Supplementary Information

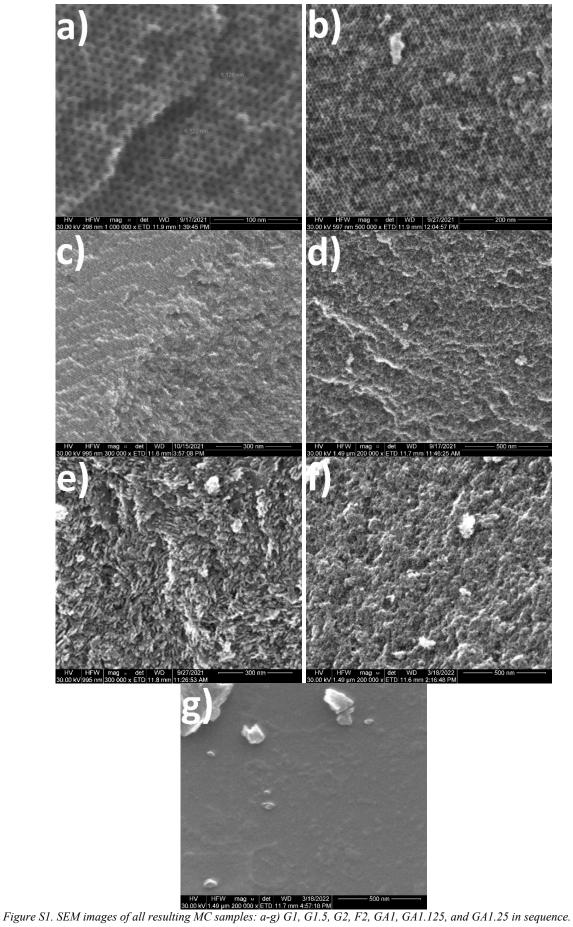
Advancing Mesoporous Carbon Synthesis for Supercapacitors: A Systematic Investigation of Cross-linking Agent Effects on Pore Structure and Functionality

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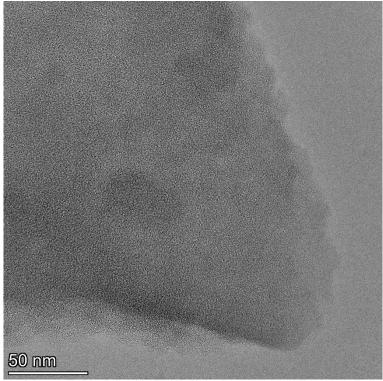


Figure S2. TEM image for sample GA1.25.

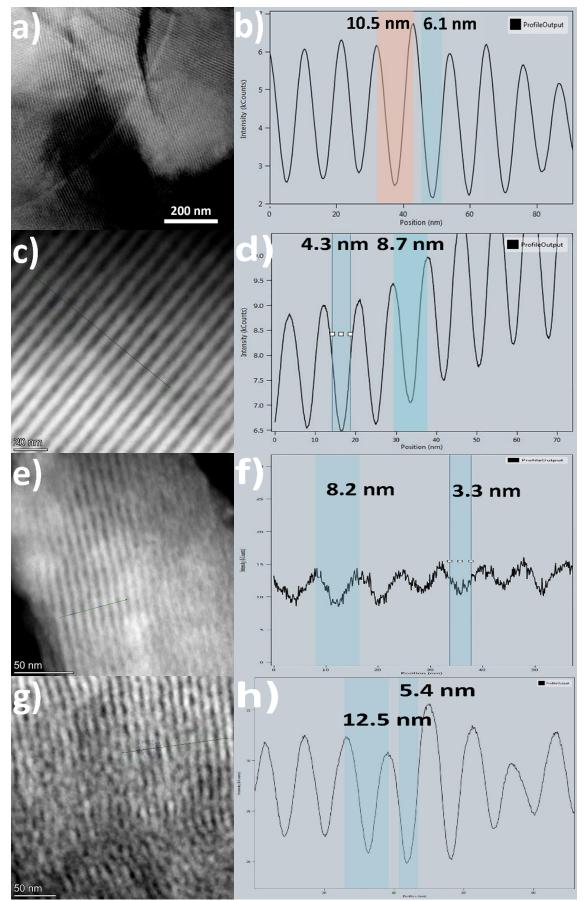


Figure S3 HAADF STEM images and line profile of processed HAADF intensity for ordered mesoporous samples: a, b) G1, c, d) G1.5, e, f) G2, and g, h) GA1, respectively.

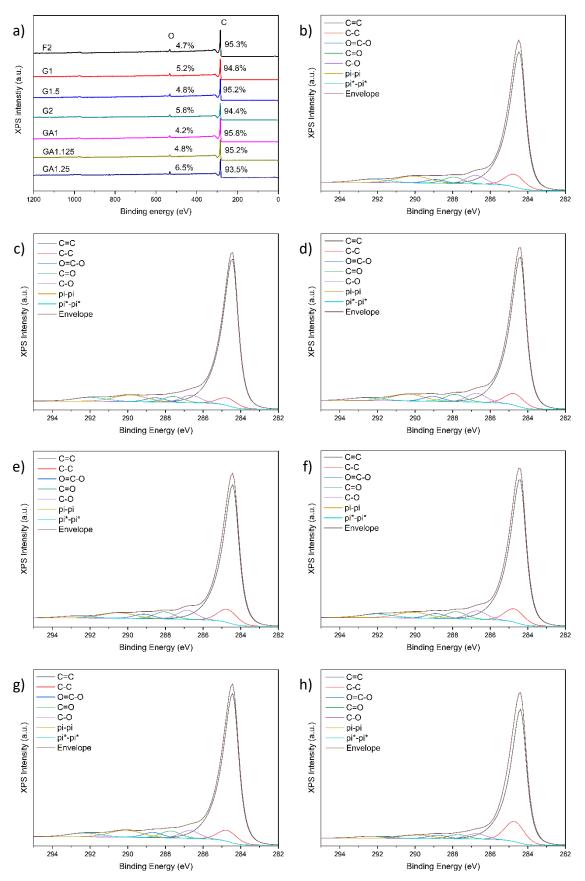


Figure S4. Summary of wide-scan XPS spectra (a) and deconvolution of C 1s peaks for MCs (b-h): F2, G1, G1.5, G2, GA1, GA1.125, and GA1.25, respectively.

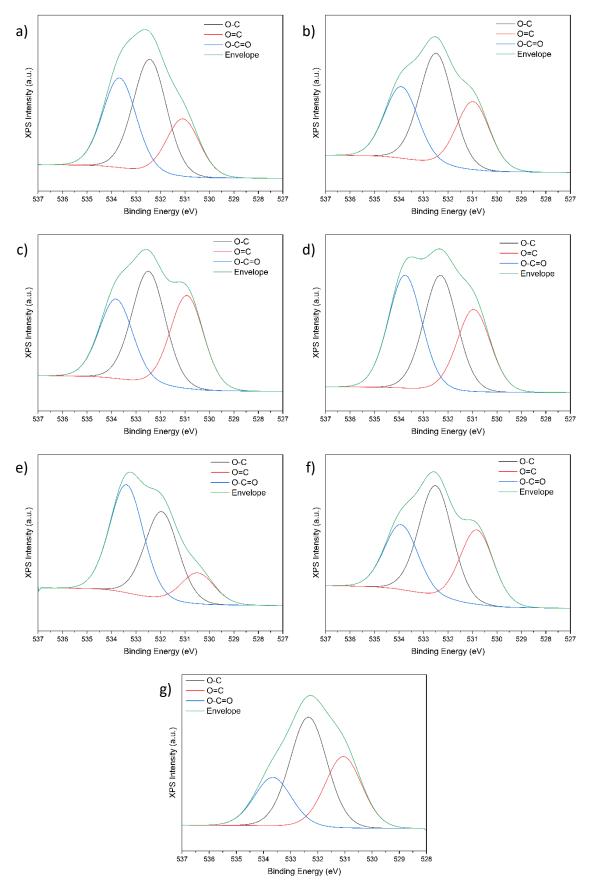


Figure S5. Deconvolution of XPS O 1s peaks for MCs (a-g): F2, G1, G1.5, G2, GA1, GA1.125, and GA1.25, respectively.

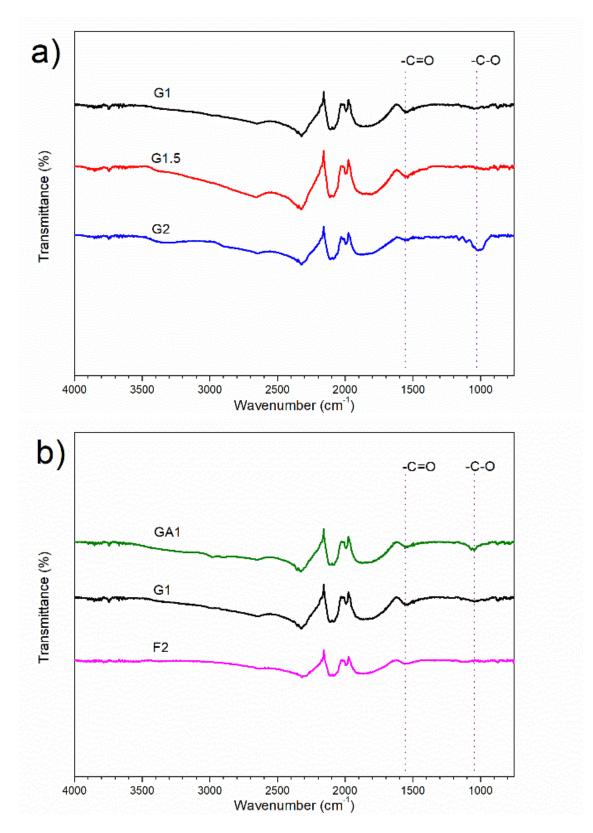


Figure S6. IR spectra: a) MCs prepared by glyoxal cross-linker, and b) comparison of MCs with different cross-linkers.

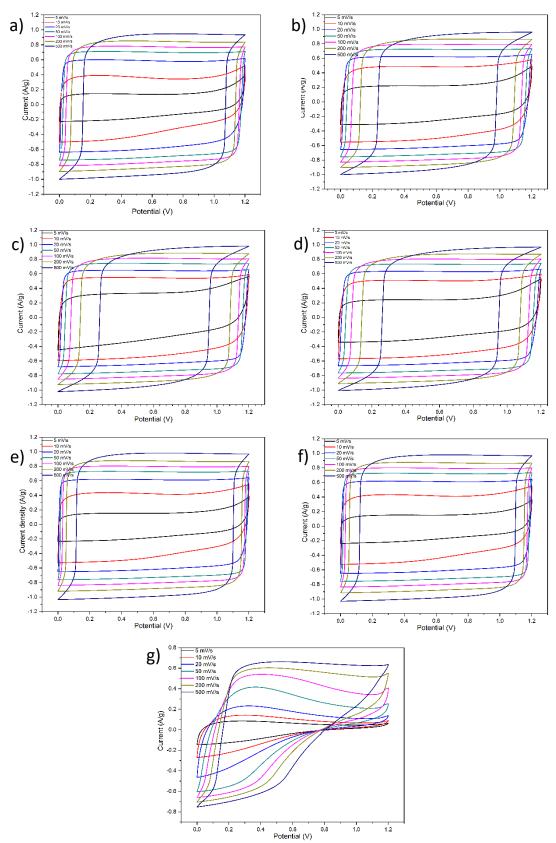
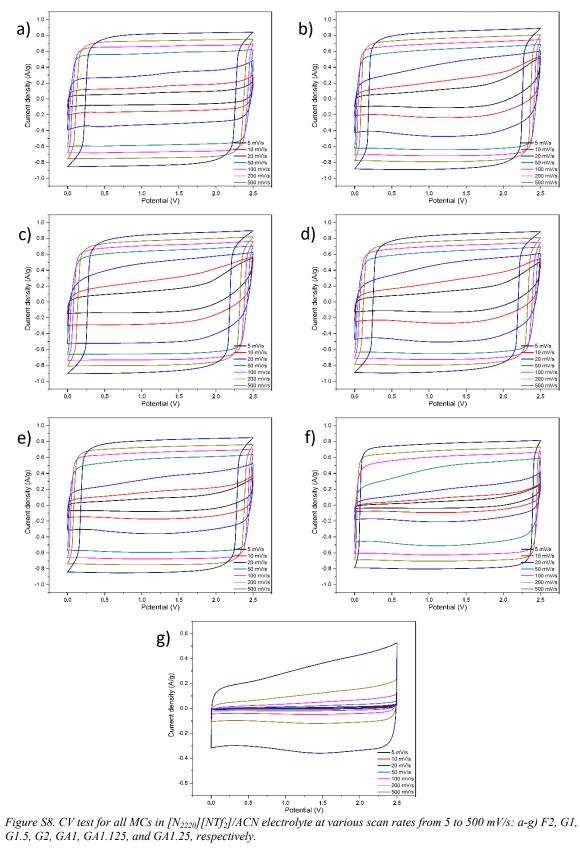


Figure S7. CV test for all MCs in 6 M KOH electrolyte at various scan rates from 5 to 500 mV/s: a-g) F2, G1, G1.5, G2, GA1, GA1.125, and GA1.25, respectively.



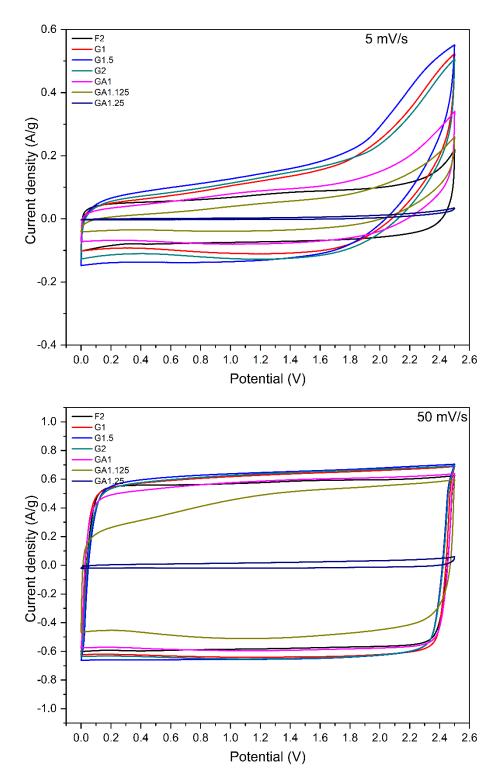


Figure S9. CV test for all MC electrodes in $[N_{2220}]$ [NTf₂]/ACN electrolyte at a scan rate of 5 and 50 mV/s, respectively.

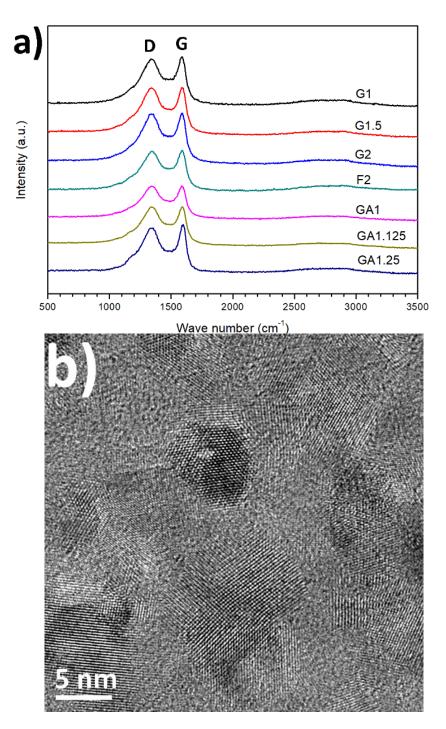


Figure S10. a) Raman spectra of resultant MC samples, and b) representative TEM image of random graphite crystalline structures in sample F2.