

## Supporting Information

### Multi-role conductive hydrogels for flexible transducers regulated by MOFs for monitoring of human activities and electronic skin functions

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#### Text 1

#### Synthesis of Zn-MOF

First the Zinc metal organic framework (Zn-MOF) was developed utilizing the hydrothermal synthesis method. Trimesic acid was chosen as the organic linker, while  $\text{Zn}(\text{NH}_3)_2$  employed as the precursor metal. The standard procedure involved dissolving of 3g  $\text{Zn}(\text{NH}_3)_2$  in a 60ml solution composed of water, DMF, and ethanol, maintaining a 1:1:1 ratio. Subsequently, 1.5g of TMA was introduced into the mixture and stirred for 30 minutes at room temperature. Following this, the homogeneous blend was carefully transferred into a 100ml autoclave. The autoclave was subjected to a controlled temperature of 120°C within an oven for a duration of 12 hours. This thermal

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treatment yielded distinctive white crystals, which were subsequently isolated, subjected to multiple rounds of washing with ethanol and water, and then carefully filtered. The end product was subjected to vacuum drying and set aside for subsequent comprehensive characterizations.

### Characterization

Fourier-transformed spectroscopy was utilized to discern the functional groups of TMA within the Zn-MOF, and their respective spectra are depicted in Figure S1. As per existing literature, the absorption peak at 3454  $\text{cm}^{-1}$  signifies the OH stretching vibration of a water molecule, complemented by deformation vibrations at 719  $\text{cm}^{-1}$ . These observations suggest that water not only acts as a solvent but also assumes a pivotal role in the Zn-MOF preparation[1]. Furthermore, the discernible bands at 1627-1579  $\text{cm}^{-1}$  and 1453-1378  $\text{cm}^{-1}$  are attributed to the asymmetric and symmetric vibrations of the interconnected carboxylic groups, affirming the successful synthesis of Zn-MOF. This conclusion gains further substantiation from the presence of a distinctive band at 578  $\text{cm}^{-1}$ , indicating the characteristic vibration of metal ions and TMA (O-Zn)[2-4].

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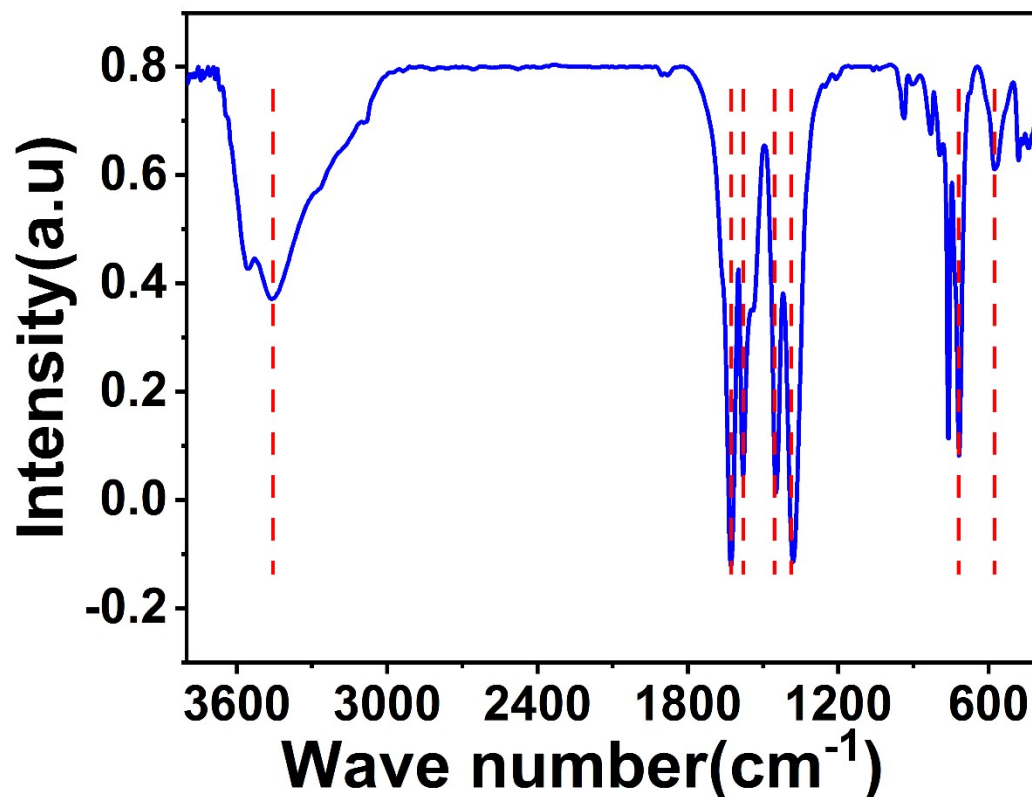


Figure S1 FTIR spectra of the prepared Zinc based metal organic frame work.

The X-ray diffraction analysis of the synthesized MOFs is depicted in Figure S1. This graphical representation reveals and affirming the successful preparation of the Zn-MOF. Importantly, there is no evidence of peak splitting, indicating the presence of a single-phase crystallographic structure within the Zn-MOF. According to the previously reported literatures, in the XRD patterns Zn peaks can be observed at 2 thetas of 15.6, 18.5, 19.7, 20.2, 22.4, 25.6, 31.8, 34.6, and 36.2, All these values suggested the successfully synthesis of the Zn-MOF[5-7].

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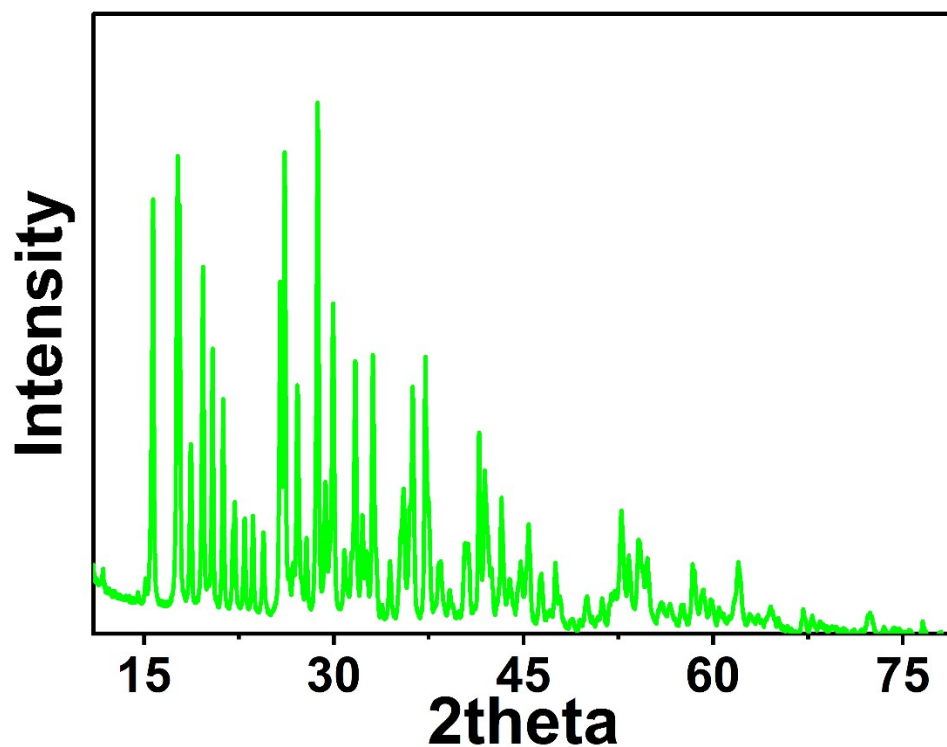


Figure S2 XRD analysis of the prepared Zn-MOF.

Furthermore, the Zn-MOF underwent characterization using Energy-Dispersive X-ray Spectroscopy (EDS) and scanning electron microscopy (SEM) as given in Figure. EDS analysis, as depicted in Figure a and table b confirmed the presence of Zinc metal with the 24% in the MOF, as evidenced by the identified elemental peaks. Moreover, SEM images of the MOFs revealed a distinct cylindrical morphology, as illustrated in Figure c. However, according to the SEM image the size of the Zn-MOF was different in length.

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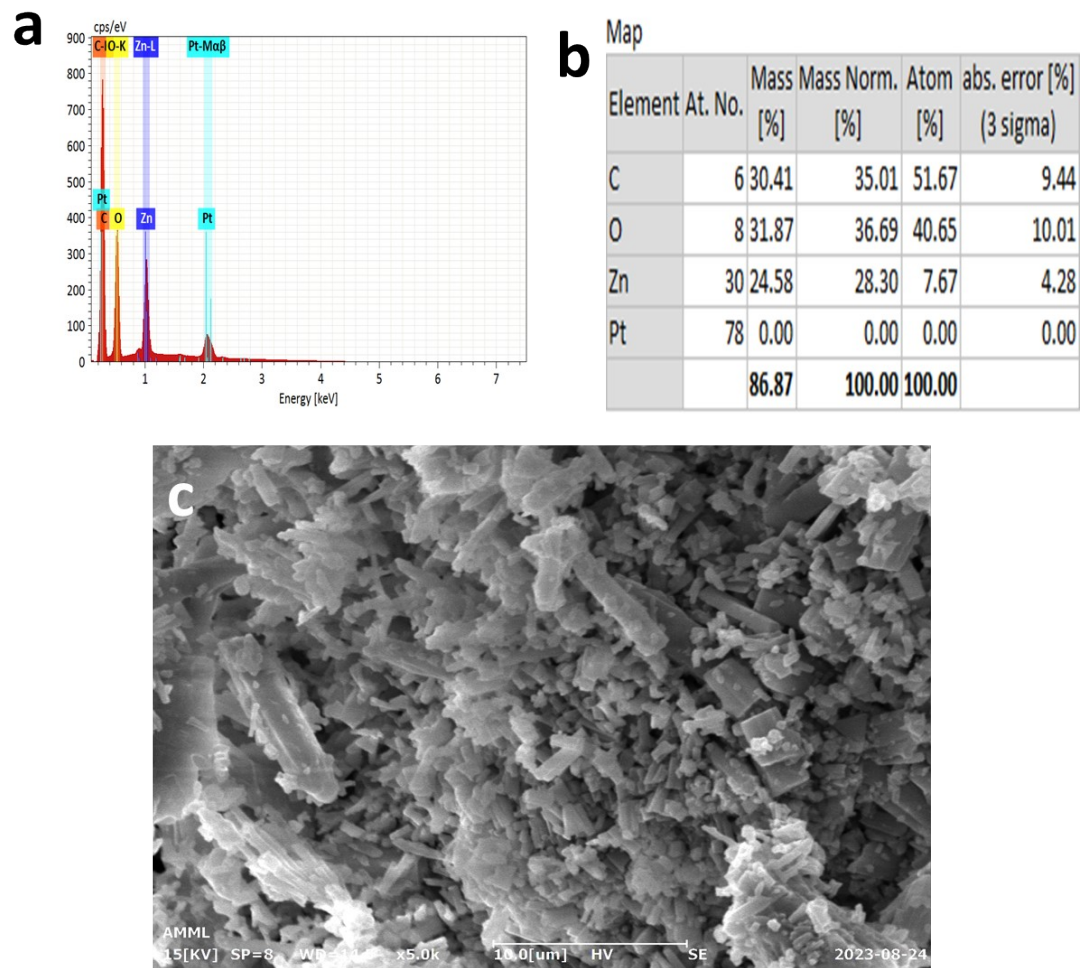


Figure. S3 (a-b) Illustration of the EDS analysis of the Zn-MOF (c) SEM study of the Zn-MOF.

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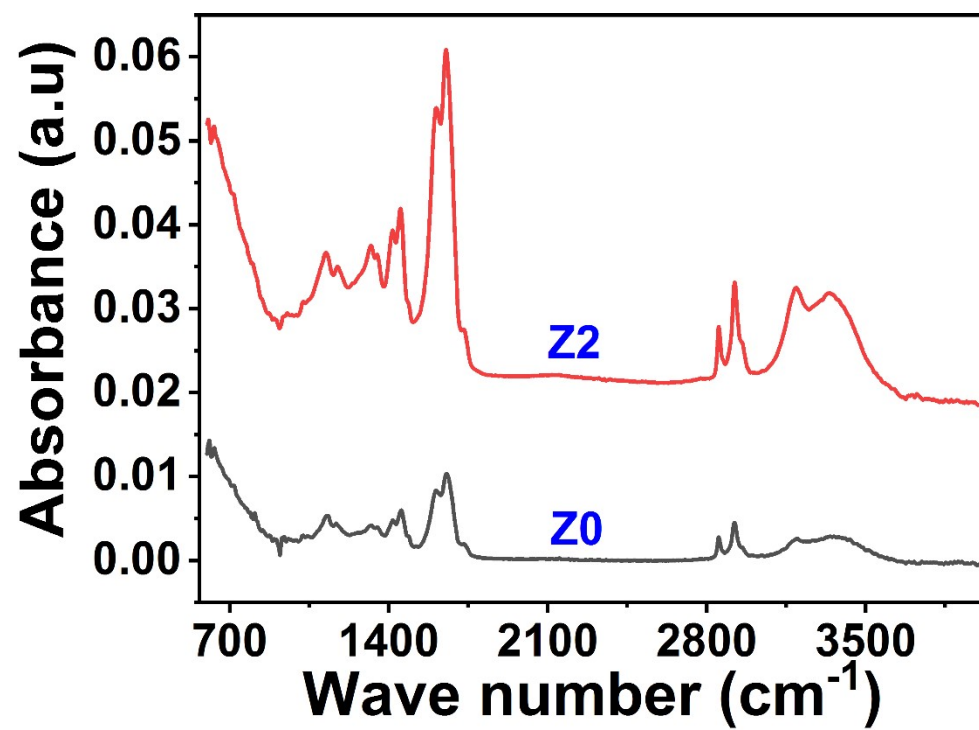


Figure S4 FTIR spectroscopy of the hydrogels

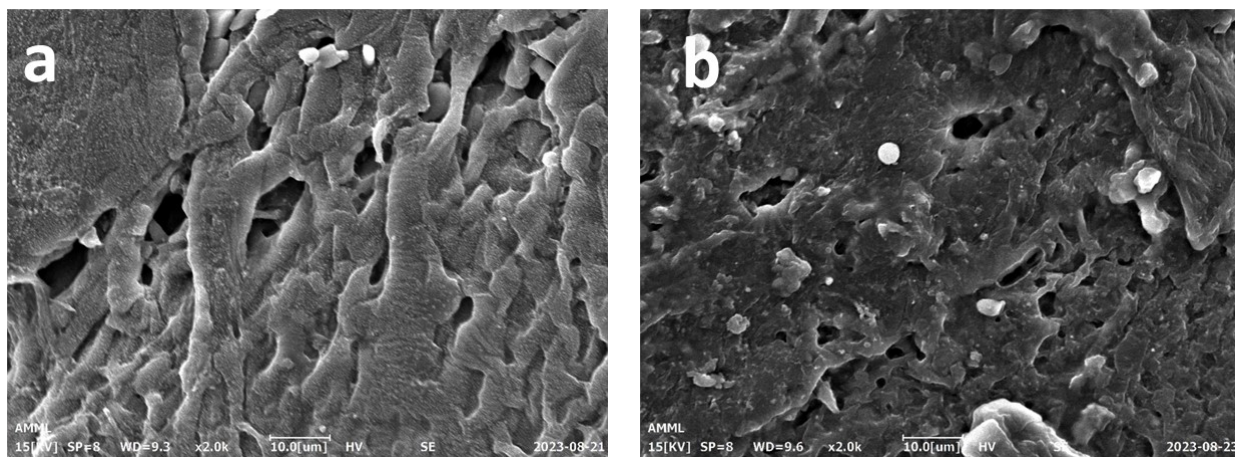


Figure S5 SEM images of the prepared hydrogels (a) Z0 and (b) Z2

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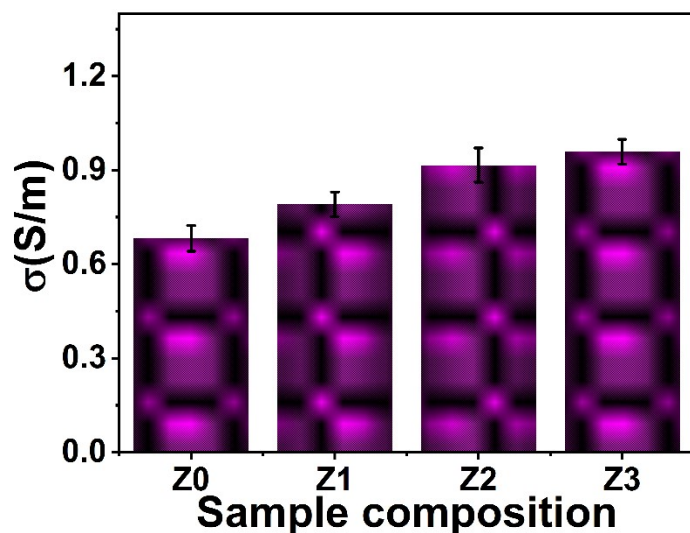


Figure S6 conductivity of the prepared p(DDMA-AM- DMAEMC) Zn-MOFs hydrogels.

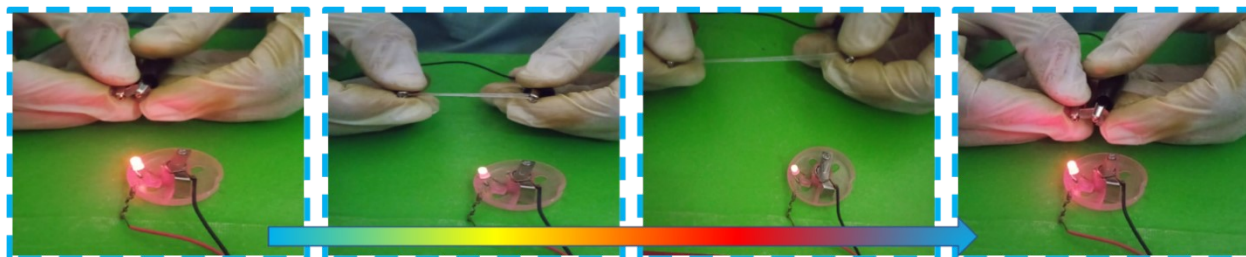


Figure S7. Visualization of the strain sensitivity of the p(DDMA-AM- DMAEMC) Zn-MOF hydrogels

## Reference

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