

## PAPER

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# Novel MOF-derived 3D hierarchical needlelike array architecture with excellent EMI shielding, thermal insulation and supercapacitor performance†

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The upcoming 5G era will powerfully promote the development of intelligent society in the future, but it will also bring serious electromagnetic pollution. Thus, the development of efficient, lightweight and multifunctional electromagnetic shielding materials and devices is an important research hotspot around the world. Herein, a novel needlelike  $\text{Co}_3\text{O}_4/\text{C}$  array architecture is constructed from MOF precursor *via* a simple pyrolysis process, and its microstructure is controllably tailored by changing the pyrolysis temperature. The unique 3D hierarchical structure and multiphase components enable the architecture to provide high-efficiency electromagnetic interference (EMI) shielding, along with good thermal insulation. More importantly, the architecture possesses fast ion transport channels, which can be used to construct supercapacitors with high specific capacitance and excellent cycle stability. Obviously, this work offers a new inspiration for the design and construction of multifunctional electromagnetic materials and devices.

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## 1. Introduction

With the advent of the 5G era, electromagnetic pollution emitted by increasingly popular electronic devices has posed a great threat to the global society, which not only seriously threatens the national military security, but also creates potential hazards to human health.<sup>1–8</sup> Thus, the development of high-efficiency electromagnetic shielding materials and devices has become a research hotspot around the world. Meanwhile, the rapid development of electronic devices towards miniaturization and intelligence has also put forward new requirements for thermal insulation and energy storage performance. Currently, although considerable efforts have been devoted to searching for new materials capable of achieving electromagnetic interference (EMI) shielding, thermal insulation and energy storage, few works have been reported on the integration of these three functions together.<sup>9–12</sup> Therefore, the exploration of high-efficiency and multifunctional EMI shielding materials is an important research topic.<sup>13,14</sup>

Metal–organic framework (MOF) is gathering momentum as versatile materials in many fields, such as electromagnetic

protection, battery, catalysis, and supercapacitor.<sup>15–21</sup> In particular, benefiting from the designable components, variable structures and tunable electromagnetic properties, it is highly favored in electromagnetic materials and functional devices.<sup>22,23</sup> For example, Ji' group<sup>24</sup> reported a highly efficient NC- $\text{Co}_3\text{O}_4/\text{CP}$  absorber with a minimum reflection loss (RL) of  $-41.38$  dB at 2.3 mm. Che and co-workers<sup>25</sup> fabricated  $\text{Co}@\text{NC}$  composites that displayed a minimum RL of  $-53.0$  dB at 1.8 mm. Chen *et al.*<sup>26</sup> produced a core-shell  $\text{ZnO}@\text{C}/\text{Co}_3\text{ZnC}$  and its RL reached  $-62.9$  dB with a bandwidth of 5.5 GHz. Xie' group<sup>27</sup> developed MOFs/SiC nanowire hybrids, and its RL achieved  $-47$  dB. Han *et al.*<sup>28</sup> synthesized hollow carbon microcubes, with a minimum RL of  $-60.7$  dB. Zhang' group<sup>29</sup> prepared ferrite/Co/porous carbon materials, and the optimal RL was  $-31.05$  dB at 1.5 mm. These fantastic MOF-derived functional materials have brought continued prosperity to the field of electromagnetic wave absorption. However, the development of MOFs has barely begun for EMI shielding, let alone the multifunctional shielding materials.

In this work, a novel and hierarchical needlelike  $\text{Co}_3\text{O}_4/\text{C}$  array architecture is successfully constructed from zeolitic imidazolate framework-67 (ZIF-67) precursor on Ni foam ( $\text{NF}@\text{Co}/\text{C}$ ), and the effect of pyrolysis temperature on the microstructure is dissected in depth. Benefitting from the unique 3D hierarchical structure and multiphase components, this  $\text{NF}@\text{Co}/\text{C}$  architecture integrates three functions, including excellent EMI shielding and good thermal insulation as well as stable and efficient supercapacitor performance. Importantly,

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a deep insight into the connection between structure and functions is given, and the internal response mechanism is reasonably excavated. Undoubtedly, this multifunctional architecture is of great significance to the construction of portable and intelligent devices.

## 2. Results and discussion

The preparation route of needlelike  $\text{Co}_3\text{O}_4/\text{C}$  arrays on nickel (Ni) foam is given in Fig. 1. Initially, the MOF-based precursors (NF@ZIF-67) are obtained by a simple self-assembly process. Afterwards, the obtained precursors are pyrolyzed in the air to produce 3D hierarchical needlelike  $\text{Co}_3\text{O}_4/\text{C}$  arrays. The microstructure of  $\text{Co}_3\text{O}_4/\text{C}$  arrays can be flexibly adjusted by chan-

ging the pyrolysis temperature. Fig. S1a† shows the XRD pattern of ZIF-67 stripped from the NF@ZIF-67, which is in good agreement with the simulated XRD pattern.<sup>30</sup> In Fig. S1b,† eight typical diffraction peaks can be indexed to the cubic structure of  $\text{Co}_3\text{O}_4$ , confirming the successful preparation of NF@ $\text{Co}_3\text{O}_4/\text{C}$  architecture from NF@ZIF-67 precursor by the pyrolysis process.

SEM images clearly show the grown ZIF-67 and  $\text{Co}_3\text{O}_4/\text{C}$  arrays on the surface of Ni foam, as shown in Fig. 2. The obtained products pyrolyzed at different temperatures possess various morphologies. The NF@ $\text{Co}_3\text{O}_4/\text{C}$  architectures pyrolyzed at relatively low temperatures inherit the polyhedral shape of ZIF-67, but their surfaces become rough. As the pyrolysis temperature increases, the fracture and decomposition of 2-methylimidazole (2-MIM) ligand aggravate, clearly enlarging the

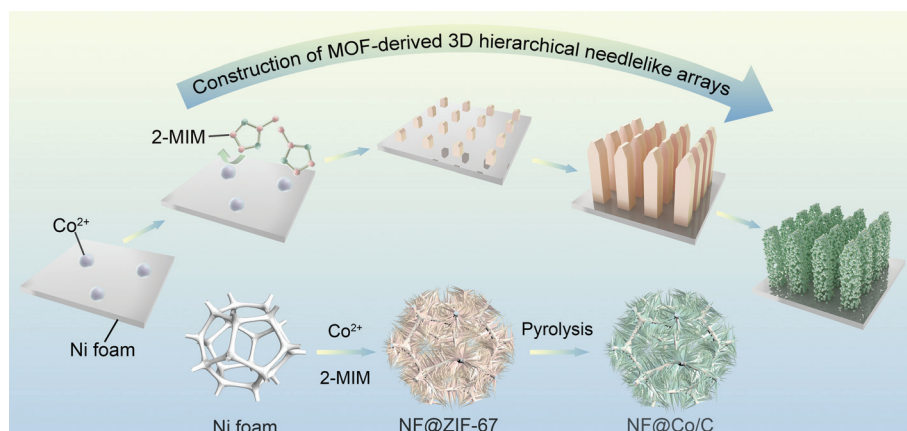


Fig. 1 Schematic illustration of the preparation route of NF@ $\text{Co}_3\text{O}_4/\text{C}$  architecture.



Fig. 2 SEM images of (a) NF@ZIF-67, (b) NF@ $\text{Co}_3\text{O}_4/\text{C}$ -450, (c) NF@ $\text{Co}_3\text{O}_4/\text{C}$ -500, (d) NF@ $\text{Co}_3\text{O}_4/\text{C}$ -550, (e) NF@ $\text{Co}_3\text{O}_4/\text{C}$ -600 and (f) NF@ $\text{Co}_3\text{O}_4/\text{C}$ -650. (g) SEM and elemental mapping images of NF@ $\text{Co}_3\text{O}_4/\text{C}$ -550.



pores on the surface of the product.<sup>31–33</sup> Meanwhile, the structure further shrinks, breaks, and tends to form small nanoparticles. As shown in Fig. 2b–f, the resulting  $\text{Co}_3\text{O}_4/\text{C}$  arrays consist of numerous  $\text{Co}_3\text{O}_4/\text{C}$  nanoparticles. In addition, high temperature will destroy the connection between  $\text{Co}_3\text{O}_4/\text{C}$  arrays and Ni foam, causing them to flake off. Fig. 2g shows the distribution of Co, O, C and N elements in the selected area. A uniform distribution of Co and O elements confirms the uniform growth of  $\text{Co}_3\text{O}_4$  arrays on the surface of Ni foam.<sup>34</sup> Meanwhile, nitrogen atoms are demonstrated to be successfully doped, and a small amount of carbon is retained in the final products.<sup>35,36</sup>

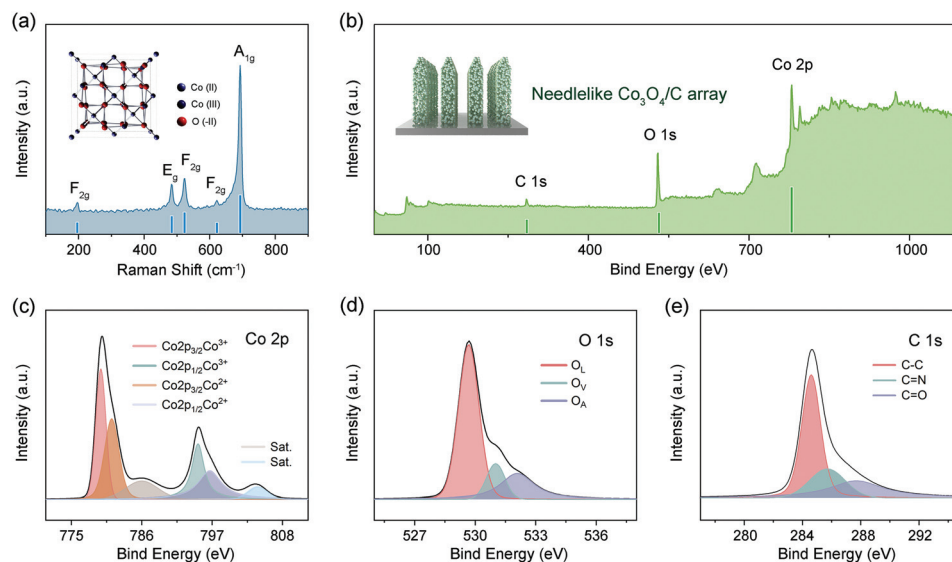
An insight into the morphologies and microstructures of NF@ZIF-67 and NF@Co/C architectures is further given by TEM, HR-TEM and SAED images (Fig. 3). As displayed in Fig. 3a and b, ZIF-67 shows a needlelike morphology with a diameter of  $\approx 1.2 \mu\text{m}$ . The TEM images in Fig. 3c–f show the morphology of the  $\text{Co}_3\text{O}_4/\text{C}$ -550 sample, once again confirming that the needlelike  $\text{Co}_3\text{O}_4/\text{C}$  arrays consist of small nanoparticles. A thin layer of carbon is observed at the edge of the  $\text{Co}_3\text{O}_4$  nanoparticle, which is consistent with elemental mapping (Fig. 3g).<sup>37,38</sup> Furthermore, two typical lattice fringes of 0.23 and 0.45 nm are observed in Fig. 3h, corresponding to (311) and (111) crystal planes, respectively.<sup>37,38</sup> The size of nanoparticles ranges from 25 nm to 85 nm (Fig. 3i). Fig. 3j illustrates the conversion process from ZIF-67 to  $\text{Co}_3\text{O}_4/\text{C}$ . In this process, the ZIF-67 generates cracks and gradually disintegrates, where Co ion and 2-MIM ligand are converted into  $\text{Co}_3\text{O}_4$  nanoparticles and carbon layers, respectively, under the action of high temperature and oxygen.<sup>32</sup>

The microstructure of NF@Co/C architecture is investigated by Raman spectrum, as shown in Fig. 4a. The peaks appearing around 194, 482, 522, 618, and  $691 \text{ cm}^{-1}$  are attributed to  $\text{Co}_3\text{O}_4$ .<sup>37–39</sup> The XPS spectra further confirm the successful preparation of  $\text{Co}_3\text{O}_4/\text{C}$  (Fig. 4b). The Co 2p spectrum shows two prominent peaks, including 779.8 eV for Co 2p<sub>3/2</sub> and 794.9 eV for Co 2p<sub>1/2</sub>, respectively (Fig. 4c).<sup>31,40</sup> Two weak peaks at 786.2 and 804.1 eV may correspond to the shakeup satellite peaks. The energy difference between Co 2p<sub>3/2</sub> and Co 2p<sub>1/2</sub> is about 15 eV, revealing that the Co element exists mainly in the form of  $\text{Co}_3\text{O}_4$ .<sup>31,38</sup> The peaks of Co 2p<sub>3/2</sub> and Co 2p<sub>1/2</sub> can be deconvoluted into four fitting peaks. The peaks at 781.2 and 796.7 eV are assigned to  $\text{Co}^{2+}$ , while the other two peaks at 779.7 and 794.8 eV are attributed to  $\text{Co}^{3+}$ .<sup>34–37</sup> The O 1s spectrum can be resolved into three fitting curves with binding energies at 529.7, 531.1 and 532.1 eV (Fig. 4d). They are assigned to the metal–oxygen bonds (Co–O bonds,  $\text{O}_\text{L}$ ), oxygen vacancies ( $\text{O}_\text{V}$ ) and adsorbed oxygen species ( $\text{O}_\text{A}$ ), respectively.<sup>41</sup> In addition, the C 1s spectrum can be deconvoluted into three major peaks (Fig. 4e). The peaks at 284.6, 285.7 and 287.8 eV are assigned to C–C bonds, C=N bonds and C=O bonds, respectively.<sup>31,40</sup>

The total shielding effectiveness ( $\text{SE}_\text{T}$ ), as well as the contribution of absorption ( $\text{SE}_\text{A}$ ) and reflection ( $\text{SE}_\text{R}$ ), of NF@Co/C architectures is investigated in the frequency range from 8.2 to 12.4 GHz (Fig. S2a–c†). The obtained products demonstrate the fluctuant  $\text{SE}_\text{T}$  in X band, which may result from the uneven distribution of internal material. The NF@Co/C-550 sample has the optimal  $\text{SE}_\text{T}$  in the investigated frequency band due to the appropriate structure and component. Among them, Ni



**Fig. 3** TEM and HR-TEM images of (a), (b) ZIF-67, (c–h)  $\text{Co}_3\text{O}_4/\text{C}$ . Inset in (g) is the corresponding SAED pattern. (i) Size distribution of  $\text{Co}_3\text{O}_4/\text{C}$  nanoparticles. (j) Schematic illustration for the pyrolysis from ZIF-67 to  $\text{Co}_3\text{O}_4/\text{C}$ .

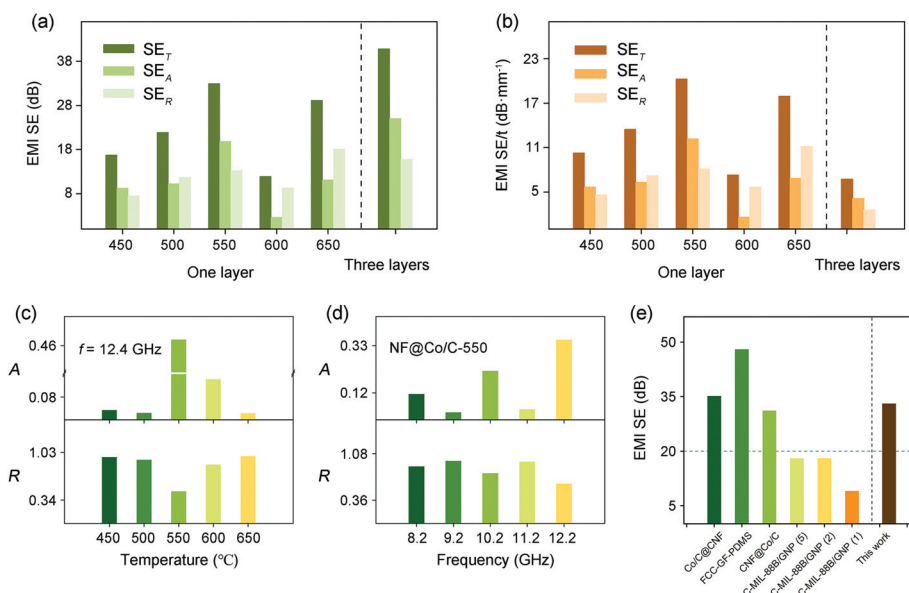


**Fig. 4** (a) Raman spectrum of NF@Co/C, and inset is the crystal structure of  $\text{Co}_3\text{O}_4$ . (b) Broad scan XPS spectrum of NF@Co/C, and inset is the corresponding model. (c) Co 2p spectrum. (d) O 1s spectrum. (e) C 1s spectrum.

foam with high electrical conductivity significantly contributes to the reflection of electromagnetic waves, promoting the improvement of shielding performance (Fig. S3†). The absorption coefficient ( $A$ ), reflection coefficient ( $R$ ) and transmission coefficient ( $T$ ) represent the capability of a shielding material to absorb, reflect and transmit electromagnetic waves, respectively. As shown in Fig. S2d–f,† both  $R$  and  $A$  coefficients vary to different degrees, and the  $A$  coefficient closes to 0.5 at most, indicating the possibility of green EMI shielding. Meanwhile,

the  $T$  coefficient is less than 0.1, suggesting the high-efficiency EMI shielding of the architectures.

The  $\text{SE}_T$ ,  $\text{SE}_A$  and  $\text{SE}_R$  values of the NF@Co/C architectures are evaluated in Fig. 5a. It can be observed that the pyrolysis temperature has a significant effect on the EMI shielding of the NF@Co/C architecture. The NF@Co/C-550 sample achieves the optimal shielding performance with a maximum EMI SE of 33 dB, suggesting that it can shield 99.95% of electromagnetic waves. The proportion of  $\text{SE}_A$  reaches over 60%. As a



**Fig. 5** (a) Comparison of  $\text{SE}_T$ ,  $\text{SE}_A$  and  $\text{SE}_R$  values of the NF@Co/C architectures with different pyrolysis temperatures. (b) Comparison of specific  $\text{SE}_T$ ,  $\text{SE}_A$  and  $\text{SE}_R$  values. The thickness of one-layer samples is 1.631, 1.625, 1.626, 1.634 and 1.623 mm, respectively. The thickness of the three-layer sample is 4.882 mm. Comparison of  $A$  and  $R$  coefficients at (c) different temperatures and (d) frequencies. (e) Comparison of the EMI SE performance of the MOF-derived materials ever reported with that of this work.

result, the pyrolysis temperature should be selected around 550 °C for further application. In addition, the EMI SE of both NF@Co/C-500 and NF@Co/C-650 samples also exceeds 20 dB, enabling them to meet commercial application standards. The EMI SE of three-layer sample achieves 41 dB. More importantly, the NF@Co/C architectures can be directly applied to various electronic devices, which has a greater advantage than powder MOF materials in practical applications. The specific shielding effectiveness (EMI SE/t) can exclude the influence of thickness on the performance. As plotted in Fig. 5b, the NF@Co/C-550 sample also exhibits the highest EMI SE/t.

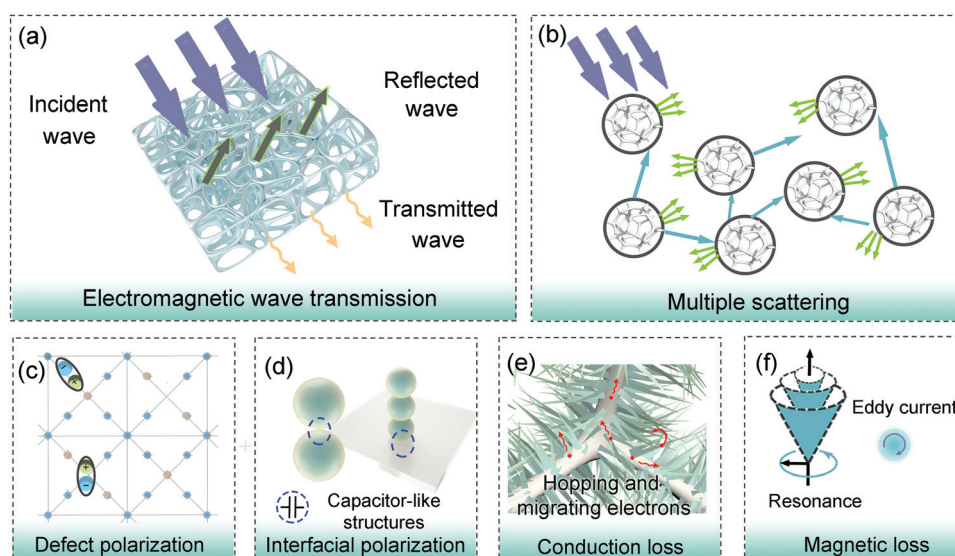
Absorption and reflection are the two main mechanisms affecting EMI shielding. The influence of pyrolysis temperature on the *A* and *R* coefficients is investigated in Fig. 5c. At 12.4 GHz, the NF@Co/C-550 sample shows the largest *A* coefficient, higher than *R* coefficient, which means that its green shielding index ( $g_s$ ) is greater than 1.<sup>41,42</sup> Fig. 5d shows the *A* and *R* coefficients of the NF@Co/C-550 sample at different frequencies. The high electrical conductivity ( $1.2588 \times 10^5 \text{ S m}^{-1}$ ) is conducive to the reflection of electromagnetic waves, resulting in the larger *R* coefficient. The  $g_s$  is adopted to evaluate the environmental friendliness of shielding materials. The result suggests the possibility to achieve green shielding by optimizing the pyrolysis temperature under certain conditions. Compared with other MOF-based shielding materials previously reported, the NF@Co/C-550 sample shows excellent electromagnetic shielding performance and competitiveness, opening new possibilities for the design of green EMI shielding materials (Fig. 5e).<sup>43–46</sup>

Fig. 6a shows the electromagnetic shielding mechanism of the NF@Co/C architecture. The porous Ni foam and needlelike  $\text{Co}_3\text{O}_4/\text{C}$  arrays provide a large surface area, which promotes the scattering of electromagnetic waves and enhances energy dissipation (Fig. 6b).<sup>47,48</sup> Therefore, the constructed 3D hierarchical needlelike architecture can significantly improve the

EMI shielding performance. In addition, by tailoring the microstructure and component, the effective absorption of electromagnetic waves by the NF@Co/C architecture can also be reasonably optimized.

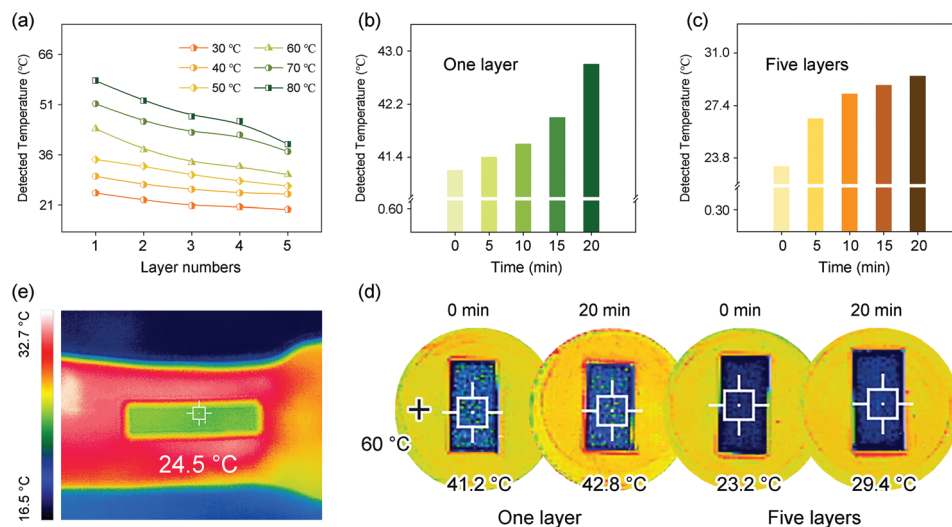
To further study the electromagnetic response mechanism, the NF@Co/C-550 sample was mixed with wax and its electromagnetic parameters were measured (Fig. S4†). First, the defects inside the needlelike  $\text{Co}_3\text{O}_4/\text{C}$  arrays cause the charge asymmetric distribution, forming dipoles (Fig. 6c).<sup>38,49–54</sup> Meanwhile, the interfaces between  $\text{Co}_3\text{O}_4$  core and carbon shell, between  $\text{Co}_3\text{O}_4/\text{C}$  nanoparticles, and between  $\text{Co}_3\text{O}_4/\text{C}$  nanoparticles and Ni foam can be equivalent to capacitor-like structures and generate interfacial dipoles (Fig. 6d). Under an alternating electromagnetic field, these dipoles convert electromagnetic energy into thermal energy due to relaxation loss (Fig. S4a and b†).<sup>49,50</sup> Second, electrons are capable of absorbing electromagnetic energy to migrate or hop in the architectures, and cause the energy conversion by colliding and scattering with lattice, *i.e.* conduction loss (Fig. 6e).<sup>1,54</sup> Third, magnetic  $\text{Co}_3\text{O}_4$  and Ni foam can bring additional magnetic loss (Fig. 6f). As shown in Fig. S4c,† the eddy current coefficient ( $\mu''(\mu')^{-2}f^{-1}$ ) varies with frequency, suggesting that the eddy current loss isn't the only source of magnetic loss within the architectures. A peak appears in the complex permeability (Fig. S4d†), which may be indexed to magnetic resonance.<sup>38,39,55,56</sup>

The NF@Co/C architecture also has good thermal insulation performance, which can protect electronic devices from high temperature. The temperature variation of the upper surface of the NF@Co/C architecture with the number of layers is compared at different heating temperatures. The heating temperature is set to 30, 40, 50, 60, 70 and 80 °C, respectively, and the number of layers is increased from one to five. As shown in Fig. 7a, the increment of the number of layers effectively improves the thermal insulation performance. Fig. 7b



**Fig. 6** EMI shielding mechanism of NF@Co/C architecture. (a) Electromagnetic wave transmission. (b) Multiple scattering. (c) Defect polarization. (d) Interfacial polarization. (e) Conduction loss. (f) Magnetic loss.





**Fig. 7** (a) Variation of the temperature detected on the upper surface of the NF@Co/C architecture with the number of layers at different heating temperatures. Variation of the temperature detected of (b) one-layer and (c) five-layer samples with heating time, as well as (d) the corresponding thermal infrared images. (e) Thermal infrared image of the NF@Co/C architecture placed on the arm.

and c present the detected temperatures of one-layer and five-layer samples that are recorded in the range of 0 to 20 min on a heated platform of 60 °C. The temperature difference maintains a narrow range of variation, reflecting the stable thermal insulation performance of the architecture. The thermal infrared images of one-layer and five-layer samples at 0 and 20 min are captured, respectively, by infrared detection device, as shown in Fig. 7d. Compared with the chartreuse presented by heating platforms, the covered area by the sample turns obvious dark blue, intuitively confirming the excellent thermal insulation performance of the architecture.

Benefiting from the good thermal insulation performance, the NF@Co/C architecture is endowed with infrared stealth function that is worth expecting. Fig. 7e emerges the thermal infrared image of the NF@Co/C architecture placed on the arm. The apparent red of the arm suggests its strong infrared radiation. After the NF@Co/C architecture is placed, the color of the covered area turns from red to green in the thermal infrared image, indicating its excellent radiation suppression. In general, thermal conductivity depends on solid-phase conduction, gas-phase conduction and radiation heat transfer.<sup>18</sup> The NF@Co/C architecture features low density and high porosity, and has air with low thermal conductivity insides simultaneously, which is considered to have a great advantage in significantly reducing both the thermal conduction and the heat transfer by radiation. The thermal conductivity of the NF@Co/C-550 sample measured at 60 °C is low to 0.5108 W m<sup>-1</sup> K<sup>-1</sup>, confirming the above analysis.

The capacitance performances of the NF@Co/C architecture are evaluated in a 6 M KOH aqueous electrolyte through a three-electrode system, with Pt foil as counter electrode and Hg/HgO as reference electrode (the inset in Fig. 8a). The NF@Co/C architecture can be directly used as an electrode without additional conductive additives and binders, thereby effectively reducing

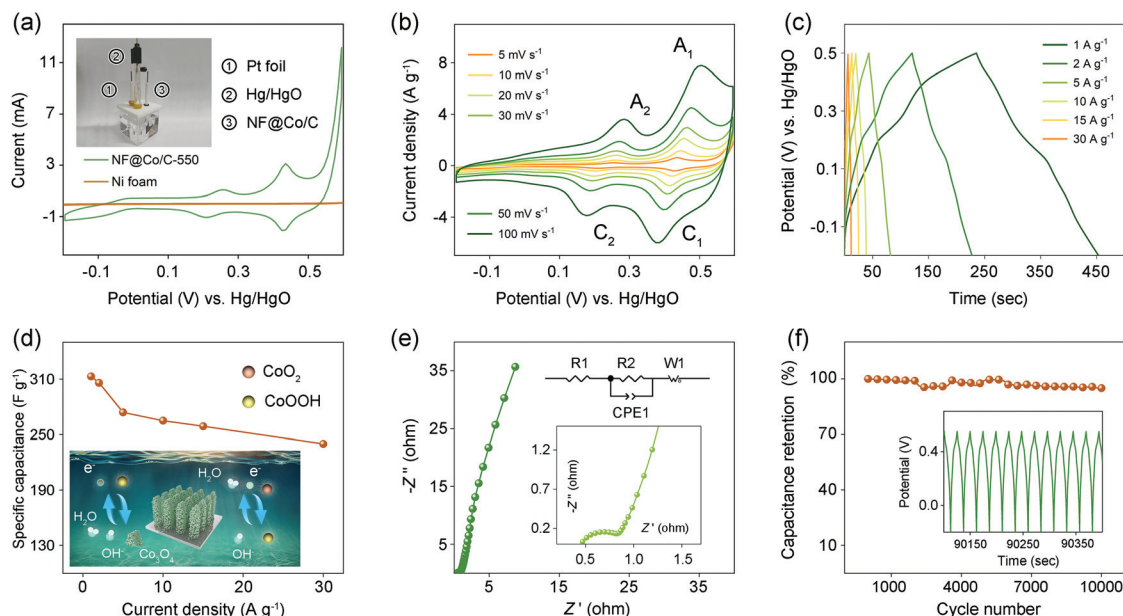
the internal resistance of the electrode and increasing the overall specific capacitance.<sup>57</sup> Fig. S5† compares the cyclic voltammetry (CV) curves of five samples at a scan rate of 5 mV s<sup>-1</sup>. The NF@Co/C-550 sample exhibits the largest CV curve area, the strongest redox peak, and the highest specific capacitance, which is presumably given credit to its appropriate structure and component. Fig. 8a compares the CV curves of the NF@Co/C-550 sample and Ni foam at 5 mV s<sup>-1</sup>, revealing the little contribution of Ni foam to specific capacitance.

In Fig. 8b, the CV curves of the NF@Co/C-550 sample exhibit two pairs of redox peaks. The kinetics analysis of the NF@Co/C-550 sample is conducted with the data of CV curves. As shown in Fig. S6,† the contribution of surface capacitance increases with the scanning rate. The electrochemical active surface area (ECSA) is calculated according to equation:

$$ECSA = \frac{C_{dl}}{C^*} \quad (1)$$

where the general specific capacitance  $C^*$  in alkaline electrolyte is 0.04 mF cm<sup>-2</sup>.<sup>58</sup> The calculated  $C_{dl}$  values of NF@Co/C-550 sample from its CV curve is 15.79 mF cm<sup>-2</sup>, and the calculated ECSA reaches 394 cm<sup>-2</sup> (Fig. S7†). The large specific surface area and the unique porous structure can reduce charge transfer resistance and accelerate ion diffusion/transportation kinetics.<sup>59</sup> One pair of redox peaks ( $A_1/C_1$ ) corresponds to the conversion between CoOOH and CoO<sub>2</sub>, and the other pair of peaks ( $A_2/C_2$ ) is indexed to the reversible reaction between Co<sub>3</sub>O<sub>4</sub> and CoOOH.<sup>31,60</sup> This result is consistent with previously reported works, but the position and strength of the peaks are slightly different, which may be related to the morphology and crystallinity of the electrode materials.

The galvanostatic charge-discharge (GCD) test is conducted in the potential range from -0.2 to 0.5 V at different current densities to evaluate the electrochemical performance



**Fig. 8** (a) CV curves of Ni foam and NF@Co/C-550 sample at a scan rate of  $5 \text{ mV s}^{-1}$ . Inset is a three-electrode system. (b) CV curves of NF@Co/C-550 sample at different scan rates. (c) GCD curves at different current densities. (d) Specific capacitance. Inset is the schematic illustration for electrode reaction. (e) Nyquist plots and the corresponding equivalent circuit. (f) Cycling performance at a charge–discharge current density of  $10 \text{ A g}^{-1}$ .

(Fig. 8c). The obvious deviation of these charge–discharge curves from the ideal triangular shape reveals the pseudo-capacitive nature of the NF@Co/C-550 electrode. The charge–discharge platforms are in good agreement with the redox couples in the CV curves.<sup>57,61</sup> Based on the GCD curves, the capacitances at different current densities are readily determined, as shown in Fig. 8d.<sup>62</sup> The NF@Co/C-550 sample shows the highest specific capacitance, reaching  $313 \text{ F g}^{-1}$  at  $1 \text{ A g}^{-1}$ . With the increment of current density, the specific capacitance gradually drops, obtaining  $306 \text{ F g}^{-1}$  at  $2 \text{ A g}^{-1}$ ,  $274 \text{ F g}^{-1}$  at  $5 \text{ A g}^{-1}$ ,  $265 \text{ F g}^{-1}$  at  $10 \text{ A g}^{-1}$ ,  $259 \text{ F g}^{-1}$  at  $15 \text{ A g}^{-1}$  and  $240 \text{ F g}^{-1}$  at  $30 \text{ A g}^{-1}$ , respectively. Nonetheless, 76% of the capacitance can still be retained at a current density as high as  $30 \text{ A g}^{-1}$ , indicating the excellent rate capability of the architecture.

Electrochemical impedance spectroscopy (EIS) is employed to characterize the NF@Co/C-550 electrode (Fig. 8e). The equivalent series resistance ( $R_s$ ) is determined from the  $x$ -intercept in the high-frequency region, and the charge transfer resistance ( $R_{ct}$ ) is calculated from the diameter of semicircle. The values of capacitive and resistive components fitting the experimental impedance spectrum based upon the proposed equivalent circuit are given in Table S1.† At a low frequency range, a vertical spike parallel to the  $Z''$  axis indicates typical capacitance behavior.<sup>63</sup> More importantly, the NF@Co/C-550 sample demonstrates extraordinary stability in an alkali solution environment, retaining 95% of the initial capacitance at  $10 \text{ A g}^{-1}$  after 10 000 cycles (Fig. 8f). After cycling tests, the  $R_s$ ,  $R_{ct}$  and Warburg impedance slightly increase (Fig. S8†). Table 1 shows the electrochemical performance of some MOF-

**Table 1** The evaluation of the electrochemical performance of  $\text{Co}_3\text{O}_4$ -based electrodes

| Materials                           | Capacitance ( $\text{F g}^{-1}$ ) | Current density ( $\text{A g}^{-1}$ ) | Cycle number | Cycle stability (%) | Ref.      |
|-------------------------------------|-----------------------------------|---------------------------------------|--------------|---------------------|-----------|
| $\text{Co}_3\text{O}_4$ nanomeshes  | 1216.4                            | 1                                     | 8000         | 86.4%               | 34        |
| $\text{Co}_3\text{O}_4$ @Co-MOF     | 1020                              | 0.5                                   | —            | —                   | 60        |
| $\text{Co}_3\text{O}_4$             | 410                               | 2                                     | —            | —                   | 64        |
| 3DPC/ $\text{Co}_3\text{O}_4$       | 423                               | 1                                     | 2000         | 83%                 | 65        |
| Reduced $\text{Co}_3\text{O}_4$     | 329                               | 1                                     | —            | —                   | 66        |
| $\text{Co}_3\text{O}_4$ microsphere | 261.1                             | 0.5                                   | 2000         | 90.2%               | 67        |
| Nanoporous $\text{Co}_3\text{O}_4$  | 970                               | 1                                     | 5000         | 77.5%               | 68        |
| $\text{Co}_3\text{O}_4$ particles   | 150                               | 1                                     | 3400         | —                   | 69        |
| $\text{Co}_3\text{O}_4$ @Co-MOF     | 797                               | 1                                     | 10 000       | 69%                 | 70        |
| $\text{Co}_3\text{O}_4$ @NPC        | 525                               | 1.5                                   | 8000         | 81%                 | 71        |
| $\text{Co}_3\text{O}_4$ nanotubes   | 647                               | 1                                     | 1500         | —                   | 72        |
| NF@Co/C                             | 313                               | 1                                     | 10 000       | 95%                 | This work |

derived  $\text{Co}_3\text{O}_4$ -based electrodes, where specific capacitance and cycle stability are considered.

The excellent electrochemical performance of NF@Co/C architecture can be attributed to its unique 3D hierarchical structure and multiphase components. First, the uniform distribution of needlelike  $\text{Co}_3\text{O}_4/\text{C}$  arrays on Ni foam ensures good electrochemical activity. Second, the hierarchical structure of the electrode provides a high specific surface area capable of achieving sufficiently contact with the electrolyte and improves ion transport. Third, the direct contact between  $\text{Co}_3\text{O}_4/\text{C}$  active electrode materials and Ni foam avoids the use of conductive additives and binders, reducing the internal resistance of the electrode and increasing the overall specific capacitance. Finally, the unique 3D hierarchical structure of the NF@Co/C architecture helps to alleviate the volume expansion during the cycle process, which particularly contributes to better maintaining structural integrity and providing outstanding cycle stability.

### 3. Conclusions

In summary, a novel and hierarchical needlelike  $\text{Co}_3\text{O}_4/\text{C}$  array architecture derived from a MOF-based precursor has been successfully constructed, and its microstructure is controllably tailored by controlling the pyrolysis temperature. Benefiting from the 3D hierarchical structure and multiphase components, the NF@Co/C architecture achieves highly efficient EMI SE, capable of shielding up to 99.95% of electromagnetic radiation, along with good thermal insulation. More importantly, this architecture can be directly used as an excellent electrode for supercapacitors without additional conductive additives and binders, and 95% of the initial capacitance can be retained after 10 000 cycles. In the future, the novel NF@Co/C architecture will show a bright application prospect as the core of advanced multifunctional devices.

## 4. Experimental section

### 4.1 Materials

Ni foam and cobalt nitrate hexahydrate ( $\text{Co}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$ ) were purchased from Aladdin Reagent Co., LTD (Beijing, China). 2-MIM was purchased from Innochem Technology Co., LTD (Shanghai, China). All chemicals were of analytical grade and used without further purification.

### 4.2 Synthesis of ZIF-67 arrays on Ni foam

Ni foam cut into sheets of 15 mm  $\times$  30 mm was ultrasonically cleaned with deionized water and acetone for 30 minutes, respectively, and then dried at 60 °C for 12 h. In a typical process, 40 mL of  $\text{Co}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$  solution (0.4 M) was quickly added into 40 mL of 2-MIM solution ( $25 \times 10^{-3}$  M). Then, a piece of Ni foam was immersed in the mixture solution for 4 h at room temperature. After being taken out, it was washed with

deionized water for three times, and then dried at 60 °C overnight. The obtained product was denoted as NF@ZIF-67.

### 4.3 Synthesis of $\text{Co}_3\text{O}_4/\text{C}$ arrays on Ni foam

The obtained NF@ZIF-67 precursors were pyrolyzed at different temperatures (450, 500, 550, 600 and 650 °C) for 30 min under air atmosphere with a heating rate of 1 °C  $\text{min}^{-1}$ . The resulting products were correspondingly denoted as NF@Co/C-450, NF@Co/C-500, NF@Co/C-550, NF@Co/C-600 and NF@Co/C-650, respectively.

### 4.4 Characterizations

The X-ray diffraction (XRD) patterns were recorded using Malvern Panalytical Empyrean X-ray diffractometer with a Cu-K radiation source (0.1546 nm). Raman spectrum was collected using a Renishaw inVia Raman microscope with a 532 nm laser beam as the light source. The chemical elements and valence states were characterized by X-ray photoelectron spectrometer (XPS) (ESCALAB 250Xi, Thermo Fisher). The morphology, crystal structure and component of the as-synthesized products were characterized by transmission electron microscopy (TEM, Tecnai G2 F30), high-resolution TEM (HR-TEM), and scanning electron microscope (SEM, Hitachi, S-4800) with energy-dispersive X-ray (EDS) analysis function. The thermal infrared images were visualized with infrared detection device (Sigma ST9660). The electrical conductivity was measured by the four-point probe method (FT-330). The thermal conductivity was measured by the transient plane source method (Hotdisk).

### 4.5 Electromagnetic and electrochemical measurements

The tested NF@Co/C samples were made into a rectangle with a size of 22.86 mm  $\times$  10.16 mm and the thickness was measured. The  $S$  parameters of the NF@Co/C samples were characterized by the waveguide method by a vector network analyzer (VNA, Anritsu 37269D) in X band (8.2–12.4 GHz). In addition, the sample (10%) was mixed with wax and pressed into a toroidal shape with an outside diameter of 7.03 mm and an inside diameter of 3.00 mm. Then, its complex permittivity and complex permeability were measured by the coaxial method.

Electrochemical measurements of the NF@Co/C samples were carried out in a three-electrode system in a 6 M KOH aqueous electrolyte, with Pt foil as counter-electrode and Hg/HgO as reference electrode. CV, EIS and GCD measurements were performed by CHI660E electrochemical workstation. The CV curves were conducted at scan rates of 5, 10, 20, 30, 50, and 100 mV  $\text{s}^{-1}$  with the potential window between  $-0.2$  and  $0.6$  V. The GCD curves were conducted at current densities of 1, 2, 5, 10, 15, and 30 A  $\text{g}^{-1}$ . The EIS measurements were recorded in the frequency range from  $10^{-2}$  to  $10^5$  Hz with an AC potential amplitude of 5 mV.

## Conflicts of interest

There are no conflicts to declare.



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