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Supporting Information

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3 3D Ti₃C₂T_x MXene-matrigel with electroacoustic stimulation 4 to promote the growth of spiral ganglion neurons

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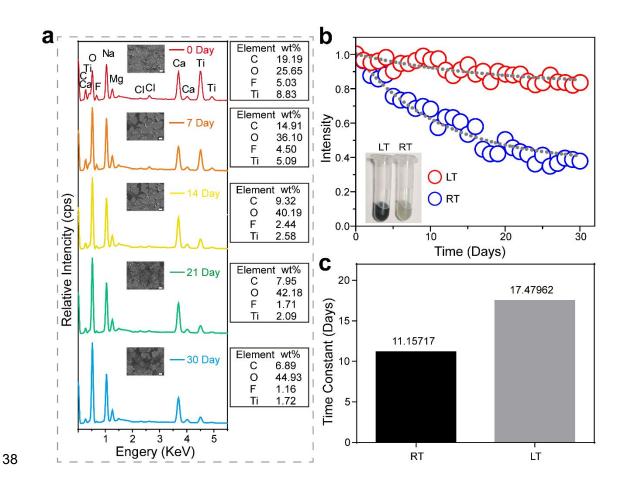
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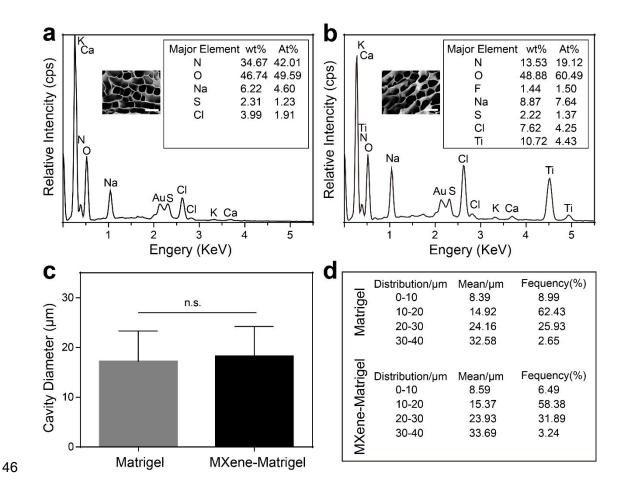
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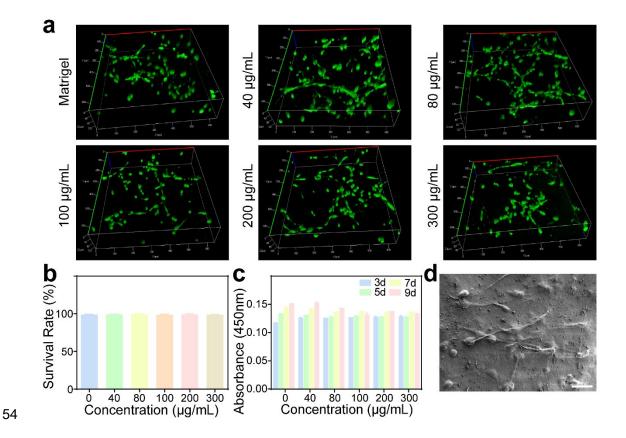
37 <u>Supplementary figures and figure legends</u>



Supplementary Figure 1. The $Ti_3C_2T_x$ MXene nanosheet degraded slowly over time. (a) SEM-EDX results of $Ti_3C_2T_x$ MXene stored for different times. Scale bar=2 µm. (b) Absorbance attenuation curve of $Ti_3C_2T_x$ MXene aqueous solution stored at low temperature (LT) and room temperature (RT) in the dark. The dotted lines were fitted according to the exponential decay function $f(x) = y_0 + Ae^{-x/\tau}$. Lower left was the color contrast of $Ti_3C_2T_x$ MXene (0.3mg/mL) stored in LT and RT at day 30. (c) Time constants of $Ti_3C_2T_x$ MXene in LT and RT environments.



Supplementary Figure 2. $Ti_3C_2T_x$ MXene was successfully integrated into the matrigel. (a) SEM-EDX mapping of mareigel hydrogel. (b) SEM-EDX mapping of $Ti_3C_2T_x$ MXene-mareigel hydrogel. wt%: weight percentage, At%: atom percentage. Scale bar=20 µm. (c) The cavity diameter of the matrigel and $Ti_3C_2T_x$ MXene-matrigel hydrogel. (d) Statistical results of grouping the cavity diameters of the matrigel and $Ti_3C_2T_x$ MXene-matrigel hydrogel.



55 Supplementary Figure 3. Ti₃C₂T_x MXene-matrigel hydrogel was an excellent scaffold 56 for SGNs. (a) Representative fluorescence images of live/dead staining of SGNs 57 cultured in Ti₃C₂T_x MXene hydrogels with different concentrations. Entire cytoplasm 58 of live cell was labeled with Calcein-AM (green), and nuclei of dead cell was labeled 59 with EthD-1 (red). (b) The proportion of Calcein-AM positive cells. (c) The cytotoxicity 60 of matrigel hydrogel and Ti₃C₂T_x MXene-matrigel hydrogels with different 61 concentrations to SGNs from CCK-8. (d) Representative SEM image of SGNs cultured 62 in Ti₃C₂T_x MXene hydrogel for three days. Scale bar=20 µm. Data are presented as 63 mean \pm SD.