MXene Based Sensing Materials: Current Status and Future Perspectives

Vishnu Sankar Sivasankarapillai,† Tata Sanjay Kanna Sharma,‡ ³ Kuo-Yuan Hwa³ ³ Saikh Mohammad Wabaidur,⁴ Subramania Angaiah⁵ and Ragupathy Dhanusuraman¹

Abstract

MXenes are a family of two-dimensional multifunctional materials in a rapid development phase since their discovery in 2011. MXenes possess excellent features like high conductivity and surface area, improved mechanical properties, hydrophilicity, and ability to tune the surface properties with modifying functional groups. These features make MXenes suitable candidates for a wide range of applications, including biomedical and energy storage. This review is focused on various types of MXenes which are recently reported for sensor applications. The current scenario regarding fabrication and properties of MXenes are initially described, followed by a discussion of their applications as piezoresistive and biochemical sensors. This involves mechanical strain detection and detection of biomolecules, biomarkers, and drug molecules relevant to biomedical applications. Finally, the future perspectives are briefly discussed, which will help the researchers identify the limitations for the current scenario and develop new strategies that focus on developing novel, efficient and sensitive MXene based sensors.

Keywords: MXene; Biosensors; Energy storage; Multifunctional materials; Biomedical applications.

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1. Introduction

Two dimensional materials have emerged as a platform of enormous possibilities in material science due to their distinctive electronic, mechanical, chemical, and optical properties. These materials are composed either of a single element such as Silicene,¹ Graphene,² Germanene,³ and Borophene⁴ or two elements such as dichalcogenides.³ MXenes are a relatively new candidate of this family, first discovered in 2011 by introducing titanium carbide (Ti₃C₂). The synthesis method used Ti₃AlC₂ as the precursor in which the selective etching of Al atoms using hydrofluoric acid (HF) under ambient conditions giving Ti₃C₂ MXene as represented in Fig. 1.⁶

MXenes are derivatives of the three dimensional MAX phases as the parent structure and have the general formula Mₙ₊₁AXₙ.⁷ M represents member from early transition metal family like scandium, titanium, vanadium, chromium, zirconium, niobium, molybdenum, hafnium, tantalum, tungsten (Sc,Ti, V, Cr, Zr, Nb, Mo, Hf, Ta, W respectively), A indicates an element from either group 13 or group 14, and X can be either carbon or nitrogen. Here n is an integer ranging from 1 to 3. MAX phase is a unique structure in which closely packed multilayers are present which are composed of alternating layers of M and A. M-X bonds are strong compared to weak M-A bonds. MXenes are found to be superior to their parent MAX phases in terms of electronic, magnetic, optical, and electrochemical properties. The special feature of combined metallic conductivity of transition metal carbides/carbides along with the presence of hydrophilic terminated surface make MXenes as a superior candidate among other two dimensional materials.⁹

1.1 Synthesis of MXenes

Fabrication routes of MXenes are primarily focused on the removal of the more reactive “A” atoms from Mₙ₊₁Xₙ layers
without any structural alterations. This feature is attributed to the relative weakness of M-A bonds compared to the M-X bonds, which is a stronger metallic covalent bond.[6] MAX phases are tightly wired to each other by strong van der Waals interaction, which complicates exfoliation more than similar 2D materials like Graphene, Phosphorene etc. Literature suggests that around twenty MXenes are successfully produced to date from more than 70 types of MAX phases. The dissolving and etching of the A layers require aggressive acidic conditions, and hydrofluoric acid (HF) is conventionally employed. Fig. 2 represents the milestones in the history of the development of MXenes. Naguib et al. were the first group of researchers who successfully prepared MXene through etching the A layer from Ti$_3$AlC$_2$ using hydrofluoric acid (HF).[6] Due to the hazardous nature of HF, researchers are focusing on developing novel alternative methods to prepare MXenes, giving the product in good yield.[9] Here we discuss some feasible and effective methods for fabricating MXenes available in literature over the last decade.

**Fig. 1** Fabrication strategy of MXene by HF etching. Reproduced with the permission from [6], Copyright 2012 American Chemical Society.

Yang et al. developed an efficient etching technique based on anodic corrosion of Ti$_3$AlC$_2$ without using fluorine in a binary aqueous electrolyte. The authors employed an in situ intercalation of NH$_4$OH after the dissolution of Al, which resulted in the high yield, both mono- and bi-layers, extraction (over 90%) of carbide flakes with upto 18.6 μm size.[10] Similarly, Sun et al. proposed a mechanism based on intercalation-alloying expansion-microexplosion process for preparing layered fluoride-free Ti$_3$C$_2$Tx (T = O or OH).[11] This rapid technique involved an etching strategy electrochemically without fluoride, leading to the Al layers being selectively corroded in titanium aluminum carbide (Ti$_3$AlC$_2$).

Malaki et al. provide a detailed account of employing ultrasonication for preparing MXene as an effective and successful approach.[12] The key process that needs to be considered is the proper dispersion and delamination of MXene flakes into layered sheets while producing MXenes since the properties of MXenes depends on the factors like layered structure, increased lateral size, reduced defects, as well as proper degree of exfoliation.

The two major difficulties encountered in the synthesis of nitride-based MXenes are higher formation energy of M$_{n+1}$AN$_n$ and compromised stability of the M$_{n+1}$N$_n$ layers employed in the etchant (conventionally HF). Soundiraraju et al. addressed this problem through the synthesis of Titanium Nitride (Ti$_3$N), where the Al atoms were selectively etched from ternary Ti$_2$AlN.[13] The powder was immersed in a mixture of potassium fluoride and hydrochloric acid to achieve the intercalation of layers. Furthermore, the few layered Ti$_2$NTx was obtained as a result of sonicating the formerly synthesized multilayered Ti$_2$NTx.

Aluminium fluoride (AlF$_3$·3H$_2$O) is considered to be an undesirable byproduct most commonly obtained from the synthesis of MXenes. Cockreham et al. deduced the conditions leading to the formation of AlF$_3$·3H$_2$O, while employing CoF$_2$/CoF$_3$ for etching technique exploration.[14] It is found that the ionic strength of the etching reagent plays a crucial role in the formation of AlF$_3$·3H$_2$O. The AlF$_3$ complexation is stable with the ionic strength ranging between ~8.5 and 10M.
Alhabebet et al. synthesized Ti$_3$C$_2$ by selectively etching silicon from titanium silicon carbide Ti$_5$SiC$_2$, which is considered the most common MAX phase.[15]

A universal strategy was developed by Pang et al., which was based on an electrochemical etching route assisted thermally, for the synthesis of MXenes (e.g., Ti$_2$CT$_x$, Cr$_2$CT$_x$, and V$_2$CT$_x$).[16] The experimental procedure is illustrated in Fig. 3. Since the strength of M–Al and M–C bonds are different, it was essential to determine the voltage for etching for individual MXene. The experiment was performed with the aid of cyclic voltammetry (CV) measurements. Ti$_5$AlC has broad CV peaks, but unlike that, the plateaus corresponding to Al removal can also be identified. Tran et al. synthesized V$_2$C$_2$Ti$_4$ based on the chemical exfoliation of the V$_2$AlC$_3$ MAX phase by treatment with aqueous HF.[17] Limbu et al. reported an ecofriendly and facile method for synthesizing reduced Ti$_5$C$_2$TX MXene (r-Ti$_5$C$_2$TX) which involved treatment with L-ascorbic acid at room temperature.[18]

![Fig. 3 Electrochemical etching route for preparing MXene for energy applications. Reproduced with the permission from [16], Copyright 2019 American Chemical Society.](image)

Urbanowski et al. demonstrated that Mo$_5$CT$_x$ and V$_2$CT$_x$ MXenes transform to two dimensional (2D) metal nitrides Mo$_2$N and V$_2$N through ammoniation at 600 °C.[19] The N atoms that are obtained on the decomposition of ammonia molecules substitute the carbon atoms in the MXene precursor. Zhang et al. proposed two methodologies for synthesizing oxidized Nb$_2$CT$_x$ MXene, the first being a hydrothermal method (h-Nb$_2$O$_5$/C), and the second was calcination of CO(Nb$_2$O$_5$/C).[20] The authors revealed that the synergistic effect by large lipophilic surface of the disordered carbon and high dielectric constant of the Nb$_2$O$_5$ nanoparticles are the reason for superior electro-rheological performance. Zhao et al. reported a facile, single-step method for synthesizing nanosheets of Niobium carbide (Nb$_2$CT$_x$) MXene by a simplified approach in which Nb$_5$AIC powder was etched.[21]

Frey et al. reported a positive and unlabelled (PU) Machine learning approach that can successfully predict which of the theoretically proposed MXenes have the highest probability of being synthesized successfully.[22] The authors revealed 18 promising MXene candidates that can be possibly synthesized with the knowledge of elemental information and results from density functional theory (DFT) calculations of 2D transition metal carbides, carbonitrides, and nitrides, and their layered precursor MAX phases.

### 1.2 MXene based sensors

The major applications of MXenes are represented in Fig. 4. The metal and the terminal surface groups of the MXene layer has a great deal of influence on the performance and the physical properties of the material.

It is noticed that the terminal surface groups influence the performance, the optical and electronic property of the material significantly, whereas the M in MXene layer affects its electronic properties. The increasing research interest for the application of MXene for fabricating electrochemical sensors is due to the following features:[5]:

1. High conductivity
2. Good electroactive surface area
3. Ease of fabrication with other materials to give composite materials
4. Excellent hydrophilicity compared to other 2D materials

These properties allow fabricating sensors for various molecules, including bioanalytes, environmental contaminants and other relevant chemical species like heavy metal ions and anionic analytes. In general, the electroanalytical performance of an analyte is often enhanced by the adsorption of the analyte at the electroactive surface. The terminal functional groups of the MXene and the pH of the analyte solution have been demonstrated to have a significant influence on the sensing action of MXene based electrochemical sensors. For instance, MXenes have been reported for a good affinity for metal cations at higher pH due to the creation of a negative charge surface via ionization of functional groups. Thus, the intercalation and adsorption in the structural aspects of MXenes are effectively exploited on fabricating MXene based electrodes for sensing applications. Zhu et al. performed the modification of Ti$_5$C$_2$T$_x$ with –OH groups to provide an alkylated MXene which was further employed for the electrochemical detection of heavy metal ions such as Cd(II), Pb(II), Cu(II) and Hg(II), simultaneously, by using square wave voltammetry.[23] In the experiment, the sensor developed showed high sensitivity with linear ranges between 0.1 μm and 1.0 μm for all the metals with a limit of detection values at 98 nm, 41 nm, 32 nm, and 130 nm, for Cu(II), Pb(II), Cu+, and Hg(II), respectively. Another sensor fabricated with Ti$_5$C$_2$T$_x$ on glassy carbon electrode was used to detect bromate (BrO$_3^-$) ion where response between the linear range of 50 nm to 5 μm was performed limit of detection (LOD) value of 41 nM was observed.[24]

Haemoglobin (Hb) and Tyrosinase are among the other enzymes reported to be used with MXenes to detect nitrite and phenol, respectively. Liu et al. developed nitrite sensor by the immobilization of Hb on Ti$_5$C$_2$T$_x$.[27] A broad wave of reduction was observed at –0.695 V, which was attributed to the nitride being converted to N$_2$O. Here, a LOD of 0.12 μM...
was achieved along with a broad linear range between 0.5 μm to 11.8 mm. Similarly, the alkylation of Ti$_3$C$_2$T$_x$ was done to modify the surface of the MXene with -OH groups that underwent partial termination, thus providing an improved biocompatible microenvironment to immobilize Tyrosinase.[28] This resulted in achieving a rapid time of response, coupled with efficient electrocatalytic reaction, which was due to the rapid phenol diffusion onto the immobilized tyrosinase conjugated on the MXene nanoflakes, from the bulk solution. The sensor exhibited an excellent sensitivity of 414 mA m$^{-1}$, with a very low LOD of 12 nm. 

Literature suggests that oxidation of Ti$_3$C$_2$T$_x$ occurs at potentials higher than about 0.45 V vs Ag|AgCl electrode, but this oxidation is not prominent when metal nanoparticle is combined with Ti$_3$C$_2$T$_x$ suggesting the stability attributed by metal nanoparticles.[29] The mechanism of stability in the presence of metal nanoparticles is not much revealed but needs significant attention. This review briefly discusses two different aspects of MXene based sensors such as piezoresistive (mechanical) sensors and electrochemical and luminescent biosensors. MXene based mechanical sensors give a wide horizon of opportunities in the biomedical field like wearable electronics,[30] therapeutic applications,[31] etc. In contrast, the biosensors cover sensors for detecting analytes such as biomolecules, biomarkers and drug molecules.[32]

2. Piezoresistive sensors
Piezoresistive pressure sensors have got a significant interest in the biomedical field owing to their ability to monitor physiological signals. Wearable electronic sensors are the perfect examples of real-time applications of these sensors.[33] Li et al. successfully developed wearable electronic pressure sensor by incorporating multilayer Ti$_3$C$_2$T$_x$ MXene into poly(vinylidene fluoride) trifluoroethylene (PVDF-TrFE) which is a hydrophobic polymer.[34] This piezoresistive pressure sensor works on the property of Ti$_3$C$_2$T$_x$ in which the resistance changes as a function of interlayer separation and stable even after twenty-week exposure to air. Similarly, Liao et al. reported a similar wearable strain sensor prepared from conductive MXene nanocomposite organohydrogel (MNOH).[35] It is prepared by the immersion of MXene nanocomposite hydrogel (MNH) into a solution of ethylene glycol (EG) and subsequent development into hydrogel polymer networks. This material possesses advantages over the typical conductive hydrogels where they use pure water as the medium of dispersion which loses its properties due to freezing at subzero temperatures. The authors observed that MNOH could detect human physiological actions with a broad strain range (up to 350% strain) and possess outstanding features such as excellent anti-freezing property (-40°C), prolonged retention of moisture (8d), outstanding capability to self-heal, and enhanced mechanical properties.

Conducting polymers were also reported to be combined with MXenes for developing similar mechanical sensors. These polymers which possess enhanced conductivity due to the conjugated carbon framework which facilitates charge transport. Due to this reason, conducting polymers have got significant interest in energy storage devices and sensors.[36,37] One example of this category of polymer is Polypyrrole which has already been well explored for its application in electrochemical sensors.[38] Seroka et al. prepared functionalized Ti$_3$AlC$_2$-carbon nanoparticle-Polypyrrole nanocomposite having a sensitivity of 0.00017 kPa$^{-1}$ and a rapid recovery time between 3s and 5s.[39] Ti$_3$AlC$_2$ is also...
reported for fabricating MXene/ink sensor which has wide application prospects in monitoring the damage of structural composites used in aerospace applications. Proper investigation of employing conducting polymers for developing pressure sensors and correlating with their electrochemical performance can give rise to multifunctional sensors. Ti$_3$C$_2$Tx/polyurethane (PU) composite fibers were also reported as components for wearable strain sensing textiles. Uzun et al. further extended the application potential of Ti$_3$C$_2$Tx by encapsulating cellulose yarns with Ti$_3$C$_2$Tx to form knitting and washable textile components. Electrospinning technique was also reported to develop 2D Ti$_3$C$_2$T$_x$ MXene based electrospun mats for piezoresistive sensing applications. Li et al. fabricated a highly conductive MXene/cellulose nanocrystal (CNC) with super flexibility and coated onto thermoplastic polyurethane (TPU) non-woven fabric (NWF). The authors used dip-coating technique for the fabrication of wearable strain/pressure sensors. The material showed a wide range of sensing (about 83%), an excellent sensitivity (GF = 3405), and a very low limit of detection (0.1%) which was primarily attributed to the opening or closing of micro-cracks present in the composite. A similar study reported the fabrication of a multilayer fiber sensor which was developed by self-assembling silver nanowire (AgNW), MXene, and waterborne polyurethane (WP) layer-by-layer assembly. This sensor proved to be highly sensitive and a wide operating range (up to 100%), along with exhibiting great stability (1000 cycles), reliability, a rapid response (344 ms) and relaxation time of 344 ms. Similarly, Li et al. developed a sensor which exhibits ultra-high sensitivity with a rapid time of response of 26 ms, and an excellent cycle stability (5600 cycles).

Li et al. successfully developed a piezoresistive sensor by treating polyurethane (PU) with chitosan (CS) to obtain positively charged CS@PU sponge. This sponge was successively coated with negatively charged Ti$_3$C$_2$T$_x$ MXene sheets. The authors demonstrated that this sensor has promising degree of compressibility and stable piezoresistive response for compressive strains of up to 85% with a stress of 245.7 kPa.

Literature is available for employing aerogels for both electrochemical and piezoresistive sensor applications. Zhao et al. employed conductive 3D-Ti$_3$C$_2$T$_x$ porous architectures to develop hybrid aerogels by combination of graphene oxide assisted hydrothermal assembly and freeze-drying. The structure of these hybrid aerogels consists of Ti$_3$C$_2$T$_x$ walls in which inner skeleton is graphene sheets, and this hybrid aerogel possess future perspectives to be developed as a sensor material. Ma et al. reported reduced graphene oxide (RGO)/Ti$_3$C$_2$T$_x$ MXene aerogel with very good sensitivity (22.56 kPa) and fast response time (~200 ms). This study revealed that RGO nanosheets could prevent the poor oxidization of MXene by enhancing the wrapping of MXene inside the aerogel. Hydrogels were also reported for developing sensors with superior performance.

Zhang et al. prepared MXenes by Yury mild method. Poly(N-iso-pracrylamide) (PNIPAM) hydrogel and physical crosslinking hydrogel were used as the matrix to prepare composite hydrogels self-healing and stress-sensing properties. The conductivity of the composite hydrogel was about fifteen times more than that of the control hydrogel devoid of MXene. Tran et al. developed a thermal sensor using poly(2-dimethylamino) ethyl methacrylate) (PDMAEMA) which is a stimuli-responsive polymer with Ti based MXene using the OH groups on the MXene surface as a linking agent for organic moieties. The authors also provide a theoretical insight into the conductivity mechanism of the material through DFT calculations. Yue et al. fabricated piezoresistive sensor with MXene-sponge network by a simple and efficient dip-coating process with insulating Polyvinyl alcohol (PVA) nanowires as spacer. The authors achieved high sensitivity for the material for a broad pressure range with a low detection limit and excellent durability over 10,000 cycles. Zhang et al. give the possibility of developing wearable sensors based on Ti$_3$C$_2$T$_x$ MXenes modified with small biomolecules to give enhanced conductivity and self-healing properties. This conductive composite consisting of 10wt% of MXenes showed an elongation of 81%, and a mechanical strength of 1.81 MPa. Yuan et al. used Ti$_3$C$_2$T$_x$ with nylon fabric to form a sensor with gauge factor up to 24.35 and 5.98 within the strain range of 20% in the axial and radial of nylon fabric respectively. Also the material possesses high durability and good fastness to washing which gives the possibility for developing for real-life physiological sensors. Muckley et al. reported that films of K- and Mg-intercalated MXenes exhibit relative humidity (RH) detection thresholds of ~0.8% RH and showed monotonic RH response in the 0-85% RH range. This suggests the good potential of developing the material into an efficient humidity sensor.

Guo et al. presents a biodegradable pressure sensor fabricated by sandwiching porous tissue paper incorporated with MXene with polyactic acid (PLA) thin sheets and PLA thin sheet. This flexible sensor demonstrated to have high sensitivity with a low detection limit (10.2 Pa) with a fast response time.

It was demonstrated that the micro-architecture of both composites plays a significant role in the performance of piezoresistive sensors. This perspective has got attention in which the researchers are trying to design sensors with unique architecture. Shi et al. provides a good example of biomimetic sensor which is based on "Brick-and-Mortar" architecture. Microscale hierarchical architecture is constructed from the combination of Ti$_3$C$_2$T$_x$ /1D silver nanowire "brick" and poly(dopamine)/Ni "mortar". The sensor worked efficiently using the loading energy to promote the stepwise propagation of cracks while stretching. Another biomimetic spider web-inspired temperature and strain sensor is reported based on Ti$_3$C$_2$T$_x$ hydrogels. Wang et al. developed this system based on 3D Ti$_3$C$_2$T$_x$-polyacrylamide (PAM) nanocomposite hydrogels inspired by spider web architecture. Sensing
properties of the material is attributed to the continuous distribution of MXene nanosheets connected by the PAM network which creates electron transport pathways and proton transport bridges in the hydrogel. Similarly, Urchin like highly sensitive humidity sensor is also reported via alkali oxidation method to grow in situ TiO₂ nanowires on two-dimensional Ti₃C₂ MXene which creates enhanced surface area. Chen et al. developed microspinning structures by a simple abrasive paper stencil printing process. The obtained sensor showed high sensitivity (151.4 kPa), short response time (<130 ms), and excellent cycle stability over 10,000 cycles. Another facile approach for the preparation of sensor with wrinkled structure by spraying the active material onto the surface of a pre-stretched polyacrylate tape. Also the sensor was demonstrated to have sensitivity of 148.26 kPa in a wide pressure range up to 16 kPa with excellent durability.

Song et al. fabricated hollow-structured MXene-polydimethylsiloxane composites (MPC) by utilizing nickel foam as the three-dimensional substrate for dip-coating. The resultant MPC performs a wide working range with bending angles of 0° to 180° and excellent long-term reliability up to 1000 cycles which can be employed as a bendable piezoresistive pressure sensor.

Electrospinning is one of the widely accepted fabrication method of developing nanofibers with good performance for various applications like catalysis, drug delivery, energy storage, etc. Yang et al. developed MXene/polyurethane mat through electrospinning. The sensor showed high sensitivity, low detection limit (0.1%) and a large and tunable sensing range (up to 150%).

Wang et al. reported the use of zero-dimensional silicon nanoparticles with MXene in cotton fabric to form superhydrophobic textile sensor with broad sensing range. This material exhibited superior sensitivity (S = 12.23 kPa⁻¹), a stable response under press-relaxing cycles, and broad sensing ranges (pressure: 8.8Pa - 70 kPa, bending: 0 - 180°, torsion: 0 - 628 radm⁻¹). Also, the conductivity is sustained under wet and corrosive conditions. Liu et al. observed that an increase in the amount of fluoride-containing functional groups also improved the hydrophobicity of the cotton fabric surface.

Sharma et al. developed a highly sensitive sensor by sandwiching Ti₃C₂Tx/PVDF-TrFE composite nanofibrous scaffolds as a dielectric layer between biocompatible poly-(3,4-ethylenedioxythiophene) polystyrene sulfonate/polydimethylsiloxane electrode. The authors demonstrated a high sensitivity of 0.51 kPa with a detection limit of 1.5 Pa for the sensor along with high reliability of 10,000 cycles.

Recently Zheng et al. reported a novel method for the development poly(3,4- ethylenedioxythiophene) (PEDOT) and Ti₃C₂Tx decorated cotton fabric strain sensor. The authors used vapor phase polymerization (VPP) and spray-coating strategy and achieved a low sheet resistance of 3.6 Ωsq⁻¹ and a high EMI shielding effectiveness of 36.62 dB at an applied voltage of 12 V along with good strain sensitivity. Metal nanoparticle loaded MXenes are also reported as mechanical sensors. Li et al. reported such a work in which silver nanoparticles (AgNPs) were loaded on MXene nanosheets and compounded with one dimensional silver nanowires (AgNWs) with high strain sensitivity. AgNPs built a bridge between AgNWs and MXene, ensuring continuity and a high gauge factor even at a large strain.

3. Electrochemical and Luminescence based sensors

Biochemical sensors are always a 'hotspot' of electrochemical sensor research due to their vast application potential in biomedical field. Highly sensitive and selective sensors which can detect biomolecules and drug molecules are always a significant area of research interest. Further electrochemical biosensors which can detect biomarkers with good performance in detecting diseases and metabolic disorders have also got good research attention over the last decade. Here we are discussing the most recent developments in biosensors which are fabricated using MXenes for application in biomedical field.

One of the initial works reported for demonstrating the sensing action of MXene is the detection of Glucose. Rakhi et al. reported an interesting glucose biosensor based on Au/MXene nanocomposites. This study revealed that Au nanoparticles enhance the electron transfer process between the electrode and the Glucose oxidase enzyme. A linear response range for glucose detection was observed in the concentration range from 0.1 to 18 mm with a relatively high sensitivity of 4.2μAmm⁻¹cm⁻² and a detection limit of 5.9 μm with good reproducibility, repeatability and sensitivity. Li et al. used three-dimensional porous MXene/NiCo-LDH nanocomposite as Glucose sensor. Good reproducibility and selectivity were obtained for the sensor with a working potential of 0.45V (vs. SCE). The sensor also exhibited a broad linear range, a low limit of detection of 0.53μM and rapid response. It was observed that fast electron transfer rate and easy diffusion of electrolyte were attributed to porous NiCo-LDH nanosheets on the MXene substrate.

Kadirsoyet et al. successfully developed a quartz crystal microbalance (QCM) sensor for the detection of chlorpyrifos, a common broad-spectrum organophosphate insecticide using delaminated sulfur-doped MXene (d-S-Ti₃C₂Tx MXene). The sensor possesses linear response range of 1.0×10⁻¹² - 1.0×10⁻¹⁰ M with a detection limit of 3.0 × 10⁻¹³ M.

It is evident that the aggregation of MXene flakes significantly affects their electrocatalytic activity. Thus, delamination approaches like sonication are reported, which can prevent aggregation of MXene flakes. Ma et al. developed a sensor for the detection of Fisetin using self-assembly of negatively charged Ti₃C₂Tx MXene flakes and positively charged amine-functionalized carbon nanotubes (NH₂-CNTs). Fisetin is a dietary antioxidant used for various ailments in humans including enhancement of chemotherapeutic drugs. This study revealed the ability of NH₂-CNTs to form well-defined porous structures by
preventing the aggregation of MXene flakes and thereby significantly enhancing the electrocatalytic activity. The sensor was successfully used in real samples and showed a linear detection range for Fisetin concentration range 0.003 μmolL⁻¹ to 20.0 μmolL⁻¹ with a detection limit of 1.0 nmolL⁻¹.

Ammonia sensors always have a significant area of interest owing to their adverse effects in the environment. Wu et al. provided a good example of employing MXene based sensors to monitor environmental pollutants like ammonia at room temperature.[79] The authors prepared Ti₃C₂ MXene by etching off Al atoms from Ti₃AlC₂ and coated on the surface of ceramic tubes to construct gas sensors. It was observed that the material has good selectivity for NH3 with linear response in concentration range from 10 to 700 ppm in room temperature. Xiao et al. used first principle simulation studies and observed that NH3 could be strongly adsorbed on O-terminated semiconducting MXenes M₂CO₃ (M = Sc, Ti, Zr, and Hf), which renders them the potential candidates as the NH₃ sensor.[80] Further, the authors observed that Zr₂CO₃ as an example, it is highly selective towards NH₃ against other common gas molecules, and the adsorption energy dramatically increases from -0.81 eV to -0.20 eV when extra two electrons are injected into the Zr₂CO₃ sheet of 3 × 3 dimension.

Chen et al. employed the excitation-dependent blue photoluminescence (PL) of Ti₃C₂ quantum dots for fabricating intracellular pH sensor by combining with [Ru(dpp)₃]Cl₂. The authors used a sonication cutting and hydrothermal approach for the surface modification of Ti₃C₂ and demonstrated high PL response attributed to the deprotonation of surface defects.[81]

Yuan et al. developed a three-dimensional (3D) MXene framework through combined electrospinning and self-assembly approach for the detection of volatile organic compounds.[82] The sensor exhibited a wide detection range from 50 ppb, fast response and recovery time (<2 min), with retention of performance for 1000 bending cycles for various species such as acetone, methanol and ethanol.

Fang et al. used negatively charged OH-functionalized composite film with Nafion for the fabrication of solid-state electro-chemiluminescent sensor.[83] The authors reported long-term stability and a very low detection limit for the sensing of Tripropylamine. Further the real-sample analysis of the sensor was demonstrated using single-nucleotide mismatch discrimination in human urine, which got promising results.

Sharma et al. reported that Ti₃C₂O₂ based probe can provide superior sensing of Cortisol with a detection limit of 15.7 fg/mL.[84] This MXene based Cortisol sensor is relevant for real-time applications since it provides a non-invasive fiber optic surface plasmon resonance (FOSPR) based sensor probe to detect salivary Cortisol at the wavelength of 830 nm.

Apart from Titanium-based MXenes, other MXenes are also in their initial stage of exploration. One of such study was reported by Lee et al., which employed the use of vanadium carbide MXene (V₂CTₓ) for non-polar gas sensing applications.[85] The sensor consists of single-/few-layer two-dimensional V₂CTₓ on polyimide film and showed ultrahigh sensitivity at room temperature.

4. Scope and future perspectives
MXenes are in a significant pace of development and got exciting research interest in various applications like hydrogen storage, catalysis, supercapacitors, batteries, and desalination etc. Apart from this, fabrication of MXene based sensors paved the way of the horizon of opportunities in the detection of different categories of target molecules. Some of the most relevant candidates of these target molecules include environmental pollutants, biomolecules, drug molecules and other harmful chemical entities. Further, the categories of piezoresistive MXene sensors are also a developing area which possesses exciting opportunities for the design and development of wearable electronics. But the area is still in a budding phase which needs enough research attention to make sensors that can be applied to a wide range of applications and to detect novel and relevant analytes. We would like to invite the attention of readers into the following aspects of MXene research to accelerate the journey which gives new opportunities for MXene based sensor devices as illustrated in Fig. 5.

- **Necessity for novel and cost-effective synthesis strategies:** Even though more studies are emerging which demonstrates modified strategies for the preparation of MXenes, the design and development of MXenes are not much shifted from conventional chemical strategies. This makes the fabrication method tedious which affects the options for getting appropriately designed MXenes for specific applications. Microwave and Utrasound assisted methods should be considered to improve the preparation process.

- **Gap between theory and practice:** There are many MXenes that are reported for their stable existence in literature through theoretical predictions. These approaches involve high throughput screening and density functional calculations which gives the idea of elemental combinations that produce stable MXenes. But most of
the studies are still focused on Titanium based MXenes like Ti$_3$C$_2$T$_x$, which are produced experimental preparation methods. Urgent attention is needed in this perspective which could give rise to novel formulations of MXene, which probably possess superior properties.

- **Lack of long term stability for sensor applications:** It has been observed that both single and multilayered MXenes like Ti$_3$C$_2$T$_x$ will degrade with humidity and an aqueous environment. This problem limits the application of MXenes for fabricating into stable sensor probes which obviously need to be operated in hydrophilic conditions. Thus, other types of MXenes have be synthesized which is having good conductivity along with high stability to be used as sensors. Weak gelation of MXene layers is another problem that significantly affects their morphology.

- **Biosafety and toxicity:** Even though there are some studies that analyzed the toxicity aspects of MXenes, this perspective is still not explored in detail. Toxicity, including both cytotoxicity and genotoxicity, along with biosafety aspect has to be thoroughly investigated prior to their real-time application as sensors.

5. Conclusion

MXenes have amazing research interest within a short time span after their discovery due to their exciting layered structure and metallic-like conductivity. Researchers across the world working on sensors are shifting their focus to novel two dimensional materials like MXenes which can show superior performance over the conventional materials. The good conductivity and high stability of MXenes makes them an ideal candidate for the fabrication of sensors that can be employed for diagnostic applications. On the other hand, this area still needs urgent attention for making new category of MXenes other than Titanium based materials. Even though theoretical studies are increasing the choice of preparing MXene based sensors from other transition metals, most of the works in this area are not successfully explored other varieties. There is a high probability and scope of developing new category of MXene candidates which might possess good performance over the reported materials. We hope this review will help the researchers understand the current scenario existing in MXene based sensors that have not attained good scientific attention and high possibility of novel discoveries.

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**Conflict of Interest**

There is no conflict of interest.

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**References**


**Author information**

Vishnu Sankar Sivasankarapillai received his B.Sc and M.Sc Chemistry degrees from Mahathma Gandhi university, Kerala, India. He is currently working towards Ph.D. degree under the supervision of Dr. Ragupathy Dhanusuraman, Department of Technology at National Institute of Technology Puducherry, India. His research interests include Biomaterials and Nanocomposites for energy and biomedical applications.

Tata Sanjay Kanna Sharma received his B.S. (2012), and M.S. (2014), degrees from Sri Venkateswara University, Tirupathi, India and acquired Doctor of Philosophy Ph.D. (2021) from National Taipei University of Technology, Taipei, Taiwan. His current research interest mainly focuses on the synthesis of transition metal/Rare earth vanadate-carbon based Aerogels, Novel metal oxides combined with green synthesized carbon sources for real-time analysis in biosensors. He is more interested in Optical properties of Graphene, Landau levels for sensor applications and drug delivery.

Hwa-Yuan Kuo is an Associate Professor and the director of the Center for Biomedical Industries at the National Taipei University ofTechnology. Dr. Hwagraduated and received her PhD from the School of Medicine, the Johns Hopkins University. She is the president of theMedical Association for Indigenous Peoples of Taiwan (MAIPT). Dr. Hwa's scientific interests are: 1) nanotechnology and biosensor, 2) newdrug discovery for human diseases by proteomics and genomics approaches and 3) glycobiology, especially on enzymes kinetics. She has published 100 + conference and journal articles and 15 patents.
Saikh Mohammad Wabaidur is working as Researcher at King Saud University, Saudi Arabia. He received his PhD in 2021, from Banasthali University, India. He has published 110 research articles in international journals. His research field includes Separation, identification and determination of biologically important compounds and pharmaceutical drugs.

Subramania Angaiah has been working in the field of Materials Electrochemistry for the last 20 years. His area of research includes M-ion batteries, Supercapacitors, Dye-sensitized solar cells, Nanocomposites, Electrospun polymer membrane electrolytes for electrochemical energy devices, Corrosion and Surface Engineering etc. He has published more than 130 research articles in the peer-reviewed international journals.

Ragupathy Dhanusuraman is Assistant professor and Head at the Department of Chemistry, National Institute of Technology Puducherry, Karaikal, India. He received his PhD in 2010, from Kyungpook National University, South Korea. He has published 75 research articles in international journals. His research field includes the development of newer nanomaterials for energy, biosensor and electro-catalytic applications.

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