adv. Energy Mater.* **7**, 1601873 (2017).
24. Zhao, J. et al. One-step synthesis of few-layer niobium carbide MXene as a promising anode material for high-rate lithium ion batteries. *Dalton Trans.* **48**, 14433–14439 (2019).
25. Pang, J. et al. Applications of 2D MXenes in energy conversion and storage systems. *Chem. Soc. Rev.* **48**, 72–133 (2019).
26. Xiong, D., Li, X., Bai, Z. & Lu, S. Recent advances in layered $\text{Ti}_3\text{C}_2\text{T}_x$ MXene for electrochemical energy storage. *Small* **14**, 1703419 (2018).
27. Lukatskaya, M. R. et al. Ultra-high-rate pseudocapacitive energy storage in two-dimensional transition metal carbides. *Nat. Energy* **2**, 17105 (2017).
28. Birkel, C. R., Roberts, M. R., McTurk, E., Bruce, P. G. & Howey, D. A. Degradation diagnostics for lithium ion cells. *J. Power Sources* **341**, 373–386 (2017).
29. Lukatskaya, M. R. et al. Probing the mechanism of high capacitance in 2D titanium carbide using in situ X-ray absorption spectroscopy. *Adv. Energy Mater.* **5**, 1500589 (2015).
30. Augustyn, V. et al. High-rate electrochemical energy storage through Li^+ intercalation pseudocapacitance. *Nat. Mater.* **12**, 518–522 (2013).
31. Kim, H.-S. et al. Oxygen vacancies enhance pseudocapacitive charge storage properties of MoO_{3-x} . *Nat. Mater.* **16**, 454–460 (2017).
32. Diard, J. P., Gorrec, B. L. & Montella, C. Linear diffusion impedance. General expression and applications. *J. Electroanal. Chem.* **471**, 126–131 (1999).

Methods

Preparation of MXenes from Lewis acid molten salt route. Various MAX phases and Lewis acid salts were used to prepare MXenes (summarized in Supplementary Table 7). We here take Ti_3SiC_2 MAX phase and CuCl_2 as an example: 1 g of Ti_3SiC_2 MAX-phase powder and 2.1 g of CuCl_2 powder were mixed (in a stoichiometric molar ratio of 1:3) and ground for 10 min. Then 0.6 g of NaCl and 0.76 g of KCl were added to this mixture, which was ground for another 10 min. The mixture was placed in an alumina boat, and the boat was then put into an alumina tube with argon flow. The powder mixture was heated to 750 °C with a heating ramp of 4 °C min⁻¹, and held for 24 h. The obtained products were washed with deionized water to remove salts, and MXene/Cu mixed particles were obtained. The mixtures of MXene/Cu were then washed by 0.1 M APS solution to remove the residual Cu particles¹³. The resulting solution was further cleaned five times with deionized water and five times alcohol and filtered with a microfiltration membrane (polyvinylidene fluoride, 0.45 μm). Finally, the MXene powders (denoted as MS- $\text{Ti}_3\text{C}_2\text{T}_x$) were dried under vacuum at room temperature for 24 h.

Materials characterizations. The phase composition of the samples was analysed by XRD (D8 Advance, Bruker AXS) with Cu K_α radiation. XRD patterns were collected with 2θ steps of 0.02° with a collection time of 1 s per step. The microstructures and chemical compositions were analysed by SEM (QUANTA 250 FEG, FEI) at 20 kV and EDS; EDS values were fitted by XPP (extended Pouchou–Pichoir) model. The chemical composition and bonding states were measured by XPS using a Kratos AXIS ULTRA^{DL} instrument with a monochromic Al K_α X-ray source ($h\nu = 1486.6$ eV). The power was 96 W, and the X-ray spot size was set to 700 × 300 μm. The pass energy of the XPS analyser was set at 20 eV. The pressure of the analysis chamber was kept below 5×10^{-9} torr. All spectra were calibrated using the binding energy of C 1s (284.8 eV) as a reference. The XPS atomic sensitivity factors involved in the atomic concentration calculation were 0.278 (C 1s), 1.833 (Ca 2p), 2.001 (Ti 2p) and 0.78 (O 1s), according to Kratos Vision Processing software. Etch conditions were defined by a beam energy of 4 kV, a current of 100 μA, and a raster size of 3 mm. TEM and high-resolution TEM images were obtained using a Tecnai F20 (FEI) electron microscope at an acceleration voltage of 200 kV. Structural and chemical analysis was carried out by high-resolution STEM imaging and STEM-EDS within Linköping's double Cs corrected FEI Titan3 60-300 microscope operated at 300 kV; STEM energy-dispersive X-ray analysis was recorded with the embedded high-sensitivity Super-X detector. Temperature-programmed desorption was performed under an inert atmosphere (Ar, 100 ml min⁻¹). The sample (10–20 mg) was placed in a thermobalance and heat treated up to 1,300 °C at a rate of 10 °C min⁻¹. The decomposition products (gas evolved) were monitored by online mass spectrometry (Skimmer, Netzsch). Quantification of Cl and S could not be achieved due to the absence of standards.

Electrochemical measurements. MS- $\text{Ti}_3\text{C}_2\text{T}_x$ MXene self-standing electrodes were prepared by mixing the MXene powder with 15 wt% carbon black and 5 wt% PTFE binder, and laminated many times to obtain films of different thicknesses. The active material weight loading was calculated by dividing the mass (mg) of MXene active material by the electrode area (cm²). Metallic lithium foil was used as the counterelectrode and reference electrode, LP30 (1 M LiPF₆ in 1:1 vol/vol ethylene carbonate/dimethyl carbonate) as the electrolyte, and two slides of 25-μm-thick cellulose as the separator. Swagelok cells were assembled in the argon-filled glovebox with oxygen and water content <0.1 ppm. All electrochemical tests were performed using a VMP3 potentiostat (Biologic). Cyclic voltammetry and galvanostatic tests were conducted in two-electrode mode versus a Li electrode. Electrochemical impedance spectroscopy was carried out with a potential amplitude of 10 mV in the range from 10 mHz to 200 kHz.

In cyclic voltammetry, the capacity (C g⁻¹) and average capacitance (F g⁻¹) of a single electrode are evaluated from the anodic scan using the following equations:

$$Q = \frac{\int_0^{V/s} |i| dt}{m} \quad (4)$$

$$C = \frac{Q}{V} \quad (5)$$

where i (A) is the current changed by time t (s), m is the mass of active material (g), V is the potential window (V) and s is the scan rate (V s⁻¹).

In galvanostatic charge/discharge plots, the capacity (C g⁻¹) is given by:

$$Q = \frac{i\Delta t}{m} \quad (6)$$

where Δt (s) is charging/discharging time and i (A) is the charging/discharging current.

In situ XRD. In situ XRD was conducted on a Bruker D8 Advance diffractometer using a Cu K_α radiation source. A two-electrode Swagelok cell system³³, with MS- $\text{Ti}_3\text{C}_2\text{T}_x$ MXene film as the working electrode, a beryllium window as the current collector and Li metal as the counterelectrode, was used to perform the electrochemical test for the in situ XRD measurements. All XRD patterns were recorded during cyclic voltammetry tests at a potential scan rate of 0.5 mV s⁻¹. The (002) peak located between 6° and 10° was recorded to calculate the interlayer d -spacing (Fig. 4b).

Data availability

Source data for Figs. 2a,d and 4 are provided with the paper. The remaining data that support the findings of this study are available from the corresponding authors upon reasonable request.

References

33. Morcrette, M. et al. In situ X-ray diffraction techniques as a powerful tool to study battery electrode materials. *Electrochim. Acta* **47**, 3137–3149 (2002).

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Author contributions

Q.H., Z.L., P.-L.T. and P.S. designed the research. Y.L. conducted material preparations and most of the characterization. H.S. and L.L. conducted the electrochemical tests and Z.L., P.-L.T. and P.S. analysed the data. H.S., B.D. and P.R. carried out the in situ XRD. J.L., P.O.Á.P., P.E. and L.H. carried out the STEM measurements. E.R.-P. performed the temperature-programmed desorption/mass spectrometry. Z.L., P.S. and Q.H. prepared the manuscript. All authors contributed to the discussion of the data and writing the paper.

Competing interests

The authors declare no competing interests.

Additional information

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