

Supplementary

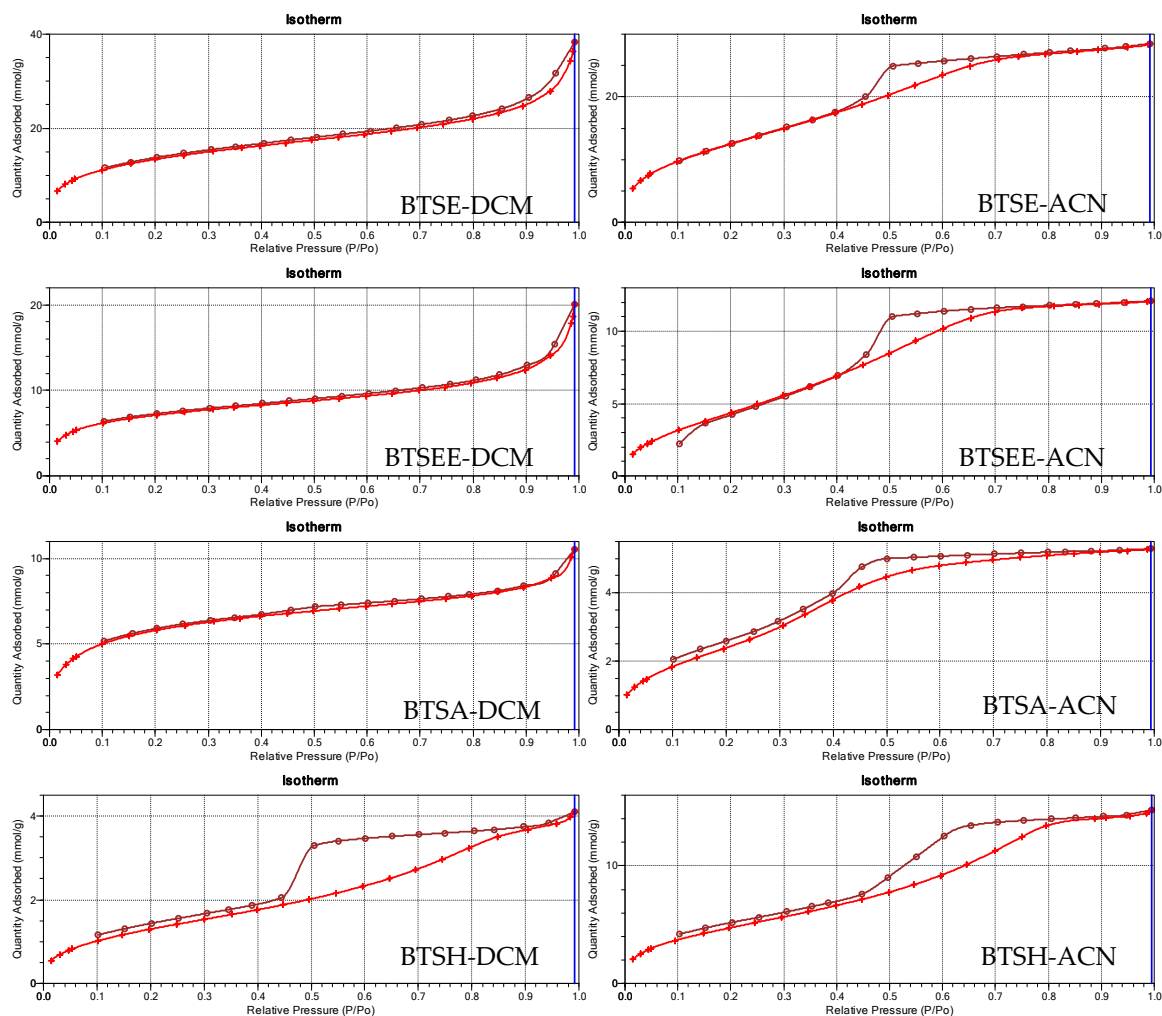
R-Silsesquioxane-Based Network Polymers by Fluoride Catalyzed Synthesis: An Investigation of Cross-linker Structure and its Influence on Porosity

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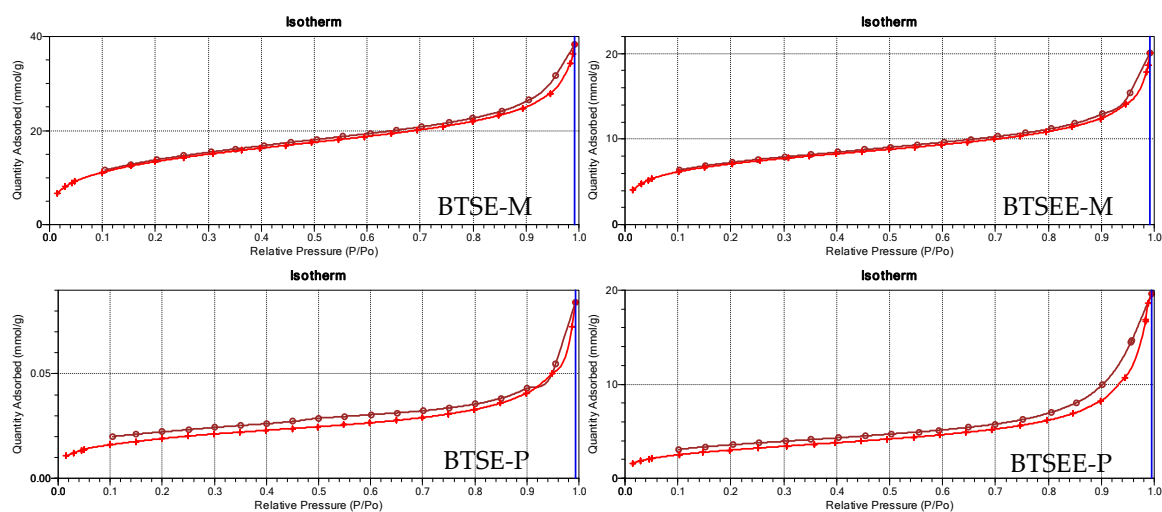
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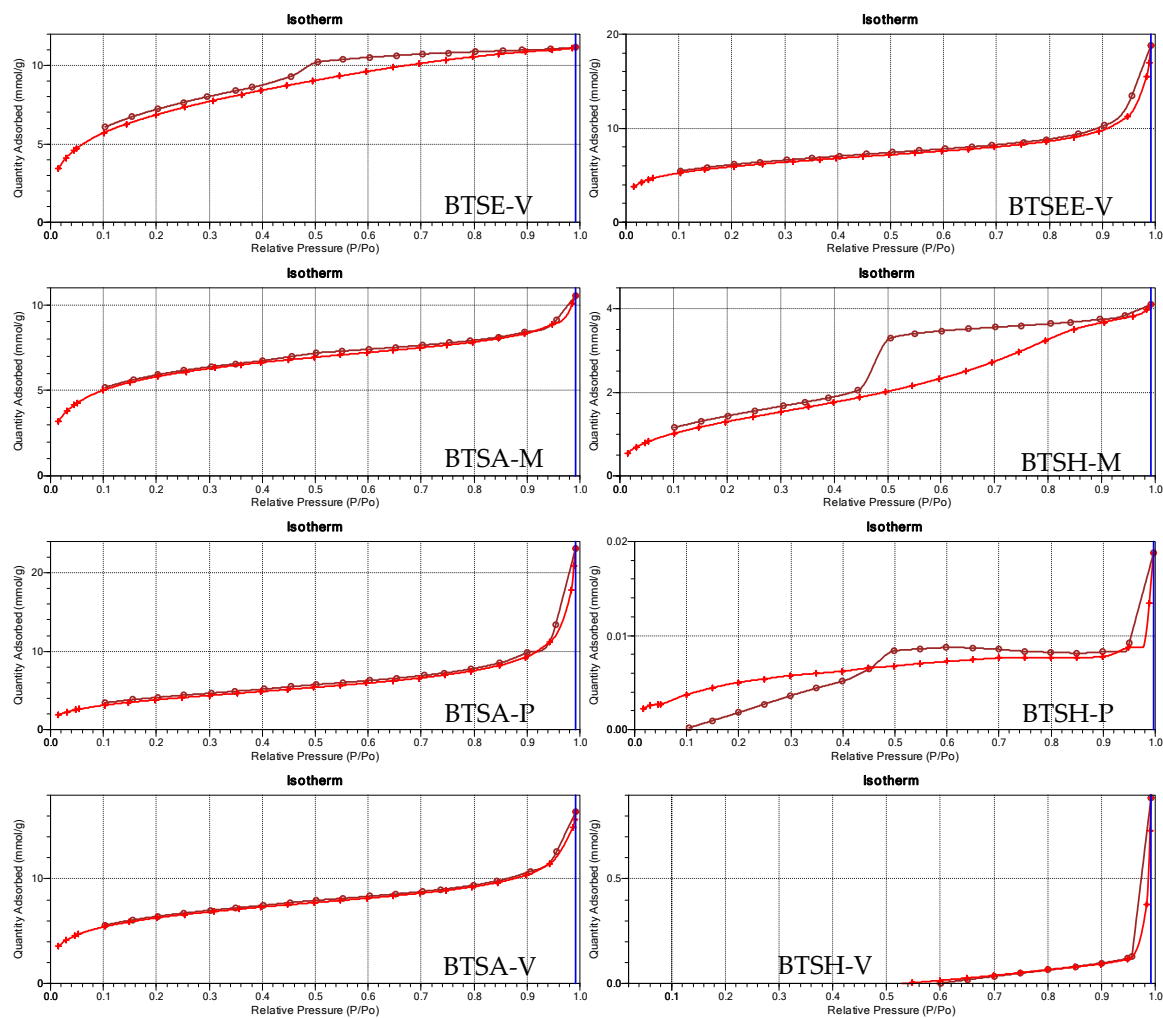
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S8	Pore size distribution graphs of materials from reactions with different corner-groups and cross-linkers.



(a)





(b)

Figure S1. (a) Isotherm graphs from reactions with different cross-linkers in dichloromethane (DCM) and acetonitrile (ACN), Plus (light red) = adsorption, circle (dark red = desorption). (b) Isotherm graphs of materials from reactions with different corner-silanes and cross-linkers, P = phenyl, V = vinyl, M = methyl from DCM, Plus (light red) = adsorption, circle (dark red = desorption).

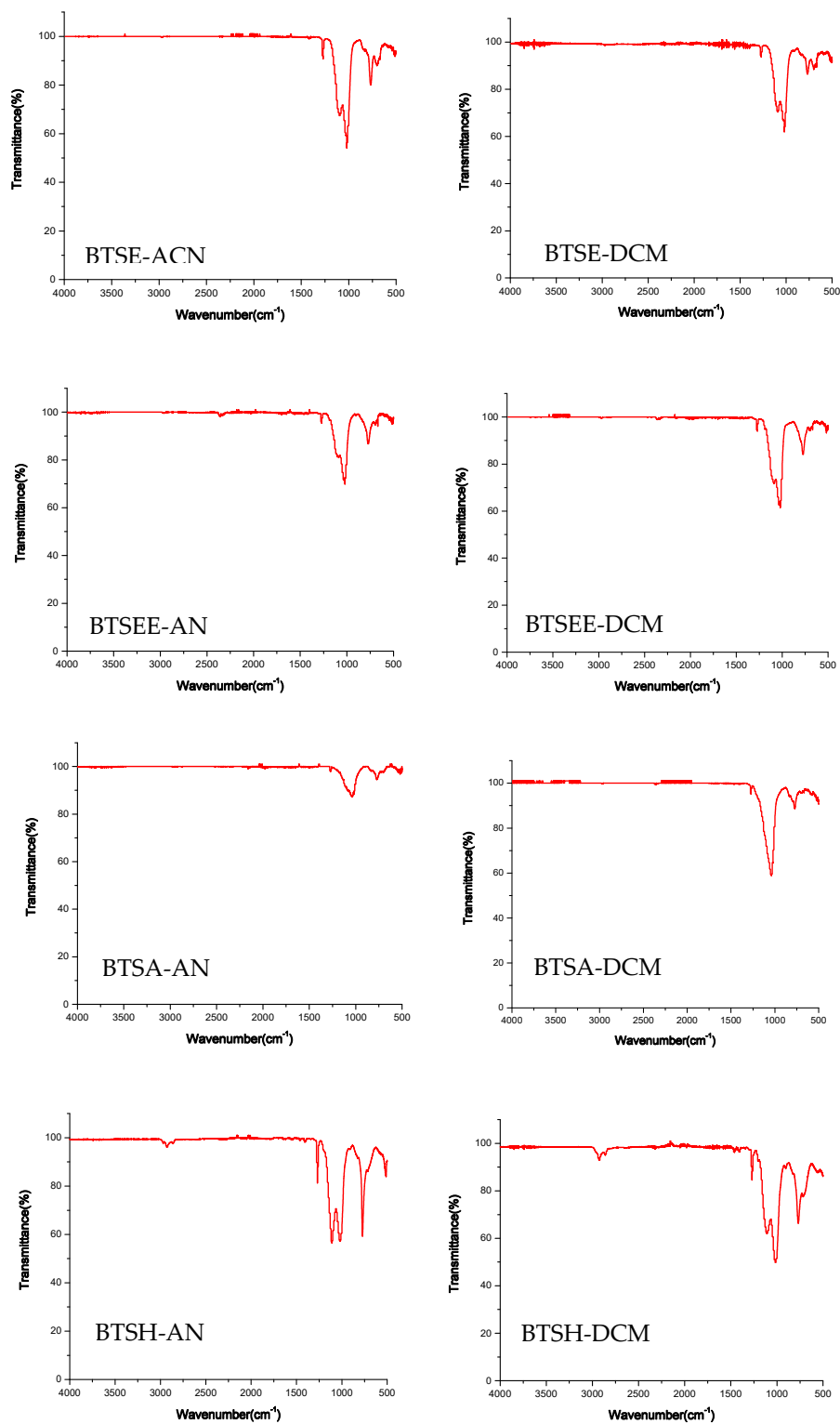


Figure S2. IR spectra of materials from reactions with different cross-linkers in dichloromethane (DCM) and acetonitrile (ACN).

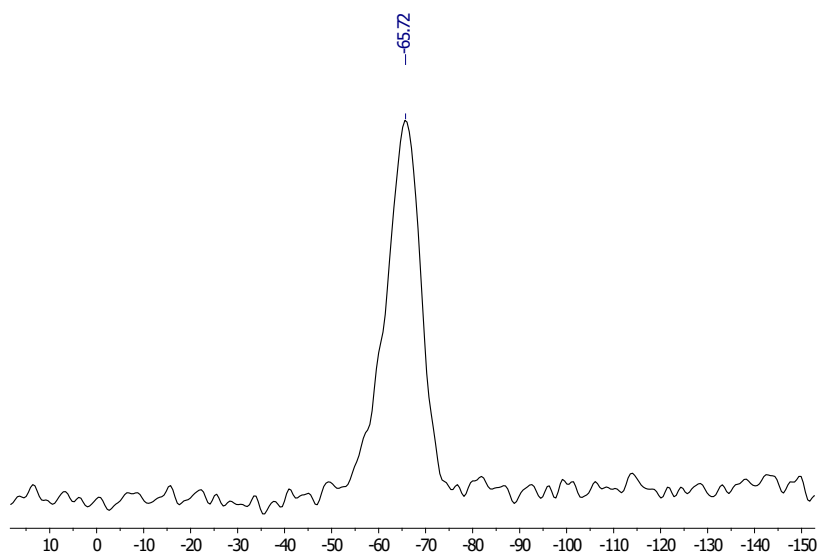
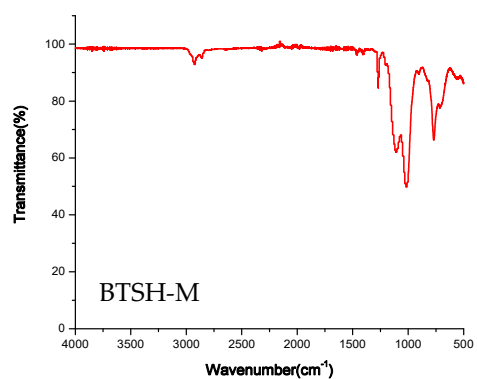
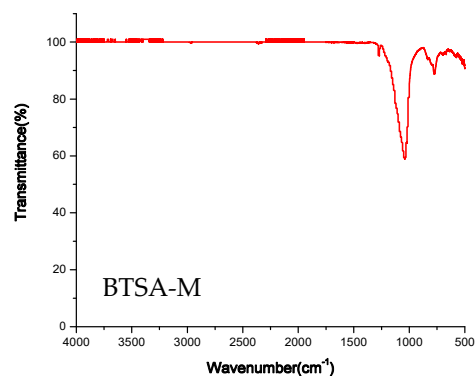
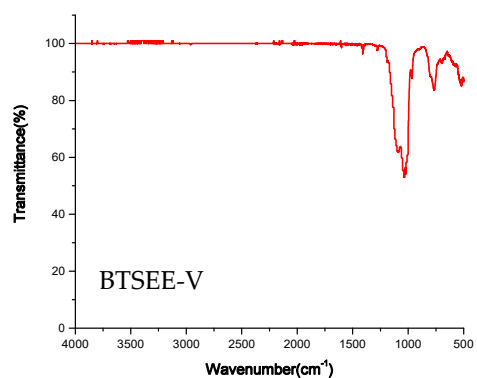
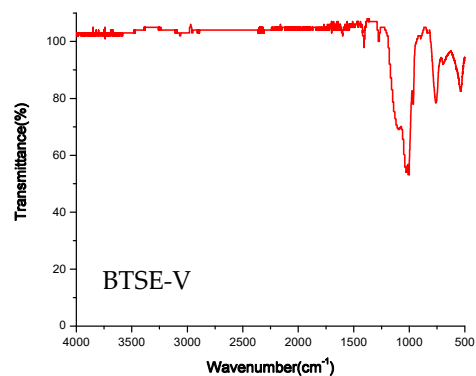
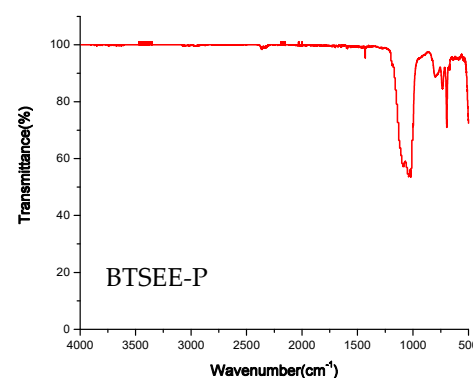
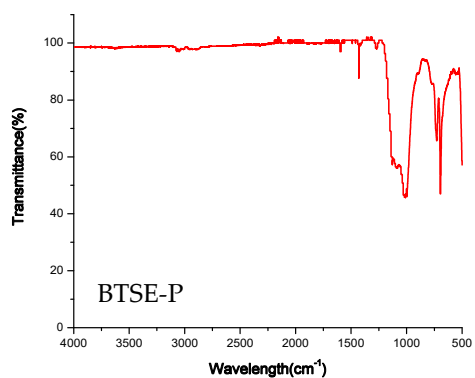
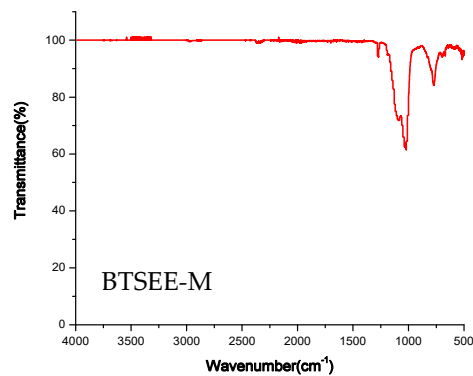
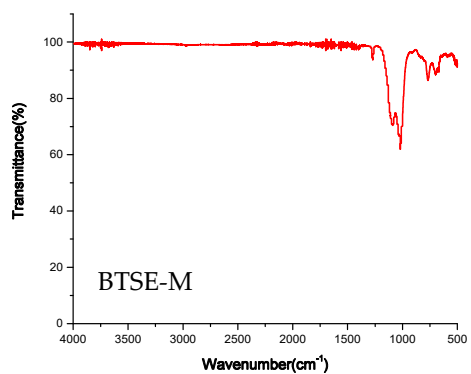


Figure S3. ^{29}Si NMR of methyl silsesquioxane based network polymer with hexyl spacer made in DCM. Note there are not extensive shoulders or side peaks indicating near complete reaction.



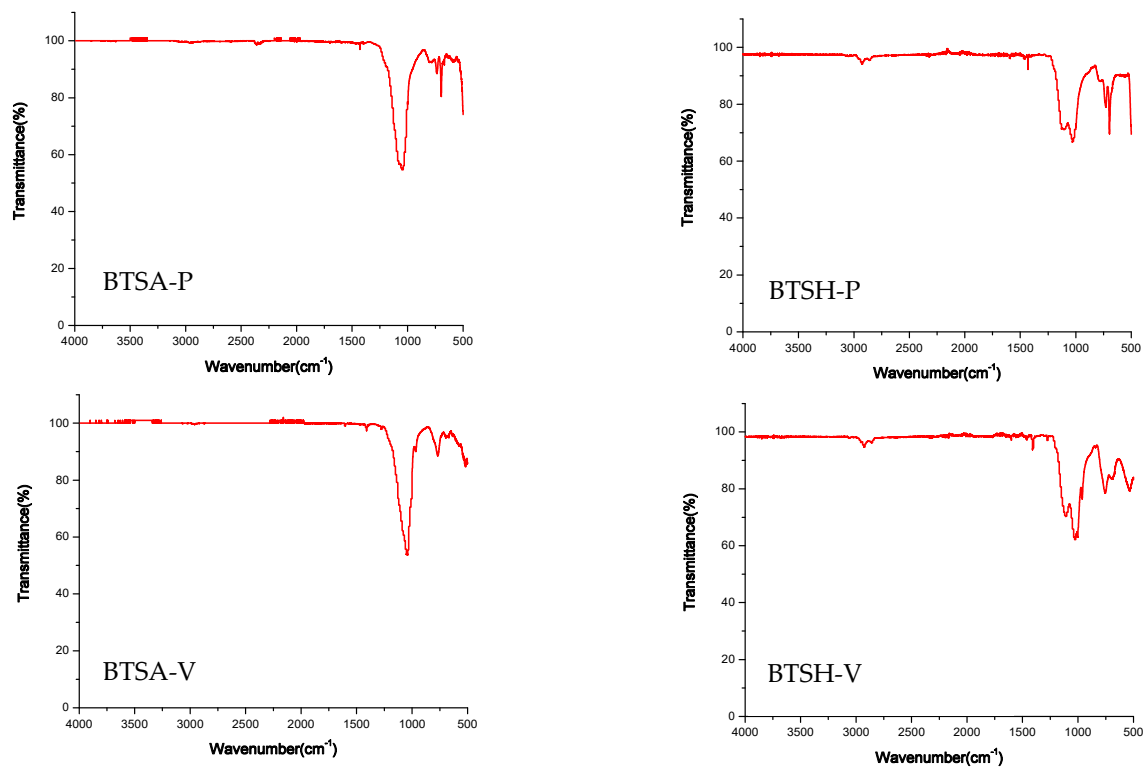


Figure S4. IR spectra of materials from reactions with different corner-silanes and cross-linkers, P = phenyl, V = vinyl, M = methyl from DCM.

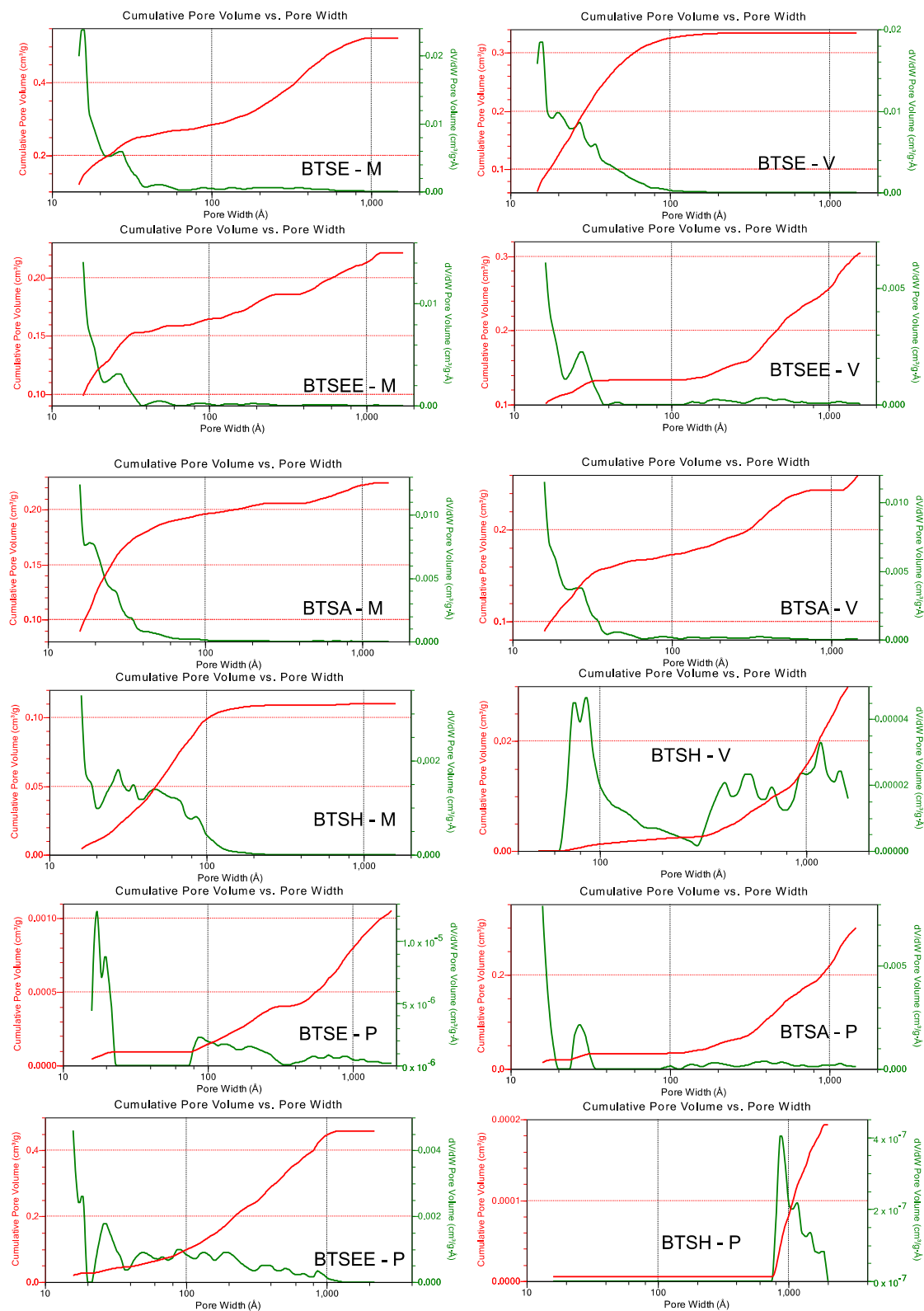


Figure S5. DFT pore size distribution plots and cumulative pore volume of materials from reactions with different corner-silanes and cross-linkers, P = phenyl, V = vinyl, M = methyl from DCM.