Developing MXenes from Wireless Communication to Electromagnetic Attenuation

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HIGHLIGHTS

• The industrial application and foundational research of MXenes at gigahertz frequency are systematically reviewed.

• The design principles of “lightweight, wide, and strong” are specifically highlighted.

• Current challenges and future directions for MXenes in wireless communication and electromagnetic attenuation are outlined.

ABSTRACT There is an urgent global need for wireless communication utilizing materials that can provide simultaneous flexibility and high conductivity. Avoiding the harmful effects of electromagnetic (EM) radiation from wireless communication is a persistent research hot spot. Two-dimensional (2D) materials are the preferred choice as wireless communication and EM attenuation materials as they are lightweight with high aspect ratios and possess distinguished electronic properties. MXenes, as a novel family of 2D materials, have shown excellent properties in various fields, owing to their excellent electrical conductivity, mechanical stability, high flexibility, and ease of processability. To date, research on the utility of MXenes for wireless communication has been actively pursued. Moreover, MXenes have become the leading materials for EM attenuation. Herein, we systematically review the recent advances in MXene-based materials with different structural designs for wireless communication, electromagnetic interference (EMI) shielding, and EM wave absorption. The relationship governing the structural design and the effectiveness for wireless communication, EMI shielding, and EM wave absorption is clearly revealed. Furthermore, our review mainly focuses on future challenges and guidelines for designing MXene-based materials for industrial application and foundational research.

KEYWORDS MXene; Wireless communication; Electromagnetic wave absorption; Electromagnetic interference shielding
1 Introduction

Wireless communication has been gaining popularity with the arrival of the age of artificial intelligence [1–3]. This is accompanied by a surge in the demand for all kinds of portable devices. These devices require a concealed integration of radio communication electronics without sacrificing lightweight and transportability [4–6]. Therefore, it is necessary to develop new routes of antenna fabrication [7, 8]. It is difficult to fabricate ultra-thin, flexible, and conformal antenna using traditional metal materials because of the skin depth limitation [9]. To overcome this shortcoming, carbon-based nanomaterials have been explored for wireless communication applications. For example, Elwi et al. prepared multi-walled carbon nanotube antennas that afforded a remarkable enhancement in the bandwidth [6]. Vacirca et al. reported an onion-like carbon antenna that showed a peak gain of −1.48 dBi, just 3 dB less than that of a copper dipole antenna [8]. However, the low conductivity of carbon inhibits these materials from achieving reasonable radio-frequency performance. Thereafter, two-dimensional (2D) nanomaterials, such as graphene, MoS$_2$, and others, have been used to fabricate antennas, thereby furnishing antennas with thinner patches. For example, Shin et al. fabricated a graphene antenna, affording a high transmitted power efficiency of 96.7% [1]. To date, it is still a challenge to find a flexible material with high conductivity for antenna fabrication.

On the other hand, the use of a large number of wireless communication devices will lead to an explosive increase in electromagnetic (EM) radiation on the scale of “big data” [10–12]. The undesired EM radiation directly affects the operation of electronic equipment and also indirectly influences human health, as long-term exposure to EM radiation may cause cancer and other health problems [13–16]. How to avoid the harm from EM radiation has always been a research hot spot. The development of electromagnetic interference (EMI) shielding and EM wave absorbing materials is the key to solving the above problem [17–22]. Recently, various materials have been globally studied as EMI shielding or/and EM wave absorbing materials, including zero-dimensional (0D), one-dimensional (1D), and 2D materials. Among them, 2D materials are the materials of choice as they are lightweight, have large aspect ratios, and offer distinguished electronic properties. For example, Cao’s group reported that chemically graphitized r-GOs exhibited high-efficiency EMI shielding effectiveness (EMI SE) at elevated temperatures. The EMI SE of the composites with 20 wt% r-GOs reached a maximum at ~38 dB [12]. Zhang et al. prepared 2D WS$_2$-rGO heterostructure nanosheets. The composite containing 40 wt% WS$_2$-rGO showed a minimum reflection loss ($RL$) of −41.5 dB, with the absorption bandwidth reaching up to 13.62 GHz [15].

MXenes, as a novel family of 2D materials, possess huge potential in the fields of wireless communication, EMI shielding, and EM wave absorption owing to their excellent electrical conductivity, numerous family members, mechanical stability, high flexibility, and ease of processability [23, 24]. At present, research on MXene antennas is still in the exploratory stage, but the excellent properties of these materials for wireless communication have been widely regarded. Moreover, since the discovery of the outstanding EMI shielding performance of Ti$_3$C$_2$T$_x$ MXene in 2016, MXenes have become the leading materials for EMI shielding and EM wave absorption with the fastest growing number of related research publications (Fig. 1). Moreover, as EM attenuation materials, the excellent chemical and physical properties of MXenes have facilitated the development of pure MXenes and hybrids with controlled structural designs such as films, foams, aerogels, and fabrics (inset of Fig. 1).

This review systematically summarizes the effects of MXenes with various structural designs on wireless communication, EMI shielding, and EM wave absorption. The most feasible strategies for high-performance wireless communication, EMI shielding, and EM wave absorption are revealed by discussing the different approaches for modifying the structures of MXenes. Finally, we provide an overview of the further development and prospects of MXenes for wireless communication and EM attenuation.
2 Antenna Mechanism and EM Attenuation Mechanism

2.1 Antenna Mechanism

Actually, the wavelength of substrate-based antennas is inversely proportional to frequency, and the length of designed antenna is selected according to antenna types and wavelength. $\lambda$ is defined as [25–30]:

$$\lambda = \frac{c}{f}$$  \hspace{1cm} (1)

Two important indexes to judge the quality of designed antenna are bandwidth and voltage standing wave ratio (VSWR) or return loss of the antenna [31–33]. The VSWR is to quantify the impedance matching at the operating frequency and different thicknesses, which can be calculated by the formula [34, 35]:

$$\text{VSWR} = \frac{1 + |S_{11}|}{1 - |S_{11}|}$$  \hspace{1cm} (2)

where $S_{11}$ represents the reflection coefficients. VSWR is the ratio between the peak amplitude and the minimum amplitude of standing wave. Standing wave is caused by any mismatch at the input of antenna, which will cause the input power to be reflected back [36, 37]. VSWR equal to 1 means that there is no standing wave ($S_{11} = -\infty$), and the antenna is an ideal match.

The research on 2D materials as patch antennas is the most extensive. Different substrates have been proposed for flexible patch antennas, such as rubber, polyethylene, cellulose nanopaper, and others, to achieve improved efficiency. The patch antenna, using natural rubber as substrate, plays a significant role in wireless communication as the mechanical properties of rubber make the antenna flexible. Moreover, an antenna on a polyethylene substrate was designed and fabricated with distinct bending curvature, affording reliable performance within the designed C-band [38].

2.2 Shielding Mechanism

The ability of a shield to against the incoming EM radiation is measured by the EMI SE, which is defined as the ratio of the transmitted and incident powers, generally on a logarithmic scale, as expressed in Eq. (3) [39]:

$$SE_T(\text{dB}) = 10\log \frac{P_T}{P_I} = 20\log \frac{E_T}{E_I}$$  \hspace{1cm} (3)

where $P_I$ and $P_T$ represent incident wave power and transmitted wave power, respectively. $E_I$ and $E_T$ represent electric field intensity of incident wave and electric field intensity of transmitted wave power, respectively. According to the theory of Schelkunoffs, the total EMI SE is the sum of

Fig. 1 Number of publications focused on MXenes for EMI shielding and EM wave absorption. Source: Web of Science Core Collection, 2020 (inside: 2D MXenes serve as EMI shielding materials and EM wave absorption materials)
reflection ($S_R$), absorption ($S_E$), and multiple reflections ($S_M$), as shown in Eq. (4) [40]:

$$S_E = S_{ER} + S_{EA} + S_{EM} \quad (4)$$

To quantify the value of $S_{ER}$ and $S_{EA}$, the concept of the absorption ($A$), reflection ($R$) and transmission ($T$) coefficient is proposed. The relation among them can be obtained as follows:

$$S_{ER} = 10\log\left(\frac{1}{1 - R}\right) \quad (5)$$

$$S_{EA} = 10\log\left(\frac{1 - R}{T}\right) \quad (6)$$

The value of $A$, $R$, and $T$ can be calculated from the measured scattering parameters ($S_{mn}$). $S_{mn}$ represents that how energy is scattered from a shield. “m” indicates the port receiving the radiation energy, and “n” indicates the port that is transmitting the incident energy. Accordingly, $A$, $R$ and $T$ can be calculated via the relationships:

$$A + R + T = 1 \quad (7)$$

$$R = |S_{11}|^2 = |S_{22}|^2 \quad (8)$$

$$T = |S_{21}|^2 = |S_{12}|^2 \quad (9)$$

Both reflection and absorption provide great contributions for excellent EMI shielding. However, in consideration of green shielding, less reflection is better. The ideal EMI shielding involves strong absorption with no reflection and transmission, as shown in Fig. 2b. Multiple reflection also plays an important role in EM wave attenuation. Multiple reflections between the front and back of the shield

![Fig. 2](https://example.com/fig2.png)

**Fig. 2** Situations in which the EMI shield responds to the EM wave: **a** allowing all waves to enter with large transmission, **b** allowing all waves to enter with no transmission, and **c** strong secondary reflection. The situations in which the absorber responds to the EM wave: **d** good impedance matching with weak attenuation, **e** an ideal absorber with good impedance matching and strong attenuation, and **f** poor impedance matching with strong attenuation.
contribute a lower EMI SE. However, when the thickness of the shield close to or larger than the skin depth
\[
(\delta = \left( \frac{\sqrt{\frac{\pi f \mu_0 \mu_r}}{\sigma}} \right)^{-1}, \text{where } \mu_0 = 4 \times 10^{-7} \text{H m}^{-1}, \mu_r \text{ is permeability of an absorber, and } \sigma \text{ is electrical conductivity}) \text{ or when the } \text{SE}_T > 10 \text{ dB, the effect of multiple reflection should be neglected.}
\]

Specific shielding effectiveness (SSE) is derived to compare the effectiveness of shielding materials taking into account the density. Lightweight materials afford high SSE. SSE can be calculated as follows:

\[
\text{SEE} = \frac{\text{EMI SE}}{\text{density}} = \text{dB cm}^3 \text{g}^{-1}
\]

(10)

To account for the thickness contribution, the following equation is used to evaluate the absolute effectiveness (SSE_t) of a material in relative terms

\[
\text{SSE}_t = \frac{\text{SSE}}{t} = \text{dB cm}^3 \text{g}^{-1}
\]

(11)

### 2.3 Absorption Mechanism

The responses of an EM wave absorption to an incident EM wave are determined by the absorber’s permittivity and permeability. The EM wave absorption capacity can be expressed by the following equations [41]:

\[
\text{RL} = 20 \log \left[ \frac{Z_{in} - 1}{Z_{in} + 1} \right]
\]

(12)

\[
Z_{in} = \sqrt{\mu_r \varepsilon_r} \tanh \left[ j \left( \frac{2\pi f d}{c} \right) \sqrt{\mu_r \varepsilon_r} \right]
\]

(13)

where \(Z_{in}\) is the normalized input impedance, \(\varepsilon_r\) and \(\mu_r\) are the relative complex permittivity and permeability of an absorber and \(d\) is the thickness of the absorber.

Excellent EM wave absorption performance of an absorber depends not only on efficient EM wave attenuation but also on impedance matching. As shown in Fig. 2e, the absorber exhibits its ideal impedance matching, which means that all EM waves are allowed to penetrate the absorber. And, its high-efficiency EM attenuation is usually derived from dielectric loss and magnetic loss, as well as multi-scattering or multi-reflection.

Generally, the dissipation pathway within the absorber is described as dielectric loss. The dielectric loss can be thought as a sort of friction to the displacement of the subatomic particles, then the passing EM wave attenuates, presenting as the subsequent generation of heat. Dielectric loss is constructed by conduction loss and polarization loss; the relation among them can be expressed by the following formula:

\[
\varepsilon'' = \varepsilon''_c + \varepsilon''_p
\]

(14)

where \(\varepsilon''_c\) and \(\varepsilon''_p\) represent conduction loss and polarization loss, respectively.

Conduction loss plays an important role in the materials with high conductivity, such as, MXene, graphene and other carbon materials. Actually, the conduction loss is the energy loss of EM wave caused by electron transition. Up to now, Cao’s group has done a lot of work to clarify the role of conduction loss. For example, the electron-hopping model (EHM) was established to explain the mechanism of conduction loss in the carbon fibers (CFs) and multi-walled carbon nanotubes (MWCNTs) [42]; a model of aggregation-induced charge transport (AICT) was proposed to illustrate electron transport in whole MWCNTs/SiO2 composites [43]; electron transition theory was used to explain the loss behavior of Ti3C2T x MXene [44]. Now, the role of conduction loss caused by electron transition has been widely recognized in the design of EM wave-absorbing materials.

Polarization loss is generated by the behavior of dipoles. Dipoles are generated in the site of functional groups, defects, and interfaces. Under a high-frequency alternating electric field, when rotation of dipoles cannot follow the change of electric field, dipole orientation polarization loss occurs, which is another key role of dielectric loss. Cao’s group made important contributions to the resources, characterization techniques, and semiquantitative methods of polarization relaxation. In 2008, they proposed a capacitor-like model and an equivalent circuit model to explain the EM wave response of CdS-Fe2O3 heterostructures [45]. In 2012, the perfect polarization relaxation was established in Fe3O4-MWCNTs and PANI-Fe3O4-MWCNTs [46]. Later, the capacitor-like model is applied to visualize the interfacial polarization in MWCNTs, graphene, MXene, or their nano-hybrids, etc. After that, a semiquantitative research strategy of multiple polarization, that is the separation of the contribution of electron transport and dipole polarization, was established by them to accurately analyze the source of polarization [47]. Recently, they proposed a new concept of polarization genes and made a semiquantitative characterization and definition [48]. In conclusion, on the premise of meeting the impedance matching, effectively rising the
conduction loss and polarization loss will greatly improve the EM wave absorbing performance of absorber. Magnetic loss is caused by eddy currents, natural resonance, and exchange resonance. The eddy current is inevitable. If magnetic loss only originates from eddy current loss, the value of $\mu''(\mu')^{-2}f^{-1}$ is constant when frequency changes. Natural resonance usually occurs at low frequency (2–10 GHz), while exchange resonance occurs at high frequencies (> 10 GHz).

3 MXene-Based Materials for Antenna

Research on MXenes for wireless communication applications is still in the exploratory stage. Gogotsi’s group has undertaken pioneering research in this field. According to the different usage frequency, MXene antennas can be divided into low-frequency antennas (< 2.4 GHz) and high-frequency antennas (> 5.6 GHz).

3.1 Low-Frequency Antenna

Gogotsi’s group firstly designed and investigated the Ti$_3$C$_2$Tx MXene dipole antenna at 2.4 GHz [49]. The translucent MXene antenna with a thickness of ~ 100 nm had a reflection coefficient of less than −10 dB. By increasing the antenna thickness to 8 µm, the reflection coefficient reached −65 dB. The VSWR was less than 2 for MXene antennas with various thicknesses despite the fact that the surface resistance increased significantly with thicknesses below 100 nm. The radiation pattern of an 8-mm-thick dipole antenna presented typical dipole radiative behavior. Moreover, the MXene antenna afforded a maximum gain of 2.11 dB at the thickness of 8 µm, which converges with the maximum gain of an ideal half-wavelength dipole antenna (2.15 dB).

Li et al. prepared a stretchable Ti$_3$C$_2$Tx nanosheets (MXene) and single-walled carbon nanotubes (SWNTs) S-MXene dipole antenna (Fig. 3a) [50]. The resonant frequencies of an S-MXene antenna were linearly dependent on the applied strains (Fig. 3b). Moreover, the S-MXene antenna afforded nearly the same reflection $|S_{11}|$ (~ −33 dB) at the same resonant frequency (1.425 GHz) during the fatigue test up to 100% uniaxial strains for 500 cycles (Fig. 3c).

3.2 High-Frequency Antenna

Gogotsi’s group reported on micrometer-thin and flexible MXene microstrip patch antennas with target frequencies of 5.6, 10.9, and 16.4 GHz, produced by a simple spray-coating fabrication method (Fig. 3d, e) [51]. The return loss values of MXene antennas with a thickness of 5.5 µm achieved −29, −25, and −48 dB at 5.6, 10.9, and 16.4 GHz (Fig. 3f), respectively, which demonstrated that MXene patches were capable of delivering RF power efficiently to the radiator. The radiation efficiency increased with the thickness of MXene patches (Fig. 3g), which was due to the decreasing conductor loss. Moreover, the outstanding performance of MXene patch antennas was comparable to their copper counterparts (Fig. 3h).

4 MXene-Based Materials for EMI Shielding

4.1 Pure MXene Matrix

4.1.1 Non-annealing

Liu et al. prepared multilayer Ti$_3$C$_2$T$_x$ by etching Ti$_3$AlC$_2$ with 40% HF at room temperature (RT) for 24 h [52]. The multilayer Ti$_3$C$_2$T$_x$/wax with the 60 wt% Ti$_3$C$_2$T$_x$ content displayed outstanding EMI shielding performance of 39.1 dB at the thickness of 2 mm. Hu et al. fabricated multilayer Ti$_3$C$_2$T$_x$ by etching Ti$_3$AlC$_2$ with 40% HF solution at 50 °C for only 0.5 h [53]. The multilayer Ti$_3$C$_2$T$_x$ prepared in such a short time also exhibited excellent EMI shielding performance. For example, the Ti$_3$C$_2$T$_x$/wax with 70 wt% Ti$_3$C$_2$T$_x$ showed EMI shielding performance of 34 dB at 18 GHz.

Previous studies focused on the EMI shielding performance of multilayer Ti$_3$C$_2$T$_x$. He et al. investigated the difference between Ti$_3$C$_2$T$_x$, nanosheet and multilayer Ti$_3$C$_2$T$_x$ in EMI shielding performance [54]. Different etchants led to different centrifugal results (Fig. 4a, b) and different morphologies (Fig. 4c–f). The Ti$_3$C$_2$T$_x$ nanosheet showed a much better EMI shielding performance compared with the multilayer Ti$_3$C$_2$T$_x$ owing to the formation of local conductive networks (Fig. 4g–l). The Ti$_3$C$_2$T$_x$/wax matrix with 80 wt% Ti$_3$C$_2$T$_x$ loading showed EMI shielding performance of 58.1 dB at the thickness of only 1 mm.
Li et al. prepared the multilayer Ti$_2$CT$_x$ MXene by HCl/LiF etching Ti$_2$AlC for 48 h at 40 °C [55]. The multilayer Ti$_2$CT$_x$ presented outstanding EMI shielding performance of 70 dB with a thickness of only 0.8 mm in the X-band, exceeding most of reported graphene-based EMI shielding composites, owing to the multilayered structure and the high electrical conductivity (0.30 S cm$^{-1}$).

The matrix of EMI-absorbing materials is usually made of wave-transparent materials. The imaginary part of the dielectric constant of the matrix is zero, which means there is no EM loss. Paraffin is generally used as the matrix of absorbing materials. There are also other wave-transmitting materials that can be used as the matrix, such as polystyrene (PS) and polyvinylidene fluoride (PVDF). A Ti$_3$C$_2$T$_x$ nanosheet@PS matrix was fabricated by electrostatic assembling of negative Ti$_3$C$_2$T$_x$ nanosheet on positive polystyrene microsphere [56]. The Ti$_3$C$_2$T$_x$@PS with 1.9 vol% Ti$_3$C$_2$T$_x$ loading exhibited an outstanding EMI shielding performance of >54 dB over the whole X-band. Such excellent EMI shielding performance was attributed to the high conductivity (1081 S m$^{-1}$) of Ti$_3$C$_2$T$_x$ and their highly efficient conducting network within PS matrix. Rajavel et al. reported that the multilayer Ti$_3$C$_2$T$_x$-PVDF with 22.55 vol% Ti$_3$C$_2$T$_x$ displayed remarkable shielding performance of 48.47 ± 3.5 dB with a thickness of 2 mm [57]. Such outstanding shielding performance
was attributed to the formation of conducting network along with the assembly of micro-capacitor network.

4.1.2 Annealing

The improvement of EMI shielding properties by annealing is also widely studied. Yin group investigated the EMI shielding performance of multilayer Ti$_3$C$_2$Tx annealed at 800 °C (Ti$_3$C$_2$Tx-$800$) in Ar atmosphere. The total SE of Ti$_3$C$_2$Tx-$800$/wax was 32 dB [58]. After that, the Ti$_3$C$_2$Tx-$200$ was fabricated by annealing Ti$_3$C$_2$Tx at 200 °C for 2 h in Ar + 5% H$_2$ atmosphere [59]. The Ti$_3$C$_2$Tx-$200$/epoxy displayed enhanced EMI shielding performance compared with Ti$_3$C$_2$Tx/epoxy. Ji et al. investigated the EMI shielding performance of multilayer Ti$_3$C$_2$Tx at different annealing temperatures (800, 950, 1100, 1250, 1400, and 1550 °C for 1 h) in Ar atmosphere [60]. Among them, the multilayer Ti$_3$C$_2$Tx annealed at 1100 °C displayed the best EMI shielding performance, showing excellent EMI shielding performance of 76.1 dB.
4.2 MXene Hybrid Matrix

It is common that MXene is doped with one-dimensional nanoparticles. Ti$_3$C$_2$T$_x$ nanosheet/Ni chain hybrid was fabricated by a one-step hydrothermal process [61]. The Ti$_3$C$_2$T$_x$/Ni hybrid with 50 wt% Ti$_3$C$_2$T$_x$ showed EMI SE of 66.4 dB. The excellent EMI shielding performance was due to the synergistic effect of conductive Ti$_3$C$_2$T$_x$ and magnetic Ni chains, by which the dielectric properties and EM loss can be easily controlled to obtain good EM wave dissipation ability. Nb$_2$CT$_x$/Nb$_2$O$_5$-Ag ternary hybrid nanostructures were fabricated by self-reduction and oxidation of Nb$_2$CT$_x$ in the presence of metallic salt (AgNO$_3$) [62]. The Nb$_2$CT$_x$/Nb$_2$O$_5$-Ag hybrid-wax matrix with a thickness of 1 mm showed excellent EMI SE of 68.76 and 72.04 dB in the X- and Ku-band region, respectively. The excellent EMI shielding performance was attributed to the strong electrical conductivity, increased interface polarization, and multiple reflection loss between the ternary interfaces.

MXene is not only doped with one-dimensional nanoparticles, but also doped with two-dimensional materials. Song et al. investigated the EMI shielding performance of honeycomb structural rGO–Ti$_3$C$_2$T$_x$ MXene/epoxy nanocomposites [63]. The introduction of rGO–Ti$_3$C$_2$T$_x$ can fully play synergistic effects of rGO and MXene, to greatly improve the electrical conductivity and EMI SE of rGO–Ti$_3$C$_2$T$_x$ MXene/epoxy nanocomposites. When the cell size of 0.5 mm with 1.2 wt% rGO + 3.3 wt% Ti$_3$C$_2$T$_x$ loading, the enhanced $\sigma$ (387.1 S m$^{-1}$) and EMI SE (55 dB) values were reached, which were, respectively, 2978 and 5 times of rGM/epoxy nanocomposites (σ of 0.13 S m$^{-1}$, EMI SE of 11 dB) at the same loading of directly blending rGO–Ti$_3$C$_2$T$_x$ fillers.

The shielding properties of multi-doped MXene have also been studied. Raagulan fabricated the Ti$_3$C$_2$T$_x$ MXene-p-aminophenol (PAT)-conductive polymer (CP) by a cost-effective spray coating technique and characterization [64]. The composite showed excellent EMI shielding performance of 45.18 dB and good electric conductivity of 7.813 S cm$^{-1}$.

4.3 Film

The EMI shielding properties of MXene as a film have been widely studied, and there are numerous research results. As a shield, MXene films can be divided into three categories: pristine MXene films, organic-hybrid MXene films, and inorganic-hybrid MXene films. The pristine MXene film refers to the film fabricated from pure MXene nanosheets, organic-hybrid MXene films contain organic substances, and inorganic-hybrid MXene films are MXene films doped with an inorganic substance.

4.3.1 Pristine

Koo’s group reported that a 45-µm Ti$_3$C$_2$T$_x$ film displayed EMI SE of 92 dB (a 2.5-µm film showed > 50 dB), which is the highest among synthetic materials of comparable thickness produced to date [65]. The outstanding electrical conductivity of Ti$_3$C$_2$T$_x$ films and multiple internal reflections led to this excellent performance. After that, they systematically studied the EMI shielding of Ti$_3$C$_2$T$_x$ MXene-assembled films over a broad range of film thicknesses, monolayer by monolayer [66]. Theoretical research showed that multiple reflection, the surface reflection, and bulk absorption become significant in the shielding mechanism below skin depth. The 24-layer film of 55 nm thickness showed EMI SE of 20 dB, revealing an extraordinarily large absolute shielding effectiveness (3.89×106 dB cm$^2$ g$^{-1}$). Meanwhile, they prepared Ti$_3$CNT$_x$ and Ti$_3$C$_2$T$_x$ MXene free-standing films of different thicknesses by vacuum-assisted filtration and investigated their EMI shielding performance under different annealing temperatures [67]. It is found that Ti$_3$CNT$_x$ film provided a higher EMI SE compared with more conductive Ti$_3$C$_2$T$_x$ or metal foils of the same thickness. This excellent EMI shielding performance of Ti$_3$CNT$_x$ was achieved by thermal annealing, owing to an anomalously high absorption of EM waves in its layered, metamaterial-like structure.

Han et al. systematically studied the shielding properties of 16 different MXene films [68]. All MXene films with micrometer thick displayed excellent EMI shielding performance (> 20 dB). Among them, Ti$_3$C$_2$T$_x$ film displayed the best EMI shielding performance. For example, Ti$_3$C$_2$T$_x$ film with a thickness of only ~40 nm showed the EMI shielding performance of 21 dB.

4.3.2 Organic-Hybrid

The organic substances for hybridization can be divided into three categories: aramid nanofibers (ANFs), cellulose nanofibers (CNFs), and others.
4.3.2.1 ANF Xie et al. prepared Ti$_3$C$_2$Tx nanosheet/ANF composite film by vacuum-assisted filtration approach [69]. The Ti$_3$C$_2$Tx/ANF composite film with different Ti$_3$C$_2$Tx addition exhibited outstanding EMI shielding performance, which was accomplished beyond the commercial standard for EMI shielding materials. The 80%-Ti$_3$C$_2$Tx/ANF composite film with an ultra-thin thickness ~17 µm possessed an EMI SE of ~28 dB in 8.2–12.4 GHz and electrical conductivity of 173.36 S cm$^{-1}$. Weng et al. further confirmed this conclusion [70]. Lei et al. found that the Ti$_3$C$_2$Tx/ANF composite film with a loading of 40 wt% Ti$_3$C$_2$Tx showed high electrical conductivity of 3661.8 S m$^{-1}$ [71], and excellent EMI shielding performance of 24.5 dB, and SEE of 8814.5 dB cm$^2$ g$^{-1}$ at the thickness of 14 µm. Wei et al. reported that the ANF/Ti$_3$C$_2$Tx film with 90 wt% displayed the EMI SE of 34.71 dB at the thickness of 11 µm and SEE of 21,971.37 dB cm$^2$ g$^{-1}$, which would be no recession after 1000 times bending [72].

After that, ternary mixed ANF-MXene film was reported [73]. A double-layered and homogeneously blended ANF-Ti$_3$C$_2$T$_x$ MXene/silver nanowire (ANF-MXene/AgNW) nanocomposite film was fabricated via the facile two-step vacuum-assisted filtration followed by hot-pressing approach, respectively. Compared with the homogeneously blended ones, the double-layered nanocomposite papers possessed greater advantages in EMI shielding performances, which was due to the massive ohmic losses, multiple internal reflections and polarization relaxation of localized defects, and abundant terminal groups.

4.3.2.2 CNF Cao et al. fabricated an ultrathin and highly flexible delaminated Ti$_3$C$_2$Tx (d-Ti$_3$C$_2$Tx) MXene/CNF composite film through a vacuum-filtration-induced self-assembly process [74]. The d-Ti$_3$C$_2$Tx/CNF film with 90 wt% d-Ti$_3$C$_2$Tx content showed high electrical conductivity (739.4 S m$^{-1}$) and outstanding EMI shielding performance (SEE$_{40}$ = 2647 dB cm$^2$ g$^{-1}$). Zhou et al. found that the CNF/Ti$_3$C$_2$Tx film showed EMI SE of ~40 dB and high SEE, up to 7029 dB cm$^2$ g$^{-1}$ with a thickness of only 0.035 mm [75]. Moreover, the EMI shielding properties could withstand the folding test more than 1000 times without obvious reduction. Cui et al. reported that a Ti$_3$C$_2$Tx/CNF film exhibited EMI SE of 42.7 dB with a thickness of 15 µm, owing to the high electrical conductivity (46.3 S cm$^{-1}$) [76].

Zhou et al. fabricated a CNF/Ti$_3$C$_2$T$_x/g$-C$_3$N$_4$ film with Ti$_3$C$_2$T$_x/g$-C$_3$N$_4$ mass ratio of 5:1 presented outstanding performance with EMI SE of 42.99 dB in X-band at the thickness of 28.20 µm [77]. Moreover, both the electrical conductivity and the EMI SE of the film remained nearly unchanged after bending at 135° for 10,000 cycles.

Xin et al. investigated the EMI shielding performance of a Ti$_3$C$_2$Tx/CNF/silver composite film [78]. The Ti$_3$C$_2$Tx/CNF/silver composite film exhibited excellent EMI shielding performance (50.7 dB) and good electrical conductivity (588.2 S m$^{-1}$), attributed to Ti$_3$C$_2$T$_x$ MXene, self-reduction of silver nanoparticles, and the brick-like structure. An ultrathin and flexible carbon nanotubes/Ti$_3$C$_2$Tx/CNF composite film was fabricated via a facile alternating vacuum-assisted filtration process [79]. The film showed a high electrical conductivity of 2506.6 S m$^{-1}$ and EMI SE of 38.4 dB. This result was attributed to the sandwich structure in improving EMI SE, and the gradient structure on regulating the contributions from reflection and absorption.

4.3.2.3 Others The flexible green multilayered Ti$_3$C$_2$Tx/hydroxyethyl cellulose (M-Ti$_3$C$_2$T$_x$/HEC) composite film was prepared via the filtration-assisted self-assembly method (Fig. 5a–c) [80]. The effect of multilayer stacking on the EMI shielding performance was investigated. The EMI SE of the film exceeded 20 dB under the stacking thickness reaching 100 mm (Fig. 5d, e). Notably, the film presented the trend of absorption-dominate green EMI shielding with the decrease of the stacking thickness (Fig. 5f).

Gogotsi’s group investigated the EMI shielding performance of a polyethylene terephthalate (PET)/Ti$_3$C$_2$Tx nanosheet film [81]. The PET/Ti$_3$C$_2$Tx film exhibited electrical conductivity values of 1080 ± 175 S cm$^{-1}$, which profoundly exceeded electrical conductivity values of other 2D materials including graphene (250 S cm$^{-1}$) and reduced graphene oxide (340 S cm$^{-1}$). Such excellent EMI shielding performance was ascribed to the high electrical conductivity and layered structure.

A free-standing, ultrathin, and flexible Ti$_3$C$_2$Tx/poly (3,4-ethylenedioxythiophene)-poly (styrene sulfonate) (PEDOT: PSS) film was prepared by a vacuum-assisted filtration process [82]. The composite film with 11.1 µm prepared a high EMI SE value of 42.1 dB. Meanwhile, the hybrid film exhibited a superior conductivity of 340.5 S cm$^{-1}$ and an excellent specific EMI shielding efficiency of 19,497.8 dB cm$^2$ g$^{-1}$. This result was due to the lamellar structure of the films and multiple interface reflection and polarization.

Luo et al. fabricated a flexible Ti$_3$C$_2$T$_x$ nanosheet/natural rubber (NR) nanocomposite film via vacuum-assisted filtration approach [83]. In the film, Ti$_3$C$_2$T$_x$ nanosheets...
selectively distributed at the interfaces of the NR particles, forming an interconnected network for efficient electron transport, which leads to excellent EMI shielding performance. The Ti$_3$C$_2$Tx nanosheet/NR film with 6.71 vol% of Ti$_3$C$_2$Tx nanosheet showed an outstanding electrical conductivity of 1400 S m$^{-1}$ and a superb EMI shielding performance of 53.6 dB.

Wang et al. prepared a flexible and ultrathin poly(vinylidene fluoride) (PVDF)/Ti$_3$C$_2$Tx/Ni chain composite film by physical mixing [84]. The PVDF/Ti$_3$C$_2$Tx/Ni chain composite film with only 0.10 mm thickness showed EMI shielding performance of 19.3 dB, which increased to 34.4 dB at 0.36 mm thickness. The outstanding EMI shielding performance was attributed to the excellent electrical conductivity (892 S m$^{-1}$).

An ultrathin Ti$_3$C$_2$Tx/calcium alginate (CA) aerogel film was fabricated via divalent metal ion-induced crosslinking, vacuum-assisted filtration, and freeze-drying. The Ti$_3$C$_2$Tx/CA aerogel film with a thickness of 26 µm presented excellent EMI SE (54.3 dB), owing to its sponge-like structure, which facilitated the dissipation of incident EM waves through multi-reflection and scattering in the Ti$_3$C$_2$Tx/CA aerogel film [85].

Liu et al. fabricated a polyurethane/Ti$_3$C$_2$Tx MXene film vacuum-assisted filtration [86], thanks to the bioinspired material design and the careful choice of polyurethane as a
polymer matrix. The polyurethane/Ti$_3$C$_2$T$_x$ with nacre-like structure showed superior electric conductivity of $\approx$ 2897.4 S cm$^{-1}$ and SEE of 33,771.92 dB cm$^2$ g$^{-1}$ with ultra-small thickness (< 10 µm).

A polyvinyl alcohol/Ti$_3$C$_2$T$_x$ (PVA/Ti$_3$C$_2$T$_x$) film with alternating multilayered structure was prepared by multilayered casting [87]. When the amount of Ti$_3$C$_2$T$_x$ was 19.5 wt%, PVA/Ti$_3$C$_2$T$_x$ multilayered film with a thickness of 27 µm displayed electrical conductivity of 716 S m$^{-1}$, EMI SE of 44.4 dB, and the SEE$_t$ of 9343 dB cm$^2$ g$^{-1}$. This excellent performance was due to the improved multiple interfacial reflection and improved absorption in the MXene layer.

Liu et al. prepared a chitosan (CS)/Ti$_3$C$_2$T$_x$ film by vacuum-assisted filtration [88]. The CS/Ti$_3$C$_2$T$_x$ film with the Ti$_3$C$_2$T$_x$ content of 75 wt% displayed high EMI shielding performance of $\approx$ 34.7 dB at the thickness of 13 µm, which was attributed to the outstanding electrical conductivity ($\approx$ 1402 ± 70 S m$^{-1}$) and multiple internal reflection.

A Ca ion cross-linked sodium alginate (SA)-montmorillonite (MMT)/Ti$_3$C$_2$T$_x$ MXene (CSA-M-T) film was fabricated by a step-by-step vacuum-assisted filtration process [89]. Compared with the pure Ti$_3$C$_2$T$_x$ layer, such kind of sandwich film can effectively maintain the EMI shielding performance (50.01 dB).

### 4.3.3 Inorganic Hybrid

#### 4.3.3.1 Carbon-Based

A assemble Ti$_3$C$_2$T$_x$ MXene-carbon nanotube (CNT) composite film was fabricated by spin spray layer-by-layer (LbL) [90]. The absolute effectiveness of the film was up to 58,187 dB cm$^2$ g$^{-1}$, which was due to the both the excellent electrical conductivity (130 S cm$^{-1}$) and the enhanced absorption with the LbL architecture of the films.

Xiang et al. prepared a lightweight and ultrathin TiO$_2$-Ti$_3$C$_2$T$_x$/graphene film with the range of 5.25–9.17 µm of thickness by vacuum filtration and pyrolysis [91]. The film displayed surface resistance of 7.5 Ω sq$^{-1}$ and EMI SE of 27 dB. Meanwhile, the value of SEE$_t$ of the film reached 30,291.43 dB cm$^2$ g$^{-1}$.

A stretchable Ti$_3$C$_2$T$_x$ nanosheet/single-walled carbon nanotube (SWNT) film was fabricated by Li et al. by pre-buckling of SWNT and Ti$_3$C$_2$T$_x$ nanosheet coatings on latex substrates [50]. A single layer of stretchable Ti$_3$C$_2$T$_x$/SWNT film demonstrated a strain-invariant EMI shielding performance of $\approx$ 30 dB up to 800% areal strain.

Zhang’s group investigated the EMI shielding performance of Ti$_3$C$_2$T$_x$ MXene-GO film [92]. Thanks to the high electronic conductivity, the Ti$_3$C$_2$T$_x$/GO film with a small film thickness of 7 µm displayed the EMI SE of 50.2 dB. Compared with other shielding materials, the MXene-GO films were obviously superior in combining excellent EMI shielding performance and good tensile strength.

A Ti$_3$C$_2$T$_x$-bonded carbon black (CB) film with a porous structure was fabricated by a vacuum-assisted filtration method [93]. With the incorporation of 70 mg of CB, the Ti$_3$C$_2$T$_x$-bonded CB film showed EMI shielding performance of 60 dB with a SE$_A$ of 15 dB and SE$_R$ of 45 dB. Moreover, the SEE$_t$ reached 8718 dB cm$^2$ g$^{-1}$. Research showed that the porous structure could improve the absorption, resulting from enhanced scattering and reflection.

#### 4.3.3.2 Others

A aluminum ion-reinforced Ti$_3$C$_2$T$_x$ MXene (Al-Ti$_3$C$_2$T$_x$) film was fabricated by Zhang group via vacuum filtration method [94]. The Al-Ti$_3$C$_2$T$_x$ film displayed a high conductivity of 265,600 S m$^{-1}$. The strong and highly conductive MXene film with a small thickness of 39 mm showed EMI shielding performances of over 80 dB in the X-band.

Ning et al. investigated the EMI shielding performance of a Mn ion-intercalated Ti$_3$C$_2$T$_x$ (MIT) film [95]. The MIT film showed an average electronic conductivity of 4268 S m$^{-1}$, which was two times than that of pure Ti$_3$C$_2$T$_x$ film (1894 S m$^{-1}$). The MIT film with a thickness of 2.5 µm showed enhanced performance of 44.3 dB compared with pure Ti$_3$C$_2$T$_x$ film (24.1 dB), owing to the additive internal absorption.

A sliver nanowire (AgNW)/Ti$_3$C$_2$T$_x$ film was fabricated by a pressured-extrusion film-forming process [96]. The MXene/AgNW composite film with a low loading of nanocellulose (0.167 wt%) showed high electrical conductivity of $\approx$ 30,000 S m$^{-1}$, and remarkable SEE$_t$ of 16,724 dB cm$^2$ g$^{-1}$.

A Ti$_3$C$_2$T$_x$/montmorillonite (MMT) film was fabricated by a simple vacuum-assisted filtration technique [97]. The EMI shielding performance of Ti$_3$C$_2$T$_x$/MMT film with different concentration ratios was investigated. The composite film with 10 wt% MMT showed high electrical conductivity (4420 S m$^{-1}$), EMI SE of 65 dB in the entire X-band.
and SEE of over 10,000 dB cm² g⁻¹ at a thickness of only 25 µm.

### 4.4 Foam

#### 4.4.1 Pristine

The inceptive study of the EMI shielding performance of pure MXene foam was reported by Yu’s group [98]. They used an efficient and facile method to prepare free-standing, flexible, and hydrophobic Ti₃C₂Tx MXene foam (Fig. 6a–g). In striking contrast to well-known hydrophilic MXene materials, the Ti₃C₂Tx foams surprisingly had hydrophobic surfaces, with outstanding water resistance and durability. Thanks to the highly efficient wave attenuation in the favorable porous structure, the lightweight Ti₃C₂Tx foam showed enhanced EMI shielding performance of 70 dB compared with its unfoamed film counterpart (53 dB) (Fig. 6h–j).

#### 4.4.2 Hybrid

A porous 3D Ti₃C₂Tx MXene/C hybrid foam (MCF) was prepared by sol–gel followed by thermal reaction [99]. And, the MCF/epoxy was obtained via vacuum-assisted impregnation followed by curing process. The MCF/epoxy with 4.25 wt% MCF displayed the EMI shielding performance of 46 dB and electrical conductivity of 184 S/m, which was 4.8 and 3.1 × 10⁴ times higher than that of MCF-0/epoxy nanocomposites (without Ti₃C₂Tx MXene), respectively.

A porous few-layered Ti₂CTₓ (f-Ti₂CTₓ) MXene/poly (vinyl alcohol) (PVA) composite foam was fabricated by a
facile freeze-drying method [100]. The f-Ti$_2$CT$_x$/PVA foam with a content of only 0.15 vol% afforded a SEE of 5136 dB cm$^2$ g$^{-1}$. Such excellent EMI shielding performance was attributed to the multi-porous structure, internal reflection, and polarization effect.

A lightweight Ti$_3$C$_2$Tx MXene/graphene (Ti$_3$C$_2$Tx-GO) hybrid foam was fabricated by freeze-drying and reduction heat treatment [101]. Thanks to the improved foam electrical conductivity and highly efficient wave attenuation in interconnected porous structures, the Ti$_3$C$_2$Tx-GO hybrid foam showed excellent EMI shielding performance of 50.7 dB and specific EMI shielding effectiveness of 6217 dB cm$^3$ g$^{-1}$, which was much higher than that most of the EMI shielding materials.

Silver nanowires (AgNWs)/Ti$_3$C$_2$T$_x$ foam was fabricated by integrating AgNWs as the skeleton and Ti$_3$C$_2$T$_x$ as the covering decoration for foaming structure [102]. The AgNWs/Ti$_3$C$_2$T$_x$ foam showed EMI shielding performance of 41.3 dB at the thickness of 1.2 µm in the X-band. The freespace created during foaming helped to obtain EM wave scattering within the skin depth.

![Fig. 7](https://example.com/fig7.png)

**Fig. 7** SEM images of Ti$_3$C$_2$T$_x$ aerogels with the densities of a 5.5, b 8.3, and c 11.0 mg cm$^{-3}$, respectively. Interlayer synapses and bridges are circled in red. d Frequency-dependent EMI SE of different MXene aerogels at a density of 11.0 mg cm$^{-3}$ and a compression thickness of 1 mm. e The average EMI SE of Ti$_3$C$_2$T$_x$ aerogels with different densities. Reproduced with permission from Ref. [104]. Copyright 2019, WILEY-VCH. f SEM images of the MXene/CNF hybrid aerogels with various angles between the cell walls’ oriented direction and electric field direction of incident EM waves. g EMI shielding performance at a fixed frequency of 10 GHz for the MXene/CNF hybrid aerogels (4 mg cm$^{-3}$) with various CNF contents at cell wall-electric field angles of 0° and 90°, respectively. h Comparison of the MXene/CNF hybrid aerogels’ shielding performance with other materials. Reproduced with permission from Ref. [106]. Copyright 2020, WILEY-VCH
A polydimethylsiloxane (PDMS)-coated Ti$_3$C$_2$T$_x$ MXene foam was fabricated by using the Ti$_3$C$_2$T$_x$ assisted with sodium alginate (SA) as template followed by coating a thin layer of PDMS [103]. The Ti$_3$C$_2$T$_x$/SA/PDMS foam with 95 wt% Ti$_3$C$_2$T$_x$ exhibited an excellent conductivity of 2211 s m$^{-1}$ and EMI shielding performance of 70.5 dB. Moreover, the foam with 74 wt% Ti$_3$C$_2$T$_x$ displayed EMI SE of 48.2 dB after 500 compression–release cycles.

4.5 Aerogel

4.5.1 Pristine

Han et al. fabricated three types of porous MXene aerogels (Ti$_3$C$_2$T$_x$, Ti$_2$CT$_x$, and Ti$_3$CNT$_x$) via a bidirectional freeze-casting technique (Fig. 7a–c) [104]. The EMI SE of Ti$_3$C$_2$T$_x$, Ti$_2$CT$_x$, and Ti$_3$CNT$_x$ aerogels reached 70.5, 69.2, and 54.1 dB at the thickness of 1 mm (Fig. 7d, e), respectively. Especially, the SEE$_t$ of Ti$_2$CT$_x$ aerogel with a density of 5.5 mg cm$^{-3}$ and a thickness of 1 mm reached 8818.2 dB cm$^2$ g$^{-1}$, which was several times higher than that of other materials. Meanwhile, Bian et al. prepared an ultralight Ti$_3$C$_2$T$_x$ aerogel by the freeze-drying method [105]. The Ti$_3$C$_2$T$_x$ aerogel with density of 6.26 mg cm$^{-1}$ exhibited electrical conductivity of 22 S cm$^{-1}$ and SEE$_t$ of 9904 dB cm$^3$ g$^{-1}$. The excellent EMI shielding performance was attributed to the high electrical conductivity and porous structures.

4.5.2 Hybrid

A cellulose nanofibril (CNF)/Ti$_3$C$_2$T$_x$ aerogel was fabricated via an ice-templated freeze-casting approach (Fig. 7f) [106]. The Ti$_3$C$_2$T$_x$ MXene “bricks” bonded by CNF “mortars” of the nacre-like cell walls induced high electrical conductivity, and interfacial polarization led to excellent EMI shielding performance. The CNF/Ti$_3$C$_2$T$_x$ aerogel with ultralow density showed EMI SE up to 74.6 dB, SEE$_t$ of as 30,660 dB cm$^3$ g$^{-1}$, and SEE$_t$ achieving 189,400 dB cm$^2$ g$^{-1}$ (Fig. 7g, h), exceeding that of other MXene-based or other shielding architectures reported so far.

Zhang et al. fabricated a 3D Ti$_3$C$_2$T$_x$ MXene/reduced graphene oxide (RGO) hybrid aerogel by directional freezing and freeze-drying [107]. The Ti$_3$C$_2$T$_x$/RGO hybrid aerogel with aligned cellular microstructure displayed a high electrical conductivity of 1085 S cm$^{-1}$ and an excellent EMI shielding performance of 50 dB in the X-band at a low Ti$_3$C$_2$T$_x$ content of 0.74 vol%, which was the best results among polymer nanocomposites with similar loading of Ti$_3$C$_2$T$_x$.

Koo’s group fabricated a 3D porous Ti$_3$C$_2$T$_x$/carbon nanotube (CNT) hybrid aerogel by a bidirectional freezing method [108]. The Ti$_3$C$_2$T$_x$/CNT aerogel showed excellent electrical conductivity of 9.43 S cm$^{-1}$ and superior EMI shielding performance of 103.9 dB at 3 mm thickness over the X-band frequency. The excellent EMI shielding performance of the Ti$_3$C$_2$T$_x$/CNT hybrid aerogel was attributed to the 3D porous structure with a high-conducting and uniform lamellar structure.

Liang et al. prepared a Ti$_3$C$_2$T$_x$/wood-derived porous carbon (WPC) aerogel via freeze-drying procedure [109]. Such wall-like “mortar-brick” structures profoundly prolong the transmission paths of the EM waves and dissipate the incident EM waves in the form of heat and electric energy, thereby exhibiting the superior EMI shielding performance. The Ti$_3$C$_2$T$_x$/WPC aerogel showed EMI SE value of 71.3 dB at density as low as 0.197 g cm$^{-3}$.

4.6 Fabric

MXene fabric usually refers to MXene-coated fabric. The main fabrics used for this purpose are cotton or polymers, although other materials are also utilized.

4.6.1 Cotton

Geng et al. prepared Ti$_3$C$_2$T$_x$ coated cotton fabrics with low Ti$_3$C$_2$T$_x$ loading (1.5–2.6 mg cm$^{-2}$) through a facile vacuum filtration process [110]. The fabric with Ti$_3$C$_2$T$_x$ loading of 2.6 mg cm$^{-2}$ showed SEE$_t$ of 2969 dB cm$^2$ g$^{-1}$. Zhang et al. reported that the Ti$_3$C$_2$T$_x$ modified fabric with a low Ti$_3$C$_2$T$_x$ loading 6 wt% exhibited excellent electrical conductivity of
5 Ω sq⁻¹ and outstanding EMI shielding performance (up to 36 dB) [111]. Cheng et al. fabricated a Ti₃C₂Tx MXene-coated cotton fabric by a simple solution impregnation and dip-coating method [112]. The cotton fabric coated by rising amount of Ti₃C₂Tx could improve the EMI shielding performance. When the amount of Ti₃C₂Tx was 5.2 mg cm⁻², the fabric afforded excellent electrical conductivity of 670.3 S m⁻¹ and EMI SE of 31.04 dB in the X-band (Fig. 8a, b). Moreover, the EMI shielding performance of the fabric was almost not changed after 800 bending times (Fig. 8c, d).

4.6.2 Polymer

A flexible and durable cellulose/Ti₃C₂Tx MXene nanocomposite fabric was fabricated by a simple dip-coating method [113]. The fabric with a Ti₃C₂Tx nanosheet loading of 1.89 vol% displayed an outstanding electrical conductivity of 2756 S m⁻¹ (Fig. 8e). After a polydimethylsiloxane (PDMS) coating, EMI SE of the fabric could achieve over 43 dB in the X and Ku band (Ti₃C₂Tx loading of 1.07 vol%, and no apparent decline was observed after 2000 bending–releasing cycles in the durability test (Fig. 8f).

Zhang’s group fabricated a Ti₃C₂Tx MXene-decorated polyester fabric (Ti₃C₂Tx-fabric) by depositing in situ polymerized polypyrrole (PPy)-modified Ti₃C₂Tx nanosheets onto poly (ethylene terephthalate) fabric, followed by silicone coating [114]. The modified fabric displayed a high electrical conductivity of 1000 S m⁻¹ and EMI SE of 90 dB, with a thickness of 1.3 mm. Benefiting from the contribution of PPy-to-EM wave absorption due to introduction of the polar group, the EMI shielding performance of the Ti₃C₂Tx-fabric was better than that of the fabric modified by MXene at similar conductivities.

Yuan et al. prepared a flexible and stretchable Ti₃C₂Tx MXene/polyurethane (PU) fabric [115]. The Ti₃C₂Tx/PU fabric with sandwich structure exhibited EMI SE of ~20 dB at a stretching process within 30% deformation.

Poylaniline (PANI)/Ti₃C₂Tx/carbon fiber (CF) fabric was fabricated based on the LbL assembly approach [116]. The fabric with a thickness of 0.55 mm possessed a high EMI shielding performance of 26 dB, SSE of 135.5 dB cm³ g⁻¹ and electrical conductivity of 24.57 S m⁻¹.
4.6.3 Others

Yu’s group prepared the silk fabric with biomimetic leaf-like MXene/silver nanowire by depositing in situ polymerized PPy-modified Ti$_3$C$_2$Tx MXene sheets onto poly (ethylene terephthalate) textiles followed by a silicone coating [117]. The flexible fabric displayed a low sheet resistance of 0.8 $\Omega$ sq$^{-1}$, excellent EMI shielding performance of 54 dB in the X-band at the thickness of 120 $\mu$m.

Ti$_3$C$_2$Tx MXene-decorated wood-pulp fabric was fabricated by depositing highly conductive Ti$_3$C$_2$Tx networks onto wood-pulp fabric grid (FG) followed by hydrophobic methyltrimethoxysilane (MTMS) coating with multi-scaled roughness via a simple vacuum-filtration approach and sol–gel process [118]. The fabric possessed superior EMI SE up to ~57.8–90.2 dB.

5 MXene-Based Materials for EM Wave Absorption

5.1 Pure MXene Matrix

5.1.1 Non-annealing

The earliest research was concerned with the EM wave-absorbing properties of multilayer Ti$_3$C$_2$Tx. Qing et al. prepared multilayer Ti$_3$C$_2$Tx MXene by etching Ti$_3$AlC$_2$...
with 50% HF for 3 h (Fig. 9a–d) [119]. Compared with the Ti$_3$AlC$_2$/wax, the multilayer Ti$_3$C$_2$T$_x$/wax showed high EM wave absorption at the same filling concentration of 50 wt% (Fig. 9e). This result was due to the unique two-dimensional (2D) morphology of multilayer Ti$_3$C$_2$T$_x$, MXene, such as the large number of defects and larger internal boundary layer capacitance. Feng et al. further confirmed that the excellent EM wave-absorbing property of multilayer Ti$_3$C$_2$T$_x$/wax was due to the high dielectric loss and the strong multi-reflections [120]. Luo et al. studied the influence of different etching time on the EM wave absorption of multilayer Ti$_3$C$_2$T$_x$ etched for 24 h afforded the best EM wave-absorbing properties among etching sample. A minimum RL value of − 42.5 dB (40 wt% HF etching). The Ti$_3$C$_2$T$_x$ showed excellent wave-absorbing properties. They found that multilayer Ti$_3$C$_2$T$_x$ further treated in ethanol showed much more superior absorption capability. This result could be due to the enlarged interlayer spacing, and increased surface functional groups after ethanol-based solvothermal treatment.

The influence of different etching time on the EM wave-absorbing performance of multilayer Ti$_3$C$_2$T$_x$ has also been studied. Tong et al. investigated the effect of different etching time in 40 wt% HF acid on the EM wave absorption of multilayer Ti$_3$C$_2$T$_x$, MXene (0, 6, 24, 48, 96, 144, and 192 h) [122]. With the increase of etching time, the morphology of Ti$_3$C$_2$T$_x$ was destroyed gradually. Ti$_3$C$_2$T$_x$ etched for 24 h afforded the best EM wave-absorbing properties among etching sample. A minimum RL value of − 42.5 dB was achieved at the thickness of 1.7 mm. This result could be attributed to multiple reflections between MXene layers and interfacial polarizations. Zhao’s group etched the multilayer Ti$_3$C$_2$T$_x$ in ≥ 40 wt% HF acid for different time (1, 2, and 3 h) [123] and then explored their EM wave absorption. Multilayer Ti$_3$C$_2$T$_x$ etched for 3 h (Ti$_3$C$_2$T$_x$-3) had excellent EM wave-absorbing property. Forty percentage ratio multilayer Ti$_3$C$_2$T$_x$-3/wax showed a minimum RL value of −36.3 dB with a thickness of 4.5 mm. Cui et al. produced multilayer Ti$_3$C$_2$T$_x$ etched by HCl/LiF with different diverse etching times (12, 24, 36, 48, and 60 h) [124] and studied their EM wave absorbing properties. They found the same result as HF etching. The Ti$_3$C$_2$T$_x$ etched 24 h showed best EM wave absorption, owing to multilayer scattering between the laminate structures.

The influence of different etchants on the EM wave-absorbing properties was reported by Xu’s group [125]. Multilayer Ti$_3$C$_2$T$_x$, MXenes were obtained by ultrasonication-solvothermal treatment in different solvents including dimethylformamide (DMF), ethanol, and dimethyl sulfoxide (DMSO), respectively. Research showed that multilayer Ti$_3$C$_2$T$_x$ treated with DMF showed excellent wave-absorbing properties due to the larger layer space and diminished oxidation effects.

Cao’s group was firstly investigated the EM wave absorption performance of the delaminated Ti$_3$C$_2$T$_x$ (d-Ti$_3$C$_2$T$_x$) nanosheet etched by HCl/LiF (Fig. 9f, g) [44]. All Ti$_3$C$_2$T$_x$ nanosheet/wax composites with different concentrations showed excellent EM wave absorption (Fig. 9h–k). Especially for 40 wt% composite, a minimum RL value of −47.9 dB and a corresponding absorption bandwidth of 3.6 GHz were achieved at a thickness of 2.5 mm. Moreover, they found the transformation mechanism between EM energy and thermal energy in the composite. The higher the concentration of delaminated Ti$_3$C$_2$T$_x$ nanosheet in the composite, more was the conversion of EM energy to thermal energy. Yin’s group originally reported that the multilayer Nb$_2$CT$_x$, MXene by 49 wt% HF etching and solvothermal/hydrothermal treatment [126]. They found that multilayer Nb$_2$CT$_x$, further treated in ethanol showed much more superior absorption capability. This result could be due to the enlarged interlayer spacing, and increased surface functional groups after ethanol-based solvothermal treatment.

5.1.2 Annealing

Yin’s group originally reported that the multilayer Ti$_3$C$_2$T$_x$, MXene annealed at 800 °C for 2 h in Ar atmosphere had excellent EM wave-absorbing properties [58]. Such excellent EM wave-absorbing properties were due to the surface functional groups of MXene modified by annealing. The annealing led to the formation of a local sandwich structure composed of TiO$_2$ nanocrystals and amorphous carbon, which enhanced the EM wave absorption. After that, they investigated the EM wave-absorbing properties of multilayer Ti$_3$C$_2$T$_x$ at different annealing temperatures (600, 700, and 800 °C) for 1 h in CO$_2$ [127]. The multilayer Ti$_3$C$_2$T$_x$ annealed at 800 °C showed best EM wave absorption performance with its RL value achieving −36 dB and absorption bandwidth of 5.6 GHz. Meanwhile, they explored the EM wave absorption performance of multilayer Ti$_3$C$_2$T$_x$ annealed at 500, 800, and 900 °C for 1 h in CO$_2$ [128]. The microwave absorption of multilayer Ti$_3$C$_2$T$_x$, annealed at 800 °C was best, which was due to the enhanced polarization loss and stronger conduction loss.
Fan et al. fabricated the multilayer Ti$_3$C$_2$T$_x$, and annealed it in O$_2$ at different temperatures (100, 200, 300, 400, and 500 °C) for 2 h [129]. The multilayer Ti$_3$C$_2$T$_x$ calcined at 100 °C showed excellent EM wave-absorbing properties, with a minimum RL value of −40.07 dB at 19.2 GHz and the absorption bandwidth of 3.8 GHz. This result was attributed to the appropriate complex permittivity and matching impedance.

### 5.2 MXene Hybrid Matrix

EM wave absorption can be improved by increasing the magnetic loss or dielectric loss. The magnetic loss of the absorber can be improved by doping with magnetic materials. Certain carbon-based materials with high conductivity can be doped into the absorber to enhance the dielectric loss.

#### 5.2.1 Magnetic Hybrid

**5.2.1.1 Fe-Based** Liu et al. investigated the EM wave absorption of the multilayer Ti$_3$C$_2$T$_x$ doped with different concentrations of Fe$_2$O$_4$ (3, 5, and 10 wt%) (Ti$_3$C$_2$T$_x$-3, Ti$_3$C$_2$T$_x$-5, Ti$_3$C$_2$T$_x$-10) [130]. By tuning the doping concentration of Fe$_2$O$_4$, the sample showed improved microwave absorption performance. Among them, Ti$_3$C$_2$T$_x$-10/wax showed excellent absorption performance, with a maximum RL value of −57.3 dB. Zhao et al. further investigated the EM wave absorption of multilayer Ti$_3$C$_2$T$_x$/Fe$_2$O$_4$ [131]. The multilayer Ti$_3$C$_2$T$_x$/Fe$_2$O$_4$ exhibited enhanced EM wave absorption compared with pure multilayer Ti$_3$C$_2$T$_x$, which is due to the outstanding impedance matching and efficient attenuation. Yang’s group studied the EM wave-absorbing properties of multilayer Ti$_3$C$_2$T$_x$ doped with Fe$_2$O$_4$ nanoparticles (Fe$_2$O$_4$@Ti$_3$C$_2$T$_x$) with different concentrations [132]. The sample contained 25 wt% Fe$_2$O$_4$ nanoparticles displayed outstanding EM wave absorption, with a minimum RL value of −57.2 dB at 15.7 GHz and bandwidth of 1.4 GHz, caused by enhanced interface polarization. Che’s group fabricated magnetized multilayer Ti$_3$C$_2$T$_x$ MXene microsphere by embedded Ti$_3$C$_2$T$_x$ MXene into a confined and magnetized Fe$_2$O$_4$ nanospheres (designated as M/F) [133]. This structure could enhance the specific interfaces and dielectric polarization. Meanwhile, these Fe$_2$O$_4$ magnetic led to the optimized impedance balance and EM coordination capability. As expected, the M/F composite with 15 wt% Fe$_2$O$_4$ content hold distinct EM wave absorption property with the strong reflection loss (−50.6 dB) and absorption bandwidth (4.67 GHz) at the thickness of 2 mm.

The multilayer Ti$_3$C$_2$T$_x$/flaky carbonyl iron (FCI) composite with different mass ratios were fabricated by the ultrasonic mixing method [134]. An excellent EM wave-absorbing properties can be realized by optimizing the Ti$_3$C$_2$T$_x$ and FCI content. Beneficial from the good impedance matching and moderate attenuation ability, the composite with 20 wt% Ti$_3$C$_2$T$_x$ and 40 wt% FCI loading presented the absorption bandwidth of 8.16 GHz with a thickness of 1.0 mm.

**5.2.1.2 Ni-Based** A Ti$_3$C$_2$T$_x$/Ni-nanoparticle hybrid was synthesized by in situ hydrothermal treatment. The Ti$_3$C$_2$T$_x$-Ni hybrid showed a RL value of −47.06 dB with a thickness of 1.5 mm and bandwidth of 3.6 GHz [135]. The combined effect of magnetic loss, conduction loss and dielectric loss is the key to achieving such excellent EM-absorbing ability. Liang et al. prepared a Ti$_3$C$_2$T$_x$ MXene/ Ni-nanochain (Ni@MXene) hybrid via a facile and moderate co-solvothermal method [36]. The Ni@MXene hybrid displayed a minimum RL of −49.9 dB at the thickness of 1.75 mm when the Ni-nanochain content was 90 wt%. It was further proved that the synergistic effect of conductive MXene and the magnetic Ni-nanochain led to the excellent EM wave-absorbing ability. Che’s group also investigated the EM wave-absorbing properties of a multilayer Ti$_3$C$_2$T$_x$/Ni hybrid [136]. The Ni nanoparticles were uniformly distributed on the surface and in the multilayered gaps of Ti$_3$C$_2$T$_x$. This unique structure led to excellent EM wave-absorbing properties. The hybrid showed a minimum RL of −50.5 dB at 5.5 GHz. Liang et al. investigated the EM wave-absorbing properties of Ni-, Co- and NiCo-doped multilayer Ti$_3$C$_2$T$_x$ [137]. Among them, multilayer Ti$_3$C$_2$T$_x$ doped with Ni nanoparticles (Ni@Ti$_3$C$_2$T$_x$) in a polyvinylidene fluoride (PVDF) matrix showed strong EM wave absorption. With 10 wt% Ni dopant, the sample exhibited the optimal EM wave absorption, with a minimum RL of −52.6 dB at 8.4 GHz and absorption bandwidth of 3.7 GHz. Liu et al. found that compared with the individual Ti$_3$C$_2$T$_x$ and Ni powders [138], hybrid Ti$_3$C$_2$T$_x$/Ni afforded the most favorable EM wave absorption performance with a minimum RL value of −24.3 dB at 9.8 GHz.

**5.2.1.3 Co-Based** Deng et al. fabricated the Co$_3$O$_4$/Ti$_3$C$_2$T$_x$ by the two-step method. When the mass ratio of Ti$_3$C$_2$T$_x$ to Co$_3$O$_4$ was 1:3 [139], the Co$_3$O$_4$/Ti$_3$C$_2$T$_x$ hybrid showed better than 90% absorption from 10.8 to 17 GHz. Such excellent performance is owing to combined effects of multilayer structure, defects, conductivity of Ti$_3$C$_2$T$_x$ and equivalent capacitance of Co$_3$O$_4$. 

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5.2.1.4 Multi-based A CoFe@Ti3C2Tx hybrid was prepared by in situ reduction, and a minimum RL value of −36.29 dB could be obtained with a thickness of 2.2 mm [140]. The excellent EM wave absorption performance was due to the sandwich-like structure and enhanced interfacial polarization. He et al. further confirmed that multilayer Ti3C2Tx modified by CoFe could improve the EM wave absorption.

A FeCo@Ti3C2Tx hybrid was fabricated by in situ hydrothermal treatment [141]. The incorporation of magnetic FeCo could improve the EM wave-absorbing property. The FeCo@Ti3C2Tx hybrid exhibited a broad EM wave-absorbing bandwidth of 8.8 GHz, due to enhanced impedance matching and microwave attenuation.

A Co-doped NiZn ferrite (CNZF)/polyaniline (PANI) on Ti3C2Tx hybrid (CNZF/PANI/Ti3C2Tx) was synthesized by hydrothermal reaction and interfacial polymerization [142]. The dipole polarization, interfacial polarization, natural resonance, eddy current loss, and multiple reflections contributed to the improved EM wave absorption performance of CNZF/PANI/Ti3C2Tx hybrid. The multiple-layer hybrid exhibited excellent EM wave absorption with a minimum RL of −37.1 dB and absorption bandwidth of 4.1 GHz.

Hou et al. investigated the EM wave-absorbing properties of NiCo2O4-doped multilayer Ti3C2Tx at different annealing temperatures (350, 400, 450, and 500 °C) for 2 h in argon [143]. The NiCo2O4/Ti3C2Tx annealed at 350 °C displayed the best EM wave absorption, with the RL value of −50.96 dB. This result was attributed to the polarization behavior and multiple scattering produced by unique structures.

The EM wave-absorbing properties of Ba3Co2Fe24O41, multilayer Ti3C2Tx, and polyvinyl butyral (PVB) after physical mixing were investigated by Yang’s group [144]. The as-synthesized PVB/Ba3Co2Fe24O41/Ti3C2Tx exhibited outstanding and efficient EM wave attenuation. A minimum RL value for the PVB/Ba3Co2Fe24O41/Ti3C2Tx composite reached −46.3 dB at 5.8 GHz; the absorption bandwidth was 1.6 GHz, with a thickness of only 2.8 mm.

5.2.2 Carbon-Based Hybrid

A graphite/TiC/Ti3C2Tx (G/TiC/Ti3C2Tx) hybrid was obtained by two steps [145]. Firstly, the graphite/TiC/Ti3AlC2 (G/TiC/Ti3AlC2) hybrid was prepared in a bath of molten salts. G/TiC/Ti3C2Tx was obtained after etching the Al atoms from G/TiC/Ti3AlC2 (Fig. 10a–c). It was found that the graphite/TiC/Ti3AlC2-wax matrix with a thickness of 2.1 mm exhibited a minimum RL of −63 dB and the effective absorption bandwidth was more than 3.5 GHz (Fig. 10d, e).

Multilayer Ti3C2Tx MXene modified with in situ grown carbon nanotubes (Ti3C2Tx/CNT) was fabricated by Yin’s group (Fig. 10f, g) [146]. Compared with the pure multilayer

![Fig. 10](image-url) SEM image of a-c G/TiC/Ti3C2Tx. d RC of G/TiC/Ti3C2Tx, Ti3C2Tx, G/TiC/Ti3AlC2, G/TiC, and G/TiC+Ti3C2Tx in paraffin matrix with the loading of 50 wt% with the sample thickness of 2.1 mm. e 3D RC of G/TiC/Ti3C2Tx versus frequency and sample thickness. Reproduced with permission from Ref. [145]. Copyright 2018, WILEY–VCH. SEM images of f and g Ti3C2Tx/CNT. h 3D representations and i theoretical curves of RC versus frequency and thickness of Ti3C2Tx/CNT with a filler loading of 35 wt%. Reproduced with permission form Ref. [146]. Copyright 2017, The Royal Society of Chemistry
Ti$_3$C$_2$T$_x$ MXenes, the hierarchical microstructure makes a contribution to the outstanding EM wave absorption performance, with a minimum $RL$ value of $-52.9$ dB and absorption bandwidth of 4.46 GHz (Fig. 10h, i).

Dai et al. reported that the Ti$_3$C$_2$Tx MXenes/nano-carbon sphere hybrid exhibited $RL$ of $-54.67$ dB at 3.97 GHz [147], owing to the unique structure of Ti$_3$C$_2$T$_x$ MXenes/nano-carbon and the formation of a heterogeneous interface structure.

Nitrogen-doped graphene (N-GP) and Ti$_3$C$_2$T$_x$ composites were prepared by Qing group [148]. The values and frequency dependencies of EM properties of N-GP/Ti$_3$C$_2$T$_x$ could be tuned by the combination of the unique structure and dielectric characteristics of the N-GP and Ti$_3$C$_2$T$_x$. A minimum $RL$ of the N-GP/Ti$_3$C$_2$T$_x$ composite reached up to $-52$ dB, and absorption bandwidth could be obtained in the frequency range of 10.9–18 GHz with a thickness of only 1.4 mm.

### 5.2.3 Others Hybrid

Qian et al. prepared an urchin-like ZnO–Ti$_3$C$_2$T$_x$ hybrid through a coprecipitation process [149]. Compared with pure multilayer Ti$_3$C$_2$T$_x$, ZnO–Ti$_3$C$_2$T$_x$ hybrid showed significant enhanced EM wave absorption. The minimum $RL$ of 75 wt% ZnO–Ti$_3$C$_2$T$_x$/wax realized $-26.30$ dB, which is much better than that of pure multilayer Ti$_3$C$_2$T$_x$ ($-6.70$ dB), owing to larger interfaces and the construction of semiconductive networks.

Ti$_3$C$_2$T$_x$ MXenes/polyaniline microspheres (Ti$_3$C$_2$T$_x$/PPy) composites with delaminated structure were fabricated for significant enhancement of EM wave-absorbing properties [150]. Thanks to the synergistic effect between Ti$_3$C$_2$T$_x$ and PPy microspheres, the obtained Ti$_3$C$_2$T$_x$@PPy composite exhibited excellent EM absorption performance. The 10 wt% Ti$_3$C$_2$T$_x$@PPy composites in wax matrix displayed a minimum $RL$ of $-49.5$ dB at 7.6 GHz, and the absorption bandwidth was 5.14 GHz. Tong et al. reported that multilayer Ti$_3$C$_2$T$_x$ MXene decorated with PPy chains is a good microwave-absorbing material [151]. The 25 wt% Ti$_3$C$_2$T$_x$@PPy hybrids in a wax matrix showed a minimum $RL$ of $-49.2$ dB.

A multilayer Ti$_3$C$_2$T$_x$ modified by MoS$_2$ was fabricated by a hydrothermal method [152]. The complex permittivity of MoS$_2$/Ti$_3$C$_2$T$_x$ increased compared with that of multilayer Ti$_3$C$_2$T$_x$. This result led to enhanced EM wave-absorbing performance. When the thickness of MoS$_2$/Ti$_3$C$_2$T$_x$-wax was 2.5 mm, the corresponding absorption bandwidth was 2.6 GHz.

A multilayer Ti$_3$C$_2$T$_x$ MXene/polyaniline (PANI) was prepared by the hydrothermal reaction [153]. The EM wave absorption of sample with different PANI doping concentrations was systematically studied. When the mass ratio of Ti$_3$C$_2$T$_x$ to polyaniline is 1:2, the sample showed best EM wave absorption. The maximum $RL$ reached $-56.30$ dB at 13.80 GHz with a thickness of 1.8 mm.

The Nb$_2$O$_5$ with different morphologies was prepared in situ and implanted between the layers of the Nb$_2$CT$_x$ MXene via hydrothermal method [154]. The Nb$_2$O$_5$/Nb$_2$CT$_x$ exhibited the enhanced EM wave absorption compared with primary Nb$_2$CT$_x$, MXene. Especially, Nb$_2$CT$_x$ doped with columnar Nb$_2$O$_5$ showed a minimum $RL$ of $-44.1$ dB at 2.8 GHz, owing to the increased lamellar spacing of the Nb$_2$CT$_x$.

### 5.3 Foam

Yin’s group fabricated reduced graphene oxide (RGO)/Ti$_3$C$_2$T$_x$ hybrids foam via self-assembly and sacrificial template processes [155]. The RGO/Ti$_3$C$_2$T$_x$ foam with the density is merely 0.0033 g cm$^{-3}$ possessed outstanding EM absorption performance superior to all reported foam-based counterparts, and the absorption bandwidth covers the whole X-band at 3.2 mm, and its specific EM absorption performance value exceeds 14,299.2 dB cm$^{-2}$ g$^{-1}$. Those results were attributed to the unique heterogeneous interface associated with core–shell structures. Meanwhile, they synthesized ordered lamellar few-layered Ti$_3$C$_2$T$_x$/SiC nanowires (f-Ti$_3$C$_2$T$_x$/SiCnws) hybrid foams with ultralow density via a combination of self-assembly and bidirectional freezing processes (Fig. 11a–d) [156]. The f-Ti$_3$C$_2$T$_x$/SiCnws hybrid foam showed a minimum $RL$ of $-55.7$ dB at an ultralow density of only about 0.029 g cm$^{-3}$ (Fig. 11e, f). Compared with most of the current foam-based counterparts, the free-standing foams exhibited enhanced EM absorption properties, owing to enhanced polarization loss and balance the conductive loss and impedance matching characteristics caused by the unique microstructure and phase compositions. After that, they prepared porous Ti$_2$CT$_x$ MXene/poly vinyl alcohol (PVA) composite foams constructed by a facile
freeze-drying method (Fig. 11g–j) [100]. Ti$_2$CT$_x$/PVA foam possessed an outstanding EM absorption performance with a minimum $RL$ of $-18.7$ dB and an absorption covering the whole X-band with any thickness from 3.4 to 3.9 mm (Fig. 11k, l).

A new ultralight carbon foam modified by Ti$_3$C$_2$Tx nanosheet (CF/Ti$_3$C$_2$Tx) with three-dimensional network structure was prepared by vacuum impregnation and freeze-drying process [157]. The CF/Ti$_3$C$_2$Tx foam with ultralow density of only 5–7 mg cm$^{-3}$ showed excellent flexibility and steady compression-resilience properties. Meanwhile, the foam showed higher EM absorption than most foam-based EM absorbers, with a minimum $RL$ of $-45$ dB at 8.8 GHz.

5.4 Aerogel

Yang et al. prepared aligned Ti$_3$C$_2$T$_x$ MXene/gelatin (M@G) nanocomposite aerogel using a unidirectional freeze-casting method [158]. The composite aerogel showed a minimum $RL$ of $-59.5$ dB at 14.04 GHz and an absorption bandwidth of 6.24 GHz in the parallel direction but presented a minimum $RL$ of $-57.3$ dB at 4.08 GHz with an absorption bandwidth covering of 3.72–4.56 GHz in the vertical direction. The aerogel exhibited significantly anisotropic EM wave-absorbing properties, owing to the unidirectional aligned microstructure.

Jiang et al. fabricated the hierarchically structured cellulose aerogels with interconnected Ti$_3$C$_2$T$_x$ MXene nanosheet networks via a freeze-casting and chemical cross-linking strategy [159]. The aerogel with a low density (0.31 g cm$^{-3}$) presented a minimum $RL$ of $-43.4$ dB at 11.2 GHz and an absorption bandwidth of 4.5 GHz, which was due to enhanced conductive loss and multiple reflection attenuating.

A TiO$_2$/Ti$_3$C$_2$T$_x$/RGO ternary composite aerogel with a three-dimensional hierarchical architecture was synthesized by a hydrothermal method [160]. The minimum RL of the aerogel reached $-65.3$ dB at the thickness of 2.5 mm. At the same time, the absorption bandwidth was
4.3 GHz, with a thickness of only 2.0 mm. The improved EM wave absorption performance was attributed to the highly porous conductive networks, multiple reflection, and scattering and defective polarization properties.

Wang et al. reported a multilayer Ti3C2Tx@RGO hybrid aerogel prepared by a hydrothermal method and freeze-drying treatment [161]. Compared with pure multilayer Ti3C2Tx, the EM wave absorption of the Ti3C2Tx@RGO aerogel improved significantly. The minimum RL achieved for the Ti3C2Tx@RGO aerogel was −31.2 dB at 8.2 GHz, and the absorption bandwidth reached 5.4 GHz. Such good performance was due to the conductive network, interface polarization, dipole polarization, and multiple scattering, as important contributors.

6 Overview and Perspectives

As shown in Table 1, MXene antennas exhibit excellent comprehensive performance compared with other materials [163–169], as clearly confirmed by Fig. 12a, b. This indicates that MXene antennas have great development prospects for the future. Research on the EM attenuation properties of MXenes was initiated in 2016. Thereafter, a large number of MXene-based structures have been designed for EMI shielding and EM wave absorption based on the principle of “lightweight, wide, and strong.” Tables 2 and 3 list the EMI shielding and EM wave absorption performances of MXene-based materials, respectively. Research on MXene-based shielding and absorption materials has mainly focused on film and hybrid materials (Fig. 12c–f), respectively. MXene-based films afford the highest conductivity among this class of materials (Table 2), which is the reason why MXene films are the most widely used materials for EMI shielding. However, the high conductivity leads to
impedance mismatch. Therefore, no research has been performed on MXene films for EM wave absorption (Table 3). MXene hybrids for EM wave absorption are the most extensively studied materials, owing to their easily tunable EM parameters. Moreover, the addition of a matrix also simplifies adjustment for impedance matching. The choice of foam and aerogel is based on their lightweight characteristics (Fig. 12d), and the porous structure can increase multiple reflections to improve the attenuation of EM waves. The advantage of using a fabric as the EM attenuation material is that it has satisfactory porosity and permeability. In addition, it is more practical.

To summarize, the core outstanding areas to be addressed in the wireless communication and EM attenuation fields include the following: conduction design, because the conduction directly affects the thickness and transmission performance of the antenna, the strength of the first interface reflection is dominated by conduction, and conduction loss plays an important role in EM wave absorption; EM transmission and attenuation mechanisms in MXenes that are not understood and may fundamentally differ from the behavior in other 2D materials, such as graphene; how to balance EM wave absorption and reflection in MXenes to achieve green shielding, and how wireless communication devices employing MXenes can be adapted to large-scale industrial production in the future.

Based on the above overview of MXenes, there are still some challenges that need to be addressed in the future. The conduction is mainly decided by the type of MXene, number and types of surface functional groups, and the construction mode of MXenes. Conduction varies greatly in different kinds of MXenes. For example, the conductivity

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Fig. 12 Comparison of a the return loss and b efficiency of MXene antenna with other materials. c EMI SE and d SEE, versus thickness for MXene-based shielding materials. e RL_{min} and f absorption bandwidth versus thickness of typical MXene-based absorbing materials.
of a free-standing Mo₂Ti₂C₃ film is 100 S cm⁻¹, while the conductivity of a free-standing Ta₄C₃ film is 0.476 S cm⁻¹ [24]. However, current research has mainly focused on Ti₃C₂Tₓ MXene. It is necessary to study the wireless communication performance and EM response mechanism of other MXenes. The number and types of surface functional groups are mainly affected by the etching method. For example, Ti₃C₂Tₓ etched with LiF/HCl has a high content of =O terminal groups compared with Ti₃C₂Tₓ etched with HF, which leads to conductivity differences [54]. The annealing temperature also has a certain influence on the type of surface functional groups [24]. Researchers need to study how to control the conductivity of MXenes by selecting different etchants and annealing temperatures. Moreover, the method used to construct MXenes also affects the conductivity of MXenes. For example, the conductivity of a pressed Ti₃C₂Tₓ disk (2 S cm⁻¹) is lower than that of a free-standing Ti₃C₂Tₓ film (1500 S cm⁻¹). Designing different methods of constructing MXenes to regulate the conduction will also be a direction for future research. The oxidation of MXenes is an unavoidable problem, especially in the case of few-layer or monolayer MXenes. After oxidation, the conductivity of

| Table 2  Typical MXene-based materials and their EMI shielding properties |
|---------------------------------|----------------|----------------|----------------|----------------|----------------|----------------|
| Type                        | Materials       | Matrix | Ratio (wt%)  | d (mm) | σ (S m⁻¹) | SE (dB) | SEE (dB  cm²  g⁻¹) | References |
| Pure MXene                  | Ti₃C₂Tₓ        | Wax    | 60          | 2     | –        | 39.1     | –                  | [52]       |
|                            | Ti₃C₂Tₓ        | SiO₂    | 60          | 1     | 0.42     | 26.7     | –                  | [54]       |
|                            | Ti₃CTₓ        | Wax    | 40          | 0.8   | 1.63 x 10⁻¹⁶ | 6        | –                  | [55]       |
|                            | Ti₃C₂Tₓ        | PS      | 1.9 vol% | 2     | 1081     | 62       | –                  | [56]       |
|                            | Ti₃C₂Tₓ        | PVDF    | 50          | 1     | 0.988    | 34.49    | –                  | [57]       |
| MXene hybrid               | Ti₃C₂Tₓ–Ni    | Wax    | 50          | 2.8   | 4        | 66.4     | –                  | [61]       |
|                            | Ti₃C₂Tₓ–Ag    | Wax    | 60          | 1     | 3.813    | 62.7     | –                  | [62]       |
|                            | Nb₂CTₓ–Ag    | Wax    | 60          | 1     | 3.123    | 72.04    | –                  | [62]       |
|                            | r GO–Ti₃C₂Tₓ  | Epoxy  | 4.5         | 0.5   | 387.1    | 55       | –                  | [63]       |
| MXene film                 | Ti₃C₂Tₓ        | –       | 100         | 0.045 | 4.665 x 10⁵ | 92       | 25.863  | [65]       |
|                            | Ti₃C₂Tₓ        | –       | 100         | 5.5 x 10⁻³ | 5 x 10⁵   | 20       | 3.89 x 10⁶ | [66]       |
|                            | Ti₃C₂Tₓ        | –       | 100         | 0.04   | 1.786 x 10⁵ | 116.2    | –                  | [67]       |
|                            | Ti₃C₂Tₓ        | –       | 100         | 0.011  | 1.6 x 10⁵   | 50       | –                  | [68]       |
|                            | Ti₃C₂Tₓ       | –       | 100         | 0.01   | 500      | 15       | –                  | [68]       |
|                            | Ti₃C₂Tₓ–SA     | –       | 90          | 0.008  | 2.9 x 10⁵   | 57       | 30.830  | [65]       |
|                            | Ti₃C₂Tₓ/ANF    | –       | 80          | 0.017  | 1.733 x 10⁴ | 28       | 1317.64 | [69]       |
|                            | ANF/Ti₃C₂Tₓ/Ag | –       | 20          | 0.045  | 9.22 x 10⁴ | 48.1     | 8907.4  | [73]       |
|                            | Ti₃C₂Tₓ/CNF    | –       | 90          | 0.047  | 739.4    | 24       | 2647    | [74]       |
|                            | Ti₃C₂Tₓ/PEDOT:PSS | –     | 87.5       | 0.0111 | 3.405 x 10⁴ | 42.10    | 19497.8 | [82]       |
|                            | Ti₃C₂Tₓ/PVA    | –       | 19.5       | 0.027  | 716      | 44.4     | 9343    | [87]       |
|                            | Ti₃C₂Tₓ/GO     | –       | 90          | 0.007  | 2.64 x 10⁵ | 50.2     | –       | [92]       |
|                            | Ti₃C₂Tₓ/MMT    | –       | 90          | 0.025  | 4420     | 65       | 10000   | [97]       |
| MXene foam                 | Ti₃C₂Tₓ        | –       | 100         | 0.006  | 58.820   | 32       | 136752  | [98]       |
|                            | Ti₃C₂Tₓ/PVA    | –       | 0.15 vol%  | 5     | 8.3 x 10⁻⁶ | 28       | 5136    | [100]      |
|                            | Ti₃C₂Tₓ/rGO    | –       | 33          | 1.5    | 1000     | 28.6     | 6217    | [101]      |
| MXene aerogel              | Ti₃C₂Tₓ        | –       | 100         | 1     | –        | 70.5     | 64.182  | [104]      |
|                            | Ti₃C₂Tₓ        | –       | 100         | 1     | –        | 69.2     | 62.909  | [104]      |
|                            | Ti₃C₂Tₓ        | –       | 100         | 1     | –        | 54.1     | 49.182  | [104]      |
|                            | Ti₃C₂Tₓ/rGO    | Epoxy  | 0.99 vol%  | 2     | 695.9    | 56.4     | –       | [107]      |
|                            | Ti₃C₂Tₓ/CNT    | –       | 25          | 3     | 943      | 103.99   | 8253.17 | [108]      |
| MXene fabric               | Ti₃C₂Tₓ        | Cotton | 6          | 0.33  | 5 Ω sq⁻¹ | 36       | –       | [111]      |
|                            | Ti₃C₂Tₓ        | Cotton | 5.2 mg/cm² | –     | 670.3    | 31.04    | –       | [112]      |
|                            | Ti₃C₂Tₓ        | Cellulose | 1.89 vol% | 0.2  | 2756     | 42.7     | –       | [113]      |
MXenes decreases significantly. How to inhibit the oxidation of MXenes is a key point for future research.

At present, research on the wireless communication performance of MXenes is still in the exploratory stage. The mechanism of transmission of EM waves in MXene antennas is not very clear. Although research on EM attenuation in MXenes is more extensive, the exact mechanism of EMI shielding and EM wave absorption is still poorly understood. Since the concept of “green shielding” was put forward, realizing green shielding by using MXene-based materials has become a major challenge. Therefore, a series of thorough studies on EM wave transmission, dielectric relaxation, and the EM response of MXene-based materials is urgently needed.

The low yield of MXenes limits their commercial application in wireless communication, EMI shielding, and EM wave absorption materials. Increasing the production of MXenes is the only way to realize commercialization. Moreover, the hydrophilic nature of MXenes limits the fabrication of composites or hybrids with polymers and other materials to only aqueous media. To broaden the path of structural design, it is essential to investigate organic dispersions of MXenes.

In summary, MXenes, as the newest and fastest growing family of 2D materials, will open new avenues for

| Table 3 Typical MXene-based materials and their EM wave absorption properties |
|-----------------------------|------------------|--------|--------|--------------|------------------|--------|
| Type                      | Materials        | Matrix | Ratio (wt%) | d (mm) | RL_{min} (dB) | Bandwidth (< − 10 dB) (GHz) | References |
| Pure MXene                 | Ti₃C₂Tₓ          | Wax    | 40          | 2.5    | −47.9          | 3.6                | [44]      |
|                            | Ti₃C₂Tₓ          | Wax    | 50          | 2      | −40            | 6.8                | [120]     |
|                            | Ti₃C₂Tₓ          | Wax    | 55          | 1.7    | −42.5          | 4.2                | [122]     |
|                            | Nb₂CTₓ          | Wax    | 70          | 2.9    | −52.2          | 0.65               | [126]     |
|                            | Ti₃C₂Tₓ          | Wax    | 45          | 1.7    | −36            | 5.6                | [127]     |
|                            | TiₓCTₓ          | Wax    | 55          | 4.5    | −54.1          | 1.1                | [128]     |
| MXene hybrid               | TiO₂/Ti₃C₂Tₓ/FcO₂ | Wax    | 70          | 1.9    | −57.3          | 2                  | [130]     |
|                            | Fe₃O₄/Ti₃C₂Tₓ    | Wax    | 60          | 4.2    | −57.2          | 1.4                | [132]     |
|                            | Ti₃C₂Tₓ/FCl      | Epoxy  | 60          | 1      | −15.52         | 8.16               | [134]     |
|                            | Ti₃C₂Tₓ/Ni       | Wax    | 50          | 1.5    | −47.06         | 3.6                | [135]     |
|                            | Ni@Ti₃C₂Tₓ      | PVDF   | 10          | 3      | −52.6          | 3.7                | [137]     |
|                            | TiₓCₓTₓ/Ni       | Wax    | 60          | 2.2    | −24.3          | 2.6                | [138]     |
|                            | TiₓCₓTₓ/CoOₓ₄   | Wax    | 50          | 2      | −34.8          | 6.2                | [139]     |
|                            | CoFe@Ti₃C₂Tₓ    | Wax    | 60          | 2.2    | −36.29         | 2.64               | [140]     |
|                            | FeCo@Ti₃C₂Tₓ    | Wax    | 70          | 1.6    | −17.86         | 8.8                | [141]     |
|                            | Ti₃C₂Tₓ−NiCoO₄ | Wax    | 50          | 2.18   | −50.96         | 0.8                | [143]     |
|                            | Co₂Z/Ti₃C₂Tₓ    | PVB    | 30          | 2.8    | −46.3          | 1.6                | [144]     |
|                            | G/TiC/TiₓCₓTₓ   | Wax    | 50          | 2.1    | −63            | 3.5                | [145]     |
|                            | CNT/TiₓCₓTₓ     | Wax    | 35          | 1.55   | −52.9          | 4.46               | [146]     |
|                            | N-GP/TiₓCₓTₓ    | Epoxy/PA | 32     | 1.4    | −52            | 7.1                | [148]     |
|                            | TiₓCₓTₓ@PPy     | Wax    | 10          | 3.6    | −49.5          | 5.6                | [150]     |
|                            | MoS₂@TiO₂/TiₓCₓTₓ| Wax    | 50          | 2.5    | −15.9          | 2.6                | [152]     |
| MXene foam                 | TiₓCₓTₓ/SiC     | –      | 100         | 2.6    | −14.7          | 3.2                | [156]     |
|                            | CF/TiₓCₓTₓ     | Wax    | –           | 4.5    | −45            | 5                  | [157]     |
| MXene aerogel              | TiₓCₓTₓ@gelatin | –      | 100         | 2      | −59.5          | 6.24               | [158]     |
|                            | TiₓCₓTₓ/Cellulose| Wax    | 24          | 2      | −43.4          | 4.5                | [159]     |
|                            | TiO₂/TiₓCₓTₓ/RGO| Wax    | 10          | 2.5    | −65.3          | 4.3                | [160]     |
|                            | TiₓCₓTₓ@RGO    | Wax    | 15          | 2.05   | −31.2          | 5.4                | [161]     |
| MXene fabric               | TiₓCₓTₓ/Ni/ZnO  | Cotton | 13.39     | 2.8    | −35.1          | 4.2                | [162]     |
realizing various classes of wireless communication and EM protection devices. This review is expected to serve as a guide to those exploring wireless communication and EM attenuation properties of MXenes.

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