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## Two-dimensional $Ti_3C_2T_x$ MXene nanosheets for $CO_2$ electroreduction in aqueous electrolytes†

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Electroreduction of carbon dioxide ( $CO_2$ ) to valuable synthons is a sustainable and viable technique to reduce the carbon footprint in the atmosphere. In this work, two-dimensional  $Ti_3C_2T_x$  nanosheets have been investigated for selective  $CO_2$  adsorption and its subsequent electroreduction to valuable products in an aqueous electrolyte medium. Various physical and electrochemical characterizations have been performed to ascertain the structural, morphological, and electrocatalytic characteristics of  $Ti_3C_2T_x$ .  $Ti_3C_2T_x$  nanosheets show a  $CO_2$  adsorption capacity of  $0.16\text{ mmol g}^{-1}$  under ambient conditions and exhibit a maximum faradaic efficiency of 96% towards  $CO_2$  electroreduction to  $C_1$  and  $C_{2+}$  products, mainly CO (42.2%), methanol (23.6%), ethanol (20.1%), and acetone (10.1%). Besides, it also shows high catalytic stability for product formation for a tested period of 72 h. This report sheds new light on the design of titanium-based advanced catalysts for efficient  $CO_2$  conversion in a benign aqueous medium.

### 1. Introduction

In this era, energy production and transportation are mainly dependent on fossil fuels which increases the emission of greenhouse gases into the atmosphere. Particularly, carbon dioxide ( $CO_2$ ) is one of the primary greenhouse gases, increasing dramatically in the earth's atmosphere. Rising  $CO_2$  levels ( $>417\text{ ppm}$ , as per NOAA-Mauna Lab Observatory, 2022) adversely affect the earth's atmosphere.<sup>1,2</sup> This mandates an urgent need to mitigate  $CO_2$  emissions in the atmosphere. Besides its antagonistic environmental impact,  $CO_2$  is also an abundant  $C_1$  source for several organic products. Therefore, efforts involving  $CO_2$  mitigation by transforming it into valuable products have currently drawn huge attention.<sup>3,4</sup> With the aim of  $CO_2$  mitigation efforts, among the different methods reported earlier, electrochemical conversion of  $CO_2$  into value-added products ( $C_1$  and  $C_{2+}$  carbon products) is considered an efficient and viable route for the sustainable ecosystem due to its ambient operating conditions.<sup>4,5</sup>

In the process of  $CO_2$  electroreduction, various  $C_1$  products (carbon monoxide, methane, formate, methanol),  $C_2$  products (ethanol, ethylene), and  $C_{2+}$  products (propanol, acetone, acetate) can be formed using different homo/heterogeneous

catalysts. Among them, methanol, ethanol, ethylene, and acetone are highly promising chemical synthons in industry as well as synthetic fuels to meet fuel scarcity. However, the chemically inert  $CO_2$  molecule significantly hinders the formation of various carbon products.<sup>6</sup> Additionally, the process of  $CO_2$  electroreduction confronts two major challenges: (i) the inferior binding capability of the  $CO_2$  reduction intermediate on electrode materials and (ii) the competitive hydrogen evolution reaction (HER) occurring during the electroreduction process.<sup>7,8</sup> To address these issues, the design of an electrocatalyst is of substantial interest. Recently, various transition metal-based oxides/hydroxides<sup>9</sup> and metal-free carbonaceous materials<sup>10</sup> have been reported for  $CO_2$  electroreduction. Among them, copper (Cu) and copper-containing electrocatalysts are well-recognized for the formation of the above-mentioned  $C_2$  and  $C_{2+}$  products due to their effective binding ability with  $*CO$  and  $*CHO$  intermediates.<sup>11,12</sup> The quest for relatively low-cost and earth-abundant highly active catalysts with enhanced charge transfer, which is highly desired to steer the enhancement in the fundamental catalytic activity towards value-added  $C_2$  and  $C_{2+}$  products, still continues.

In the last decade, new families of two-dimensional materials, namely, metal carbides, nitrides, and carbonitrides (MXenes), discovered by Gogotsi *et al.*, have aroused much attention. The applications of MXenes are extensively explored in the field of supercapacitors,<sup>13</sup> batteries,<sup>14</sup> electrochemical sensors,<sup>15</sup> and water splitting<sup>16</sup> applications due to their excellent electronic conductivity, inherent surface functional groups, and tunable interlayer spacing.<sup>17,18</sup> MXenes ( $M_{n+1}X_nT_x$ , where M is the transition element, X is carbon or nitrogen, and  $T_x$  is the surface

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functional group such as  $-O$ ,  $-F$  and  $-OH$ ) are obtained by selective etching of A elements (A represents group 13 and 14 elements) from their parent MAX phases which have a general formula of  $M_{n+1}AX_n$ .<sup>19</sup>

Apart from their wide range of applications, recent experimental and theoretical investigations proved the efficacy of layered MXenes for  $CO_2$  capture and its electrocatalytic conversion.<sup>20–22</sup> Reports have suggested that the interlayer spacing between MXene lamellar structures and its high electrical conductivity are conducive to  $CO_2$  adsorption and its efficient conversion.<sup>23–25</sup> It has been identified that the inherent surface functional groups ( $T_x = -F$ ,  $-O$ , and  $-OH$ ) in MXenes can change the MXene work function that effectively favors  $CO_2$  reduction intermediate adsorption. Also, the metal centers in MXenes can increase  $CO_2$  adsorption through non-covalent ( $O=C=O_{lp} \cdots M$ ) or covalent (surface-bound  $CO_2$ ) interactions.<sup>24,26,27</sup> For instance, Wang *et al.* reported two kinds of MXenes ( $Ti_3C_2T_x$  and  $V_2CT_x$ ) for  $CO_2$  capture and obtained a maximum  $CO_2$  adsorption capacity of  $5.79 \text{ mmol g}^{-1}$  at 40 bar pressure.<sup>25</sup> In recent years, a few reports investigated MXenes for electrocatalytic conversion of  $CO_2$ . For example, Qu *et al.* investigated that the use of N-doping and titanium vacancies in the  $Ti_3C_2$  MXene shows 92% faradaic efficiency towards CO production.<sup>28</sup> Hondoko *et al.* reported the use of  $Ti_2CT_x$  and  $Mo_2CT_x$  for the electrocatalytic conversion of  $CO_2$  into formic acid as a primary product with a 56% faradaic efficiency. They suggested that the inherent functional groups ( $T_x$ ) present in the materials are the reason for such high performance.<sup>22</sup>  $Ti_3C_2$  and  $Mo_2C$  were investigated by Attanayake *et al.* for  $CO_2$  reduction and showed a faradaic efficiency of 90% towards CO production by using ionic liquid-based electrolytes.<sup>20</sup> Also, a copper immobilized  $Ti_3C_2Cl_x$  MXene derived from a quaternary MAX phase ( $Ti_3(Al_{1-x}Cu_x)C_2$ ) was reported for electrochemical  $CO_2$  reduction, where methanol and ethanol were the major products due to the unsaturated electronic structure of Cu.<sup>28</sup> Bao *et al.* synthesized a copper single atom decorated  $Ti_3C_2T_x$  MXene for effective carbon monoxide reduction and achieved 98% of multi-carbon products. This is due to the formation of a favorable C–C coupling to generate the key intermediates ( $*CO-CHO$ ) for multi-carbon product formation.<sup>29</sup>

Of all MXene-based  $CO_2$  electroreduction processes reported so far, most use either complex organic electrolytes or high temperature or demonstrate electroreduction of the CO intermediate instead of directly using  $CO_2$ . There is enormous room available to explore the fundamental catalytic activity of MXenes using aqueous electrolytes for electrocatalytic  $CO_2$  conversion. In continuation of this exciting research field, herein, we have prepared the  $Ti_3C_2T_x$  MXene by an *in situ* etching method and used it as an active electrode material for electrocatalytic  $CO_2$  conversion using simple aqueous  $KHCO_3$  electrolyte. Laursen *et al.* and Hondoko *et al.* suggested that the use of Ti-based MXenes is beneficial in  $CO_2$  electroreduction due to their hydrogen-opposing behavior, which is beneficial for promising  $CO_2$  reduction.<sup>22,30</sup> The as-prepared material has been investigated using various physical and electrochemical characterization techniques. Unlike the product

formation obtained in organic electrolyte systems, we achieved the highest faradaic efficiency of 96% towards mono- as well as multi-carbon products such as CO, methanol, ethanol, and acetone in 0.5 M  $KHCO_3$  aqueous electrolyte. Moreover, the  $Ti_3C_2T_x$  MXene electrocatalyst exhibits excellent long-term stability. The use of an earth-abundant electrocatalyst for the transformation of  $CO_2$  in a benign aqueous medium into value-added products highlights the sustainable character of this methodology.

## 2. Experimental section

### 2.1. Chemicals

Potassium bicarbonate ( $KHCO_3$ ), methanol, ethanol, acetone, dimethyl sulfoxide (DMSO), lithium fluoride (LiF), and hydrochloric acid (HCl) were purchased from SRL Chemicals, India.  $Ti_3AlC_2$  was procured from Y-Carbon Ltd, Ukraine. All the chemicals were used without any further purification. A Nafion N-117 membrane was procured from Alfa Aaser, India. Deionized (DI) water with a resistivity of  $18.2 \text{ M}\Omega \text{ cm}$  was obtained from a Millipore ultrapure water purification system.

### 2.2. Preparation of the MXene ( $Ti_3C_2T_x$ )

$Ti_3C_2T_x$  was prepared by a top-down approach based on previous literature.<sup>19</sup> In a typical synthesis, 1.6 g of LiF was added to a solution containing 5 mL of DI water and 15 mL of 12 M HCl in a Teflon container (HF-safe container). The as-obtained mixture was stirred at 100 rpm at  $35^\circ \text{C}$  for 5 min to obtain a homogeneous solution. Afterward,  $Ti_3AlC_2$  (1 g) was added slowly and stirred for 24 h at  $35^\circ \text{C}$ . Then, the resultant mixture was repeatedly washed in a centrifuge at 4000 rpm until the pH of the suspension reached a value more than 6. Finally, the sedimented  $Ti_3C_2T_x$  was dried at  $60^\circ \text{C}$  in a vacuum oven for 8 h.

### 2.3. Modification of the glassy carbon electrode (GCE) by $Ti_3C_2T_x$ coating

5 mg of  $Ti_3C_2T_x$  and 5 wt% of Nafion solution (10  $\mu\text{L}$ ) were taken in 1 mL of ethanol and sonicated for 15 min. The obtained  $Ti_3C_2T_x$  slurry was applied to the freshly polished glassy carbon electrode (5 mm diameter) by repeatedly coating the electrode with 10  $\mu\text{L}$  drops of the suspension. The deposition of the suspension was repeated five times. Each time the coated suspension was allowed to dry before applying the next drop of suspension.

### 2.4. Physical characterization

Powder X-ray diffraction (PXRD) was performed using a Rigaku Ultima IV X-ray diffractometer with Cu  $K\alpha$  radiation (1.54 Å) in the range of  $5^\circ$  to  $60^\circ$  (step size of  $0.02^\circ$ ). Field emission-scanning electron microscopy (FE-SEM, JEOL-7610F Plus) was used to investigate the surface morphology of the as-prepared material. The elemental composition and their distribution were examined using energy-dispersive X-ray spectroscopy (EDX) and elemental mapping, respectively. The Raman spectrum was recorded using a LabRAM HR Evolution spectrometer



in the wavenumber range of 100–2000  $\text{cm}^{-1}$  with a laser wavelength of 532 nm. The surface area and textural characteristics of the MXene were evaluated using a Quantachrome gas sorption analyzer (Autosorb iQ3) under a liquid nitrogen atmosphere. The surface area was calculated using the Brunauer–Emmett–Teller (BET) method in the relative pressure ( $P/P_0$ ) region of 0.05–0.35, and the pore size was calculated using the Barrett–Joyner–Halenda (BJH) method from the desorption region of the isotherm at a relative pressure of  $\sim 0.99$ .

$\text{CO}_2$  adsorption–desorption experiments were carried out on a Quantachrome gas sorption analyzer at two different temperatures (288 and 298 K) in the relative pressure region of 0.05–1. Prior to  $\text{CO}_2$  adsorption experiments, the sample was degassed at 300 °C under a high vacuum for 12 h.

The adsorption affinity of  $\text{CO}_2$  on the MXene adsorbent was calculated from isosteric heat of adsorption ( $Q_{\text{st}}$ ) using the Clausius–Clapeyron equation:<sup>31</sup>

$$\ln\left(\frac{P_2}{P_1}\right) = \frac{Q_{\text{st}}}{R} \left(\frac{1}{T_2} - \frac{1}{T_1}\right) \quad (1)$$

where  $P_2$  and  $P_1$  are the relative pressures of  $\text{CO}_2$  at temperatures  $T_2$  (298 K) and  $T_1$  (288 K) for the same amount of  $\text{CO}_2$  adsorption,  $Q_{\text{st}}$  ( $\text{kJ mol}^{-1}$ ) is the enthalpy of adsorption, and  $R$  is the gas constant.

The  $\text{CO}_2/\text{N}_2$  selectivity was calculated at 298 K using the Henry constant ( $K_{\text{H}}$ ) equation, which is given as follows:<sup>31</sup>

$$S\left(\frac{\text{CO}_2}{\text{N}_2}\right) = \frac{K_{\text{H}}(\text{CO}_2)}{K_{\text{H}}(\text{N}_2)} \quad (2)$$

where  $S$  is the selectivity of  $\text{CO}_2$  over  $\text{N}_2$ , and  $K_{\text{H}}(\text{CO}_2)$  and  $K_{\text{H}}(\text{N}_2)$  are the Henry constants calculated from the slope of linearly fitted  $\text{CO}_2$  and  $\text{N}_2$  adsorption isotherms at 298 K, respectively.

### 2.5. Electrochemical characterization

Electrochemical experiments were performed using an Autolab 204 potentiostat/galvanostat (NOVA-2.1.4 software). The electrocatalytic activity of the as-prepared electrode material was investigated using linear sweep voltammetry (LSV) and chronoamperometry (CA) techniques. Platinum (Pt) and silver/silver chloride (Ag/AgCl (3 M KCl)) electrodes were used as the counter and reference electrodes, respectively. The  $\text{Ti}_3\text{C}_2\text{T}_x$  coated GCE ( $\text{Ti}_3\text{C}_2\text{T}_x$ -GCE) was used as the working electrode. LSV was performed in the  $\text{N}_2$  saturated ( $\text{N}_2$  bubbling for 30 min)  $\text{KHCO}_3$  (0.1 M or 0.5 M) electrolyte, followed by  $\text{CO}_2$  saturated electrolyte ( $\text{CO}_2$  bubbling for 30 min), using modified electrodes. The LSV was performed in the potential range between 0 and  $-1.6$  V (vs. Ag/AgCl) with a  $10 \text{ mV s}^{-1}$  scan rate.

Long-term  $\text{CO}_2$  electrolysis was performed by chronoamperometry (CA) using an H-type cell. The H-type cell comprised two compartments (anodic and cathodic), which were separated by a proton exchange membrane (Nafion-117 membrane), which acted as a separator and an ion exchanger. Each compartment was filled with 25 mL of  $\text{KHCO}_3$  solution of appropriate concentration (0.1 and 0.5 M). The counter electrode was placed in the anodic compartment, while the reference and working electrodes were placed in the cathodic compartment.

The schematic representation of the H-type cell is shown in Fig. S1 (ESI†). Before analysis, the cathodic compartment part was saturated with  $\text{N}_2$ , followed by  $\text{CO}_2$  gas for 30 min. Long-term  $\text{CO}_2$  electrolysis was carried out for 180 min with continuous bubbling of  $\text{CO}_2$  gas at a constant flow rate (10 sccm) throughout the process. The resultant liquid electroreduction products were analyzed by  $^1\text{H}$  NMR spectroscopy with the water suppression method using an AVANCE III 400 Ascend Bruker BioSpin spectrometer. The gaseous products were analyzed using a Thermo Fisher Trace 1110 with a TCD detector. The NMR samples were prepared by mixing 0.5 mL of cathodic part solution, 100  $\mu\text{L}$  of  $\text{D}_2\text{O}$ , and 0.05  $\mu\text{L}$  of DMSO (as the internal standard).

The faradaic efficiencies (FEs) of the products were calculated using the following equation:

$$\text{FE} (\%) = \frac{\text{Total output}}{\text{Total input}} \times 100 \quad (3)$$

$$\text{Total output} = M_{\text{product}} \times \text{number of electrons} (n)$$

$$\text{Total input} = \frac{Q}{F}$$

where  $M_{\text{product}}$  is the number of moles of product,  $n$  is the number of electrons involved in the electrode reaction,  $Q$  is the charge (C), and  $F$  is the Faraday constant ( $96485 \text{ C mol}^{-1}$ ).

## 3. Results and discussion

Synthesis of  $\text{Ti}_3\text{C}_2\text{T}_x$  was achieved by selective *in situ* etching of the Al layer from its MAX phase  $\text{Ti}_3\text{AlC}_2$  in the presence of the LiF and HCl etchant which led to an enhanced interlayer distance in  $\text{Ti}_3\text{C}_2\text{T}_x$  compared to parent  $\text{Ti}_3\text{AlC}_2$ .<sup>19</sup> The phase purity and the change in the interlayer distance were confirmed by the PXRD pattern (Fig. S2, ESI†). The sharp diffraction peak at around  $6^\circ$  (002) shows a characteristic peak of ordered  $\text{Ti}_3\text{C}_2\text{T}_x$  sheets. Compared to  $\text{Ti}_3\text{AlC}_2$ , a shift of the (002) peak towards a lower angle in  $\text{Ti}_3\text{C}_2\text{T}_x$  suggests the successful etching of the Al layer in the MAX phase, resulting in an increased interlayer  $d$ -spacing. Additionally, the diffraction peaks of (103), (104), (105), (106), (108), and (109) correspond to the TiC impurities.<sup>32,33</sup>

Further, the formation of the  $\text{Ti}_3\text{C}_2\text{T}_x$  MXene was also confirmed by Raman spectroscopy, as shown in Fig. 1(a). The low wavenumber peaks at 200, 360, 610, and  $720 \text{ cm}^{-1}$  are the characteristic peaks of  $\text{Ti}_3\text{C}_2\text{T}_x$ , which match well with the reported literature.<sup>32,34</sup> Besides, the two indistinguishable peaks at 1345 and  $1564 \text{ cm}^{-1}$  are assigned to the D and G bands of carbon. The indiscernible nature of these peaks indicates less disorder in the arrangement of the nanosheets. Moreover, no peaks were identified related to anatase  $\text{TiO}_2$ , confirming the absence of any anatase phase during MXene synthesis.

The morphology of the  $\text{Ti}_3\text{C}_2\text{T}_x$  has been investigated using field emission scanning electron microscopy (FE-SEM) and is



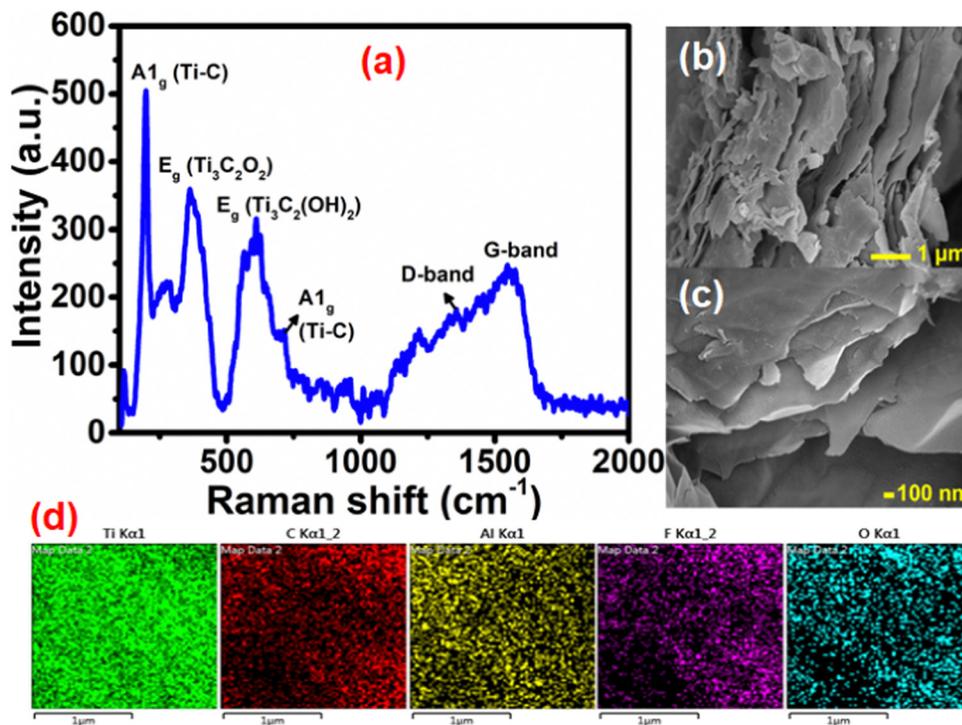


Fig. 1 (a) Raman spectrum, (b) and (c) FE-SEM images at low and high magnifications and (d) elemental mapping of  $\text{Ti}_3\text{C}_2\text{T}_x$ .

shown in Fig. 1(b) and (c). The FE-SEM images confirm the presence of a well-separated layered ultra-thin sheet-like structure of  $\text{Ti}_3\text{C}_2\text{T}_x$ . The percentage element present in  $\text{Ti}_3\text{C}_2\text{T}_x$  was studied (Fig. S3, ESI<sup>†</sup>) by energy-dispersive X-ray spectroscopy (EDX) analysis. It shows the presence of Ti and C along with surface functional group elements O and F. The presence of only a trace amount of Al (2.5 wt%) confirms the successful etching of the Al layer from the parent  $\text{Ti}_3\text{AlC}_2$ . SEM-elemental mapping (Fig. 1(d)) suggests a uniform distribution of all the elements present in the  $\text{Ti}_3\text{C}_2\text{T}_x$  MXene nanosheets.

The surface textural properties, such as the surface area and pore size characteristics of  $\text{Ti}_3\text{C}_2\text{T}_x$ , were examined by the  $\text{N}_2$  adsorption-desorption experiment. As per the IUPAC classification, the  $\text{N}_2$  adsorption-desorption isotherm (Fig. 2(a)) exhibits a type IV behavior with an  $\text{H}_3$  hysteresis loop, suggesting its mesoporous nature with uniform pore size distribution.<sup>35</sup> The Brunauer-Emmett-Teller (BET) surface area of  $\text{Ti}_3\text{C}_2\text{T}_x$  was calculated to be  $29.63 \text{ m}^2 \text{ g}^{-1}$ . Further, the Barrett-Joyner-Halenda (BJH) pore size distribution curves (Fig. 2(b)) show the narrow pore size distribution centered at 3.8 nm.

### 3.1. $\text{CO}_2$ adsorption studies

Prior to the investigation of the electrocatalytic activity of  $\text{Ti}_3\text{C}_2\text{T}_x$ , its  $\text{CO}_2$  affinity was assessed by studying the  $\text{CO}_2$  adsorption isotherm. The  $\text{CO}_2$  adsorption isotherm of  $\text{Ti}_3\text{C}_2\text{T}_x$  was investigated at two different temperatures of 288 and 298 K at a relative pressure of 1 bar (Fig. 2(c) and (d)). Fig. 2(c) depicts the adsorption-desorption isotherms at 288 and 298 K, which show maximum  $\text{CO}_2$  adsorption capacities of 0.23 and 0.16  $\text{mmol g}^{-1}$ , respectively (Fig. 2(d)). It is interesting to

observe that, at both temperatures, a significant proportion of adsorbed  $\text{CO}_2$  (*ca.* 48.9% at 288 K and 57.4% at 298 K) remains bound to the surface of  $\text{Ti}_3\text{C}_2\text{T}_x$  even after returning to zero partial pressure of  $\text{CO}_2$ . This corroborates that the  $\text{Ti}_3\text{C}_2\text{T}_x$  surface exhibits good binding affinity towards  $\text{CO}_2$ , which is an essential criterion for  $\text{CO}_2$  electroreduction. This affinity may be attributed to the strong interactions of  $\text{CO}_2$  with the existing  $-\text{OH}$ ,  $-\text{O}$  and  $-\text{F}$  surface functionalities of  $\text{Ti}_3\text{C}_2\text{T}_x$ .<sup>25</sup> To further support this observation, the FT-IR spectra of vacuum-degassed MXene and MXene exposed to  $\text{CO}_2$  at high pressure were compared to see if  $\text{CO}_2$  stays on the MXene surface after the exposure (Fig. S4, ESI<sup>†</sup>). The  $\text{Ti}_3\text{C}_2\text{T}_x$  before  $\text{CO}_2$  adsorption shows peaks at 661, 1028, and  $2012\text{--}3870 \text{ cm}^{-1}$  corresponding to the stretching vibrations of Ti-O, C-F, combined hydrogen bonded O groups, and surface hydroxyl groups.<sup>36</sup> When  $\text{Ti}_3\text{C}_2\text{T}_x$  is exposed to  $\text{CO}_2$ , the Ti-O units on the surface of  $\text{Ti}_3\text{C}_2\text{T}_x$  act as good interaction sites to facilitate the adsorption of  $\text{CO}_2$  in mono- and bidentate carbonate forms (Scheme 1).<sup>37</sup> The appearance of a new peak at  $1611 \text{ cm}^{-1}$  suggests the formation of bidentate carbonate species, while the splitting of the Ti-O-Ti peak into two separate peaks indicates the formation of monodentate carbonates.<sup>37</sup>

From the two adsorption isotherms, it is evident that decreased adsorption capacity is observed at elevated temperature, which implies that physical adsorption is the main force for the interaction between  $\text{CO}_2$  and  $\text{Ti}_3\text{C}_2\text{T}_x$  at low  $\text{CO}_2$  pressure.<sup>38</sup> Also, a linearly increasing adsorption isotherm is observed with increasing relative  $\text{CO}_2$  pressure until the measured range, implying that, owing to its mesoporous nature, the surface of  $\text{Ti}_3\text{C}_2\text{T}_x$  is still not saturated with  $\text{CO}_2$ , and even more



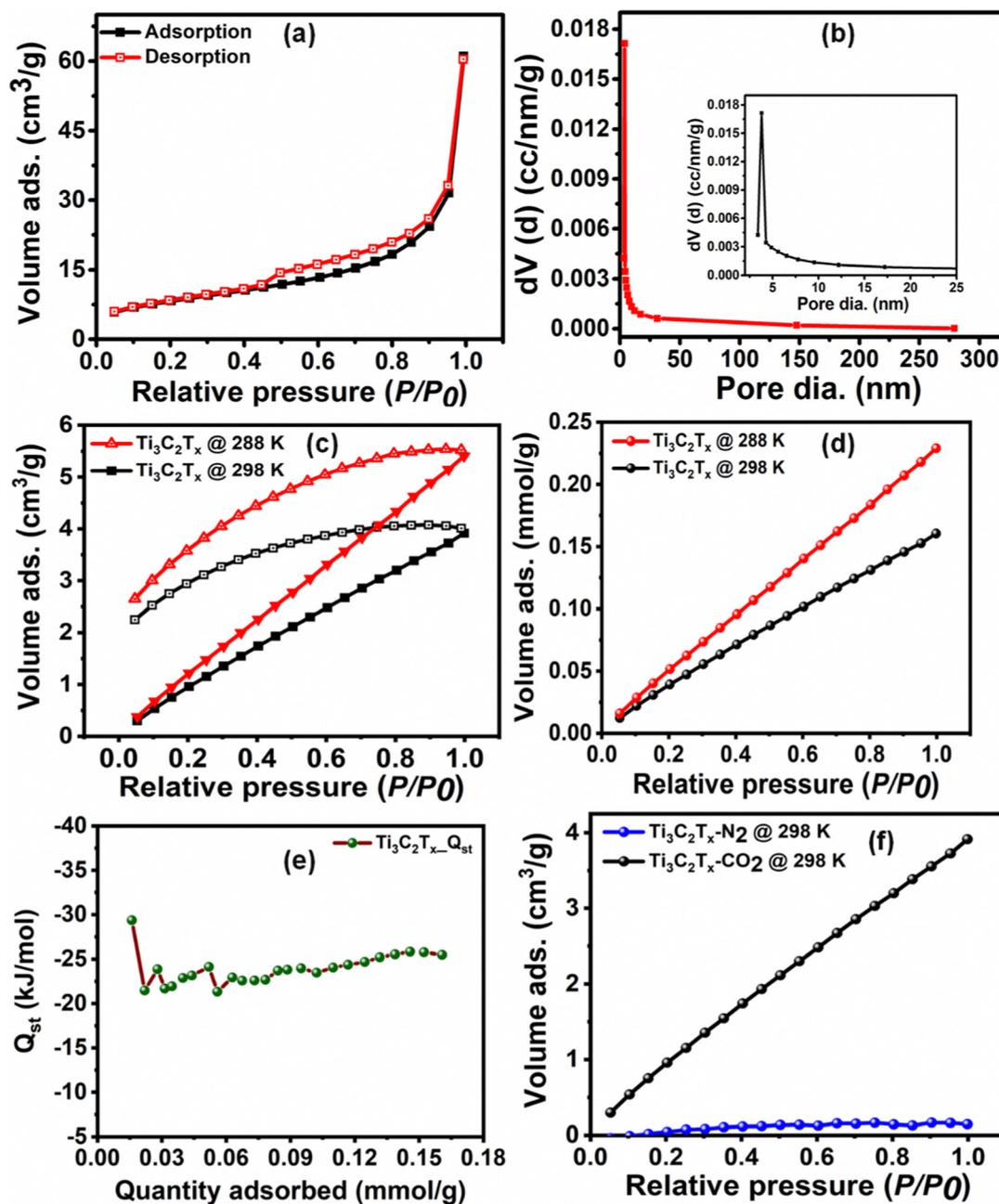
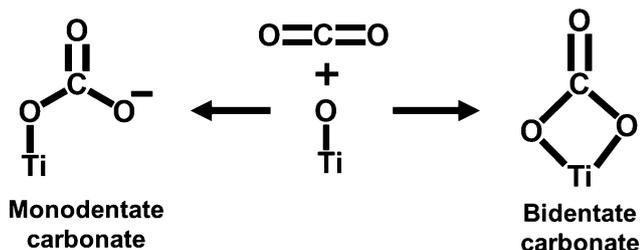


Fig. 2 (a) and (b) BET  $N_2$  adsorption–desorption isotherm and pore size distribution, (c) and (d)  $CO_2$  adsorption–desorption isotherm in  $cm^3 g^{-1}$  and  $CO_2$  adsorption isotherm in  $mmol g^{-1}$  at 288 and 298 K, (e) isosteric heat of adsorption and (f)  $CO_2$  selectivity studies over  $N_2$  at 298 K.



Scheme 1 Possible  $CO_2$  interaction on the MXene surface.

$CO_2$  adsorption can be achieved upon increasing the relative pressure.<sup>31</sup>

The adsorption affinity of  $CO_2$  on the  $Ti_3C_2T_x$  adsorbent has been calculated (Fig. 2(e)) from the  $CO_2$  adsorption isotherms at 288 and 298 K using the enthalpy of adsorption ( $-Q_{st}$ ) obtained from the Clausius–Clapeyron equation (eqn (1)). The average enthalpy of adsorption is found to be  $-23.6 \text{ kJ mol}^{-1}$ , which lies in the physisorption range ( $-10$  to  $-50 \text{ kJ mol}^{-1}$ ).<sup>39</sup> An almost constant value of the enthalpy of adsorption at all surface coverage suggests the homogeneous surface of the  $Ti_3C_2T_x$  adsorbent involving a similar type of interaction all



over its surface.<sup>38,40,41</sup> To understand the practicability of the adsorbent, the CO<sub>2</sub>/N<sub>2</sub> selectivity on Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub> has been investigated using the Henry law equation (Fig. 2(f)). The calculations indicate that the Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub> adsorbent exhibits almost 22 times selectivity for CO<sub>2</sub> over N<sub>2</sub> at room temperature, suggesting its remarkable potential as a catalyst in CO<sub>2</sub> conversion reactions. Although CO<sub>2</sub> and N<sub>2</sub> have almost similar kinetic diameters (CO<sub>2</sub>: 330 pm; N<sub>2</sub>: 364 pm), the higher electric quadrupole moment of CO<sub>2</sub> facilitates its interactions with the highly polar surface functional groups on the Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub> surface, which renders enhanced selectivity.

### 3.2. Electrochemical analysis

To evaluate the electrocatalytic activity of the Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub> modified glassy carbon electrode (GCE) (Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub>-GCE), linear sweep voltammetry (LSV) has been carried out using a standard three-electrode system in 0.5 M KHCO<sub>3</sub> aqueous electrolyte in the potential range between -1.6 and 0 V (*vs.* Ag/AgCl) at a scan rate of 10 mV s<sup>-1</sup> (Fig. 3(a)). In the N<sub>2</sub> saturated electrolyte, the obtained higher current density is attributed to the H<sub>2</sub> evolution reaction. In the case of CO<sub>2</sub>-saturated electrolyte, the current density is higher than that in the case of N<sub>2</sub>-saturated electrolyte, which is due to the simultaneous reduction of both H<sup>+</sup> and CO<sub>2</sub>. Additionally, an anodically shifted electrochemical potential has been observed for the CO<sub>2</sub>-saturated electrolyte (-1.37 V *vs.* Ag/AgCl) compared to the N<sub>2</sub>-saturated electrolyte (-1.51 V *vs.* Ag/AgCl) corresponding to a constant current density of -5 mA cm<sup>-2</sup>. These observations reveal that the electrode material active sites efficiently facilitate the electron transfer across the electrode interface and eventually help to enhance the CO<sub>2</sub> reduction process. To assess if the coated Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub> is the sole cause for this excellent electrocatalytic performance, we also investigated the catalytic activity of the bare GCE under similar conditions (Fig. 3(a)). The observed rise in the current densities for both N<sub>2</sub> and CO<sub>2</sub> saturated electrolytes is insignificant, which confirms that the overall product formation is exclusively due to the catalytic activity of Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub>.

The effect of electrolyte concentration is also an important factor for effective product formation. Therefore, we also

investigated and compared the performance of the Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub>-GCE at two different concentrations of electrolyte, namely 0.1 M and 0.5 M KHCO<sub>3</sub> (Fig. S5, ESI<sup>†</sup>). The results suggest that with increasing concentration of KHCO<sub>3</sub> electrolyte from 0.1 M to 0.5 M the current density also increases. In general, the local environment around the active materials during electrolysis is known to influence the CO<sub>2</sub> reduction reaction (CO<sub>2</sub>RR). Nakona *et al.* also observed high current density at higher KHCO<sub>3</sub> concentrations, however with decreased faradaic efficiency for CO<sub>2</sub>RR products. They suggested that, after a certain concentration of KHCO<sub>3</sub>, the K<sup>+</sup> ions generated from the electrolyte also determine the product selectivity. At higher electrolyte concentrations, the K<sup>+</sup> ions start accumulating around the negatively charged electrode materials, which hinders the CO<sub>2</sub>RR by preventing CO<sub>2</sub> from accessing the electrode, resulting in the dominance of the HER.<sup>42,43</sup> Moreover, being a basic salt (formed from a strong base and a weak acid), the higher concentration of KHCO<sub>3</sub> increases the pH of the solution. It has been suggested that > 8 pH of electrolyte causes deprotonation of the CO<sub>2</sub> carrier HCO<sub>3</sub><sup>-</sup> into CO<sub>3</sub><sup>2-</sup> and, thus, decreases the CO<sub>2</sub> carrier concentration.<sup>43,44</sup> This also favors the HER and decreases the faradaic efficiency of CO<sub>2</sub>RR products, despite the increase in the overall current density. Considering these facts, we chose a 0.5 M KHCO<sub>3</sub> concentration (causing pH ~ 7.47, just below 8; Fig. S6, ESI<sup>†</sup>) for further studies.

To examine the types of products formed during the CO<sub>2</sub> electroreduction, constant potential electrolysis, *i.e.*, chronoamperometry (CA), has been carried out in an H-type cell using a Nafion membrane as the proton exchanger and separator (Fig. S1, ESI<sup>†</sup>). Prior to the extensive investigation, we also investigated the effect of electrolyte concentration using the CA experiment at -1.1 V (*vs.* Ag/AgCl) at two different electrolyte concentrations (0.1 M and 0.5 M KHCO<sub>3</sub>). The evaluation of CO<sub>2</sub>RR products formed at two different electrolyte concentrations was done by <sup>1</sup>H NMR (Fig. S7, ESI<sup>†</sup>). Fig. S7 (ESI<sup>†</sup>) clearly suggests that the product formation in 0.1 M KHCO<sub>3</sub> is negligible. In contrast, in the case of 0.5 M KHCO<sub>3</sub> electrolyte, well-distinguished peaks for ethanol ( $\delta$  (ppm): 3.55 (q), 1.08 (t)),

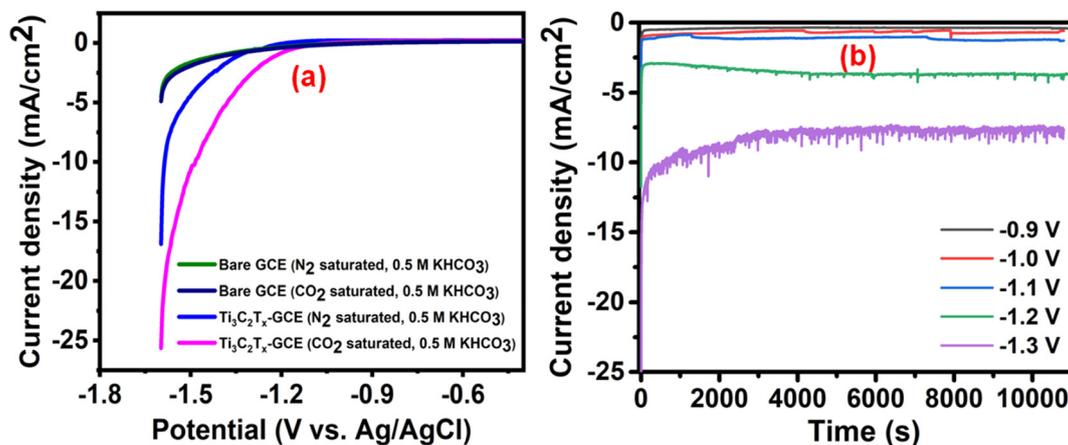


Fig. 3 (a) LSV curves in the presence of 0.5 M KHCO<sub>3</sub> at 10 mV s<sup>-1</sup> and (b) CA profiles of the Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub>-modified GCE at different potentials (*vs.* Ag/AgCl).



methanol ( $\delta$  (ppm): 3.26 (s)), and acetone ( $\delta$  (ppm): 2.13 (s)) are observable.<sup>45</sup> Therefore, choosing a 0.5 M KHCO<sub>3</sub> electrolyte concentration not only enhances the current density, as observed in the LSV experiment, but also facilitates the CO<sub>2</sub> electroreduction.

To assess the effect of potential on the CO<sub>2</sub>RR, the CA experiment was performed at different potentials (−0.9 to −1.3 V) (Fig. 3(b)). The potential choice was based on the onset potential (−1.1 V) obtained from the CO<sub>2</sub>-saturated Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub>-GCE LSV curve. The CA experiments were carried out at −0.9 V, −1.0 V, −1.1 V, −1.2 V, and −1.3 V (vs. Ag/AgCl) for 3 h, and the corresponding <sup>1</sup>H NMR spectra are shown in Fig. S8 (ESI<sup>†</sup>). The time vs. current density profiles with different potentials are shown in Fig. 3(b), which confirms the stable current densities obtained at all selected onset potentials. In the case of the CA profile at −1.3 V, some disturbance was observed, which was due to vigorous H<sub>2</sub> evolution causing bubble formation on the electrode.

Faradaic efficiency (FE) is an important measure of the catalytic activity of an electrocatalyst. The variation of the percentage FEs of the total liquid product along with the current density at different potentials is shown in Fig. 4(a). The concentrations of various products formed during the electrolysis were quantified from the standard plot obtained by plotting their relative <sup>1</sup>H NMR peak with respect to known

concentrations (Fig. S9, ESI<sup>†</sup>). From Fig. 4(a), it can be observed that, upon increasing the potential, the %FE of liquid products as well as current density gradually increases until −1.1 V (vs. Ag/AgCl) (maximum FE obtained: 53.8%), whereas a further increase in the potential causes a drastic reduction in %FE along with a sudden increase in the current density. This observation corroborates that, beyond −1.1 V, the HER is strongly favored over the CO<sub>2</sub>RR. The breakup of faradaic efficiency for individual liquid products at different voltages is shown in Fig. 4(b). It was observed that, for the highest obtained FE of 53.8%, at −1.1 V (vs. Ag/AgCl), the highest contribution came from the formation of methanol, which is comparable to most of the reported catalysts (Table S1, ESI<sup>†</sup>). In addition to liquid products, gaseous products such as CO, CH<sub>4</sub>, C<sub>2</sub>H<sub>6</sub>, and C<sub>2</sub>H<sub>4</sub> are also known to be formed in the CO<sub>2</sub>RR. Therefore, to calculate the overall %FE, we also quantified the gaseous products at −1.1 V vs. Ag/AgCl using gas chromatography (Fig. S10, ESI<sup>†</sup>), which suggested the formation of only CO (42.2%). Based on the NMR and gas chromatography studies, the overall FE of the Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub> electrocatalyst, at −1.1 V vs. Ag/AgCl, was calculated to be 96%.

In general, the Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub> electrode was observed to favor the formation of all C<sub>1–3</sub> products, suggesting a strong interaction of its surface with the CO<sub>2</sub>RR intermediates.<sup>46–48</sup> This is mainly due to the presence of −OH, −O and −F surface functional

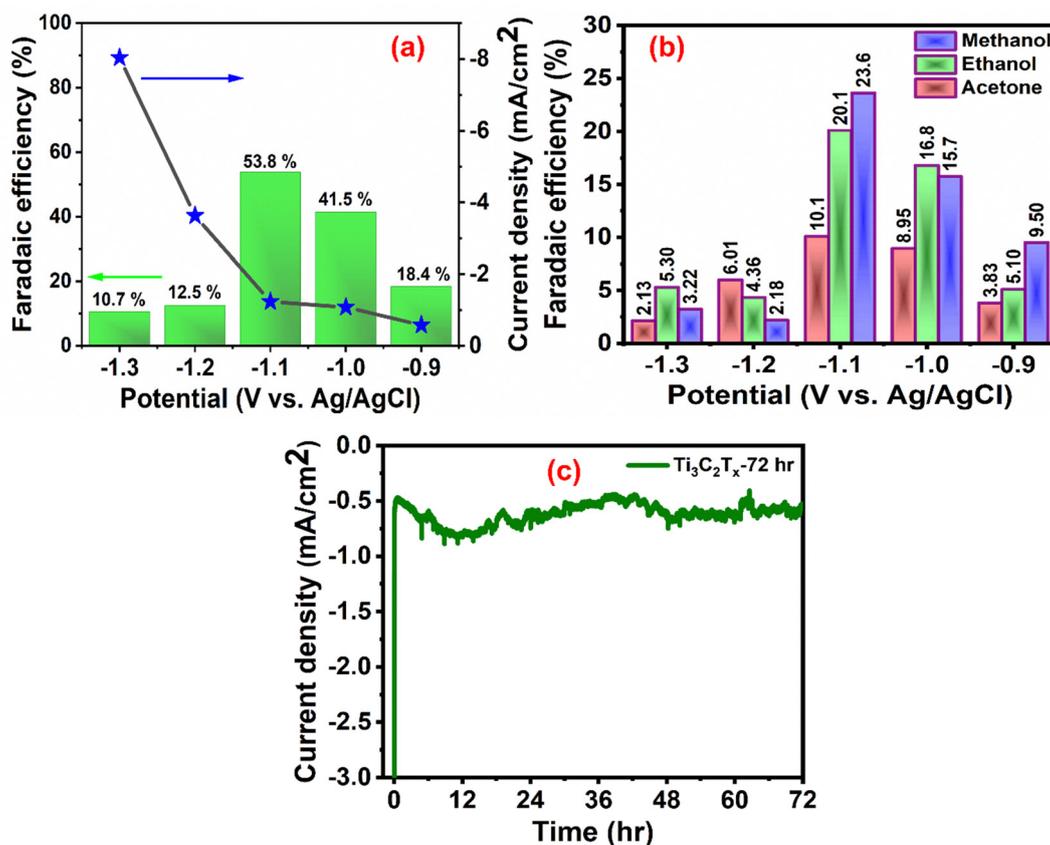


Fig. 4 (a) The total liquid product %FE and the corresponding current density, (b) individual liquid product %FE, and (c) long-term stability test at −1.1 V vs. Ag/AgCl for 72 h.



groups, which activate the electrode surface for CO<sub>2</sub> reduction.<sup>22,46</sup> It was further supplemented by calculating the electrochemical active surface area (ECSA) of Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub> in terms of double layer capacitance (*C<sub>dl</sub>*) from its cyclic voltammetry curves at different scan rates (10–50 mV s<sup>-1</sup>).<sup>49</sup> Fig. S11 (ESI†) shows the ECSA plot of Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub>, which has the maximum *C<sub>dl</sub>* value of 3.19 mF cm<sup>-2</sup>. This value is comparable to most of the recently reported electrocatalysts, confirming the presence of a higher number of electrocatalytic active sites for the excellent CO<sub>2</sub>RR performance.<sup>22,49,50</sup>

The stability of the electrocatalyst for a long-time operation is one of the important concerns for finding the best catalyst for CO<sub>2</sub> electroreduction. We investigated the long-term stability of the Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub>-GCE for 72 h at a constant potential of -1.1 V (Fig. 4(c)). After 72 h of electroreduction, no discernible change in the current density was observed, which signifies the excellent stability of the Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub>-GCE. The slight fluctuation in the chronoamperometry curve is due to bubble formation and explosion during continuous electrolysis. The same phenomena have also been observed in a previously reported investigation.<sup>29</sup> To further investigate any alteration in the structural features of the catalyst after 72 h of electrolysis, FE-SEM, EDX, elemental mapping, and PXRD studies were also carried out (Fig. S12–S14, ESI†). FE-SEM images reveal the preservation of its two-dimensional morphology with slight agglomeration. EDX and elemental mapping confirm the uniform presence of all the elements pertaining to Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub> along with electrolyte ions, confirming the overall compositional integrity of the sample after long-time electrolysis. Moreover, the PXRD pattern of the used catalysts shows all the related peaks of Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub> with reduction and broadening of some peaks, indicating partial oxidation of some Ti sites. To investigate the changes in charge transfer dynamics, electrochemical impedance spectroscopy (EIS) of the Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub> catalyst before and after the extended electrolysis was carried out (Fig. S15, ESI†). EIS measurements were performed in the frequency range of 0.1 Hz to 100 kHz with an alternating current amplitude of 10 mV. It was observed that, after prolonged electrolysis, the bulk resistance of the catalyst shows slight enhancement (*R<sub>s</sub>*: 17.2 Ω (for fresh) and 24.3 Ω (for used)), as can be observed at high frequencies. However, the appearance of a semicircle was observed for the used catalyst, which corresponds to charge transfer resistance (*R<sub>ct</sub>*) due to the presence of CO<sub>2</sub>-converted or adsorbed products (CO<sub>2</sub><sup>\*</sup>, HCO<sub>3</sub><sup>-</sup> or any liquid product) on the electrode surface which increases the impedance.<sup>51,52</sup>

## 4. Conclusion

The Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub> MXene was synthesized by a less hazardous *in situ* etching process, which was used as a selective CO<sub>2</sub> adsorbent and an electrocatalyst for CO<sub>2</sub> electroreduction. The presence of inherent functional groups, high conductivity, and metal sites in the MXene facilitates enhanced gas adsorption and endows the MXene with efficient electrocatalytic properties. Detailed volumetric CO<sub>2</sub> gas adsorption studies reveal that the MXene material exhibits a maximum adsorption capacity of 0.16 mmol g<sup>-1</sup>

at 298 K with good CO<sub>2</sub>/N<sub>2</sub> selectivity. In addition, the thermodynamic calculation suggests that the physical adsorption behavior is the main mechanism for CO<sub>2</sub> adsorption on the MXene material at low pressure. The as-prepared MXene exhibits excellent electrocatalytic properties towards CO<sub>2</sub> electroreduction in an aqueous electrolyte with an overall faradaic efficiency of 96% towards various C<sub>1</sub> (CO, CH<sub>3</sub>OH) and C<sub>2+</sub> (CH<sub>3</sub>CH<sub>2</sub>OH, CH<sub>3</sub>COCH<sub>3</sub>) products. The electrocatalyst exhibits a remarkably stable electroreduction performance for a duration of 72 hours as investigated by chronoamperometry using a laboratory-made H-type cell. These striking catalytic features of the MXene demonstrate its use as an attractive and abundantly available electrocatalyst for the conversion of CO<sub>2</sub> into multi-carbon products in a benign medium.

## Conflicts of interest

There are no conflicts to declare.

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## References

- 1 I. S. Omodolor, H. O. Otor, J. A. Andonegui, B. J. Allen and A. C. Alba-Rubio, *Ind. Eng. Chem. Res.*, 2020, **59**, 17612–17631.
- 2 P. Friedlingstein, M. O'Sullivan, M. W. Jones, R. M. Andrew, L. Gregor, J. Hauck, C. Le Quéré, I. T. Lujikx, A. Olsen, G. P. Peters, W. Peters, J. Pongratz, C. Schwingshackl, S. Sitch, J. G. Canadell, P. Ciais, R. B. Jackson, S. R. Alin, R. Alkama, A. Arneeth, V. K. Arora, N. R. Bates, M. Becker, N. Bellouin, H. C. Bittig, L. Bopp, F. Chevallier, L. P. Chini, M. Cronin, W. Evans, S. Falk, R. A. Feely, T. Gasser, M. Gehlen, T. Gkritzalis, L. Gloege, G. Grassi, N. Gruber, Ö. Gürses, I. Harris, M. Hefner, R. A. Houghton, G. C. Hurtt, Y. Iida, T. Ilyina, A. K. Jain, A. Jersild, K. Kadono, E. Kato, D. Kennedy, K. Klein Goldewijk, J. Knauer, J. I. Korsbakken, P. Landschützer, N. Lefèvre, K. Lindsay, J. Liu, Z. Liu, G. Marland, N. Mayot, M. J. McGrath, N. Metz, N. M. Monacci, D. R. Munro, S.-I. Nakaoka, Y. Niwa, K. O'Brien, T. Ono, P. I. Palmer, N. Pan, D. Pierrot, K. Pockock, B. Poulter, L. Resplandy, E. Robertson, C. Rödenbeck, C. Rodriguez, T. M. Rosan, J. Schwinger, R. Séférian, J. D. Shutler, I. Skjelvan, T. Steinhoff, Q. Sun, A. J. Sutton, C. Sweeney, S. Takao, T. Tanhua, P. P. Tans, X. Tian, H. Tian, B. Tilbrook, H. Tsujino, F. Tubiello, G. R. van der Werf, A. P. Walker, R. Wanninkhof, C. Whitehead,



- A. Willstrand Wranne, R. Wright, W. Yuan, C. Yue, X. Yue, S. Zaehle, J. Zeng and B. Zheng, *Earth Syst. Sci. Data*, 2022, **14**, 4811–4900.
- 3 A. Velty and A. Corma, *Chem. Soc. Rev.*, 2023, 150–156.
- 4 D. Gao, R. M. Arán-Ais, H. S. Jeon and B. Roldan Cuenya, *Nat. Catal.*, 2019, **2**, 198–210.
- 5 S. Garg, M. Li, A. Z. Weber, L. Ge, L. Li, V. Rudolph, G. Wang and T. E. Rufford, *J. Mater. Chem. A*, 2020, **8**, 1511–1544.
- 6 L. Fan, C. Xia, F. Yang, J. Wang, H. Wang and Y. Lu, *Sci. Adv.*, 2020, **6**, 1–18.
- 7 Q. Fan, M. Zhang, M. Jia, S. Liu, J. Qiu and Z. Sun, *Mater. Today Energy*, 2018, **10**, 280–301.
- 8 P. P. Yang, X. L. Zhang, F. Y. Gao, Y. R. Zheng, Z. Z. Niu, X. Yu, R. Liu, Z. Z. Wu, S. Qin, L. P. Chi, Y. Duan, T. Ma, X. S. Zheng, J. F. Zhu, H. J. Wang, M. R. Gao and S. H. Yu, *J. Am. Chem. Soc.*, 2020, **142**, 6400–6408.
- 9 Q. Lei, H. Zhu, K. Song, N. Wei, L. Liu, D. Zhang, J. Yin, X. Dong, K. Yao, N. Wang, X. Li, B. Davaasuren, J. Wang and Y. Han, *J. Am. Chem. Soc.*, 2020, **142**, 4213–4222.
- 10 C. Jia, K. Ching, P. V. Kumar, C. Zhao, N. Kumar, X. Chen and B. Das, *ACS Appl. Mater. Interfaces*, 2020, **12**, 41288–41293.
- 11 J. Miao, Q. Zhu, K. Li, P. Zhang, Q. Zhao and B. Xu, *J. Energy Chem.*, 2021, **52**, 243–250.
- 12 D. Gan, Q. Huang, J. Dou, H. Huang, J. Chen, M. Liu, Y. Wen, Z. Yang, X. Zhang and Y. Wei, *Appl. Surf. Sci.*, 2020, **504**, 144603.
- 13 M. K. Singh, S. Krishnan and D. K. Rai, *Electrochim. Acta*, 2023, **441**, 141825.
- 14 Y. Zhang, Z. Cao, S. Liu, Z. Du, Y. Cui, J. Gu, Y. Shi, B. Li and S. Yang, *Adv. Energy Mater.*, 2022, **12**, 2103979.
- 15 V. Kedambaimoole, K. Harsh, K. Rajanna, P. Sen, M. M. Nayak and S. Kumar, *Mater. Adv.*, 2022, **3**, 3784.
- 16 B. M. Abraham, V. Parey and J. K. Singh, *J. Mater. Chem. C*, 2022, **10**, 4096–4123.
- 17 H. Wang, Y. Wu, X. Yuan, G. Zeng, J. Zhou, X. Wang and J. W. Chew, *Adv. Mater.*, 2018, **30**, 1–28.
- 18 J. Pang, R. G. Mendes, A. Bachmatiuk, L. Zhao, H. Q. Ta, T. Gemming, H. Liu, Z. Liu and M. H. Rummeli, *Chem. Soc. Rev.*, 2019, **48**, 72–133.
- 19 M. Alhabeab, K. Maleski, B. Anasori, P. Lelyukh, L. Clark, S. Sin and Y. Gogotsi, *Chem. Mater.*, 2017, **29**, 7633–7644.
- 20 N. H. Attanayake, H. R. Banjade, A. C. Thenuwara, B. Anasori, Q. Yan and D. R. Strongin, *Chem. Commun.*, 2021, 57, 1675–1678.
- 21 X. Zhang, Z. Zhang, J. Li, X. Zhao, D. Wu and Z. Zhou, *J. Mater. Chem. A*, 2017, **5**, 12899–12903.
- 22 A. D. Handoko, H. Chen, Y. Lum, Q. Zhang, B. Anasori and Z. W. Seh, *iScience*, 2020, **23**, 101181.
- 23 I. Persson, J. Halim, H. Lind, T. W. Hansen, J. B. Wagner, L.-Å. Näslund, V. Darakchieva, J. Palisaitis, J. Rosen and P. O. Å. Persson, *Adv. Mater.*, 2019, **31**, 1805472.
- 24 Á. Morales-García, A. Fernández-Fernández, F. Viñes and F. Illas, *J. Mater. Chem. A*, 2018, **6**, 3381–3385.
- 25 B. Wang, A. Zhou, F. Liu, J. Cao, L. Wang and Q. Hu, *J. Adv. Ceram.*, 2018, **7**, 237–245.
- 26 N. Li, X. Chen, W. J. Ong, D. R. Macfarlane, X. Zhao, A. K. Cheetham and C. Sun, *ACS Nano*, 2017, **11**, 10825–10833.
- 27 A. D. Handoko, K. H. Khoo, T. L. Tan, H. Jin and Z. W. Seh, *J. Mater. Chem. A*, 2018, **6**, 21885–21890.
- 28 Q. Zhao, C. Zhang, R. Hu, Z. Du, J. Gu, Y. Cui, X. Chen, W. Xu, Z. Cheng, S. Li, B. Li, Y. Liu, W. Chen, C. Liu, J. Shang, L. Song and S. Yang, *ACS Nano*, 2021, **15**, 4927–4936.
- 29 H. Bao, Y. Qiu, X. Peng, J. A. Wang, Y. Mi, S. Zhao, X. Liu, Y. Liu, R. Cao, L. Zhuo, J. Ren, J. Sun, J. Luo and X. Sun, *Nat. Commun.*, 2021, **12**, 1–9.
- 30 A. B. Laursen, A. S. Varela, F. Dionigi, H. Fanchiu, C. Miller, O. L. Trinhammer, J. Rossmeisl and S. Dahl, *J. Chem. Educ.*, 2012, **89**, 1595–1599.
- 31 N. Guha, A. K. Gupta, S. Chatterjee, S. Krishnan, M. K. Singh and D. K. Rai, *J. CO<sub>2</sub> Util.*, 2021, **49**, 101575.
- 32 C. Zhao, Q. Wang, H. Zhang, S. Passerini and X. Qian, *ACS Appl. Mater. Interfaces*, 2016, **8**, 15661–15667.
- 33 B. Aïssa, A. Ali, K. A. Mahmoud, T. Haddad and M. Nedil, *Appl. Phys. Lett.*, 2016, **109**, 043109.
- 34 A. Sarycheva and Y. Gogotsi, *Chem. Mater.*, 2020, **32**, 3480–3488.
- 35 S. Krishnan, S. Chatterjee, A. Solanki, N. Guha, M. K. Singh, A. K. Gupta and D. K. Rai, *ACS Appl. Nano Mater.*, 2020, **3**, 11203–11216.
- 36 E. Lee, A. Vahidmohammadi, B. C. Prorok, Y. S. Yoon, M. Beidaghi and D. J. Kim, *ACS Appl. Mater. Interfaces*, 2017, **9**, 37184–37190.
- 37 L. Mino, G. Spoto and A. M. Ferrari, *J. Phys. Chem. C*, 2014, **118**, 25016–25026.
- 38 G. Singh, K. S. Lakhi, I. Y. Kim, S. Kim, P. Srivastava, R. Naidu and A. Vinu, *ACS Appl. Mater. Interfaces*, 2017, **9**, 29782–29793.
- 39 M. Senthilkumaran, C. Saravanan, P. Puthiaraj, P. Rameshkumar, G. P. Kalaigan and P. Muthu Mareeswaran, *Mater. Chem. Phys.*, 2020, **256**, 123750.
- 40 C. Chen, B. Li, L. Zhou, Z. Xia, N. Feng, J. Ding, L. Wang, H. Wan and G. Guan, *ACS Appl. Mater. Interfaces*, 2017, **9**, 23060–23071.
- 41 N. Guha, S. Krishnan, S. Gupta and D. K. Rai, *ChemNanoMat*, 2023, **9**, e202200519.
- 42 H. Zhong, K. Fujii and Y. Nakano, *J. Electrochem. Soc.*, 2017, **164**, F923–F927.
- 43 M. König, J. Vaes, E. Klemm and D. Pant, *iScience*, 2019, **19**, 135–160.
- 44 H. Zhong, K. Fujii, Y. Nakano and F. Jin, *J. Phys. Chem. C*, 2015, **119**, 55–61.
- 45 T. Chatterjee, E. Boutin and M. Robert, *Dalton Trans.*, 2020, **49**, 4257–4265.
- 46 K. Eid, Q. Lu, S. Abdel-Azeim, A. Soliman, A. M. Abdullah, A. M. Abdelgwad, R. P. Forbes, K. I. Ozoemena, R. S. Varma and M. F. Shibl, *J. Mater. Chem. A*, 2022, **10**, 1965–1975.
- 47 Y. Xiao and W. Zhang, *Nanoscale*, 2020, **12**, 7660–7673.
- 48 X. Qian, L. Li, Y. Li, Z. Liu, Z. Tian, C. Zhan and L. Chen, *Phys. Chem. Chem. Phys.*, 2021, **23**, 12431–12438.



- 49 Z. Miao, W. Liu, Y. Zhao, F. Wang, J. Meng, M. Liang, X. Wu, J. Zhao, S. Zhuo and J. Zhou, *Catal. Sci. Technol.*, 2020, **10**, 967–977.
- 50 Z. Yan, X. Wang, Y. Tan, A. Liu, F. Luo, M. Zhang, L. Zeng and Y. Zhang, *Catal. Sci. Technol.*, 2021, **11**, 6742–6749.
- 51 F. Bienen, D. Kopljar, A. Löwe, S. Geiger, N. Wagner, E. Klemm and K. A. Friedrich, *ACS Sustainable Chem. Eng.*, 2020, **8**, 13759–13768.
- 52 F. Bienen, D. Kopljar, S. Geiger, N. Wagner and K. A. Friedrich, *ACS Sustainable Chem. Eng.*, 2020, **8**, 5192–5199.

