

Supporting Information (SI)

Hydrolytically Degradable, Dendritic Polyglycerol Sulfate based Injectable Hydrogels using Strain Promoted Azide-Alkyne Cycloaddition Reaction

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1 Supplementary Results

1.1 Characterization of dPGS N₃

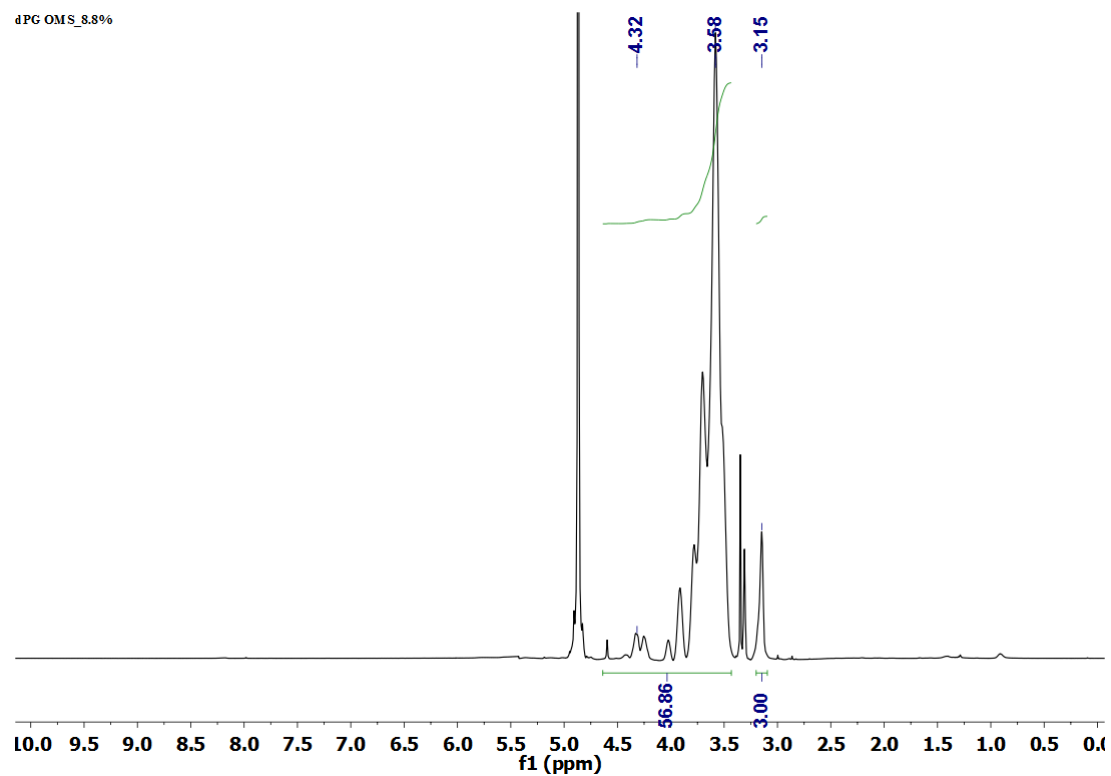


Figure S1. ¹H NMR of dPG OMs (DF 8.8 %)

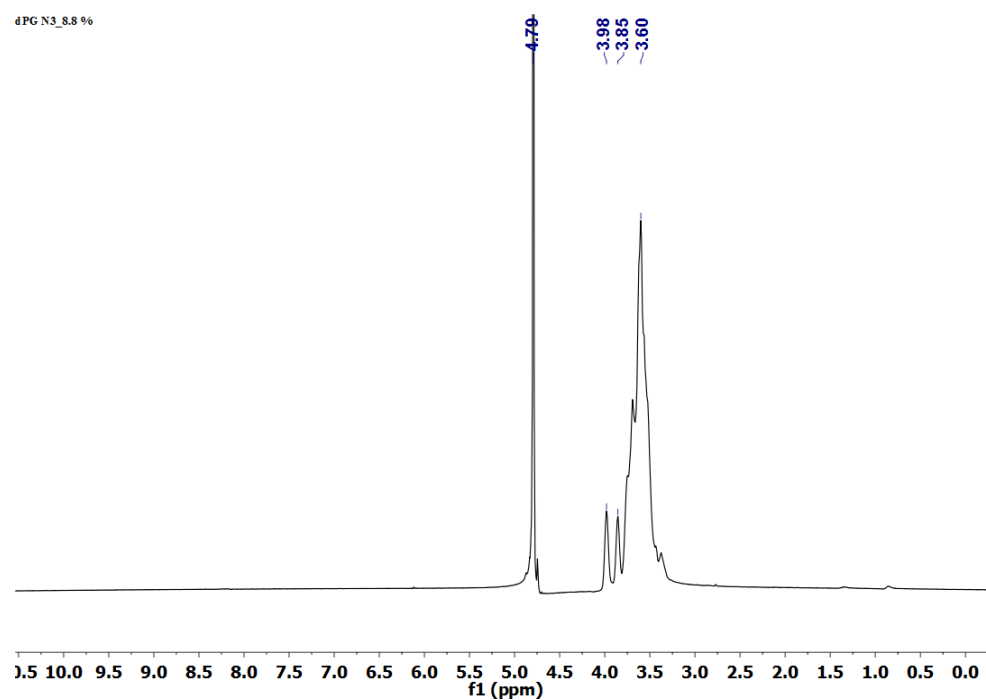


Figure S2. ¹H NMR of dPG N₃ (DF 8.8 %)

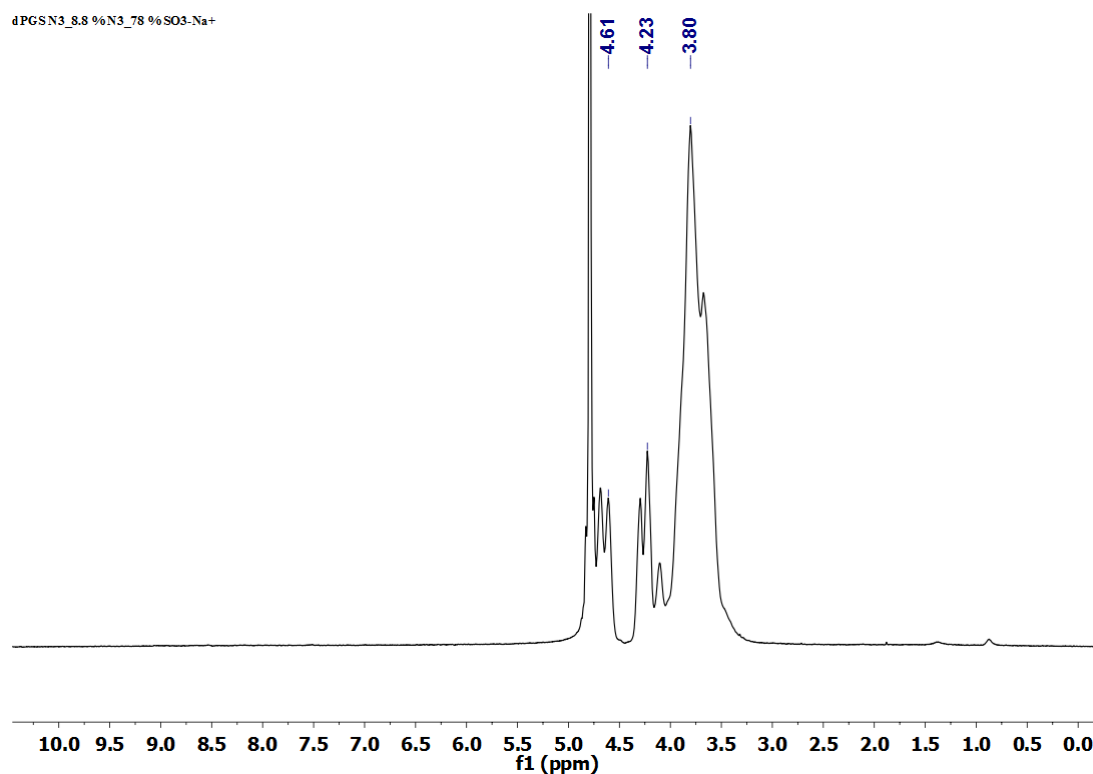


Figure S3. ¹H NMR of dPGS N₃ (DF 8.8 % N₃, 78 % SO₃⁻Na⁺)

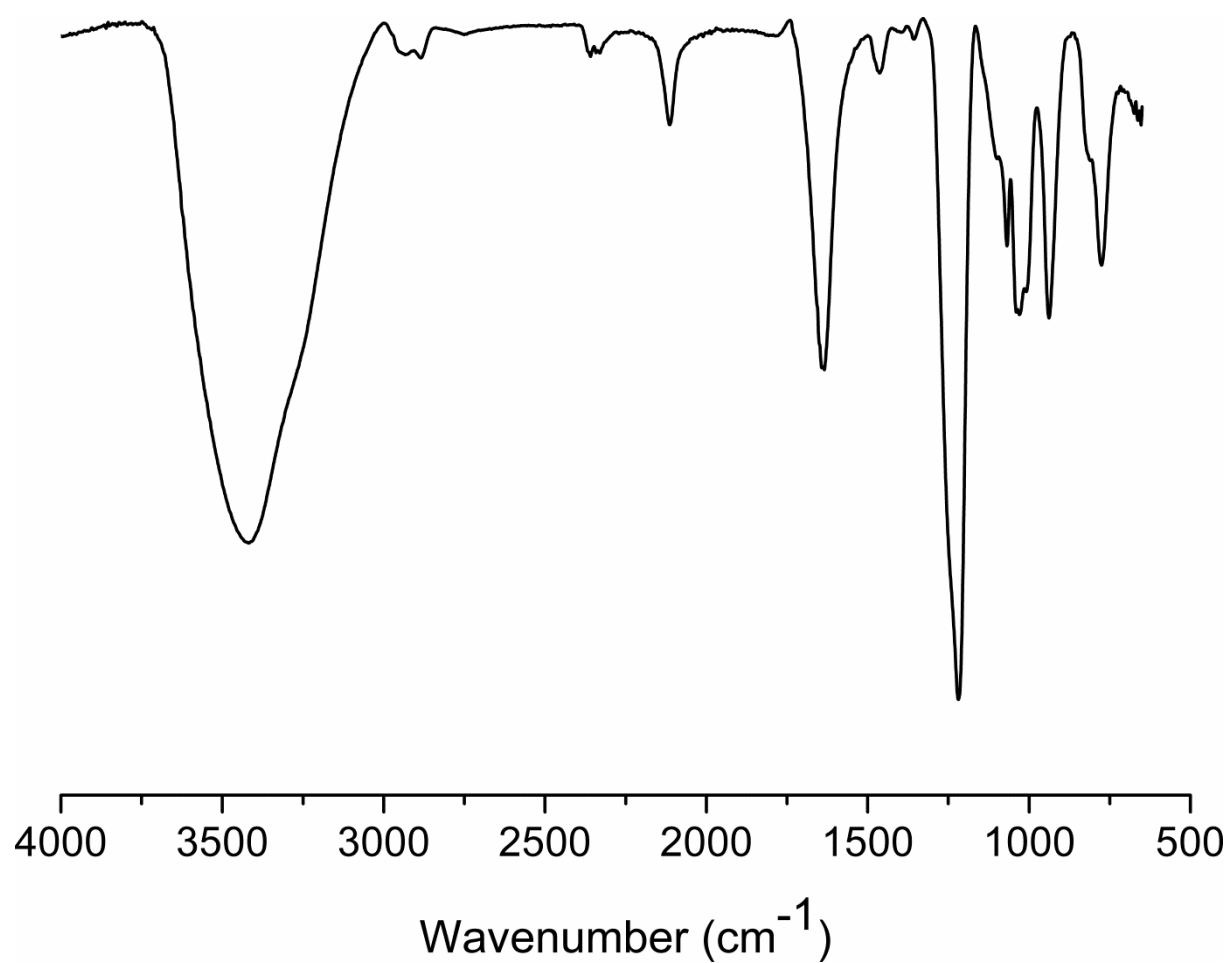


Figure S4. IR spectra of dPGS N₃.

1.2. Characterization of bicyclo[6.1.0]non-4-yn-9-ylmethyl *N*-(2-propyn-1-yl) carbamate

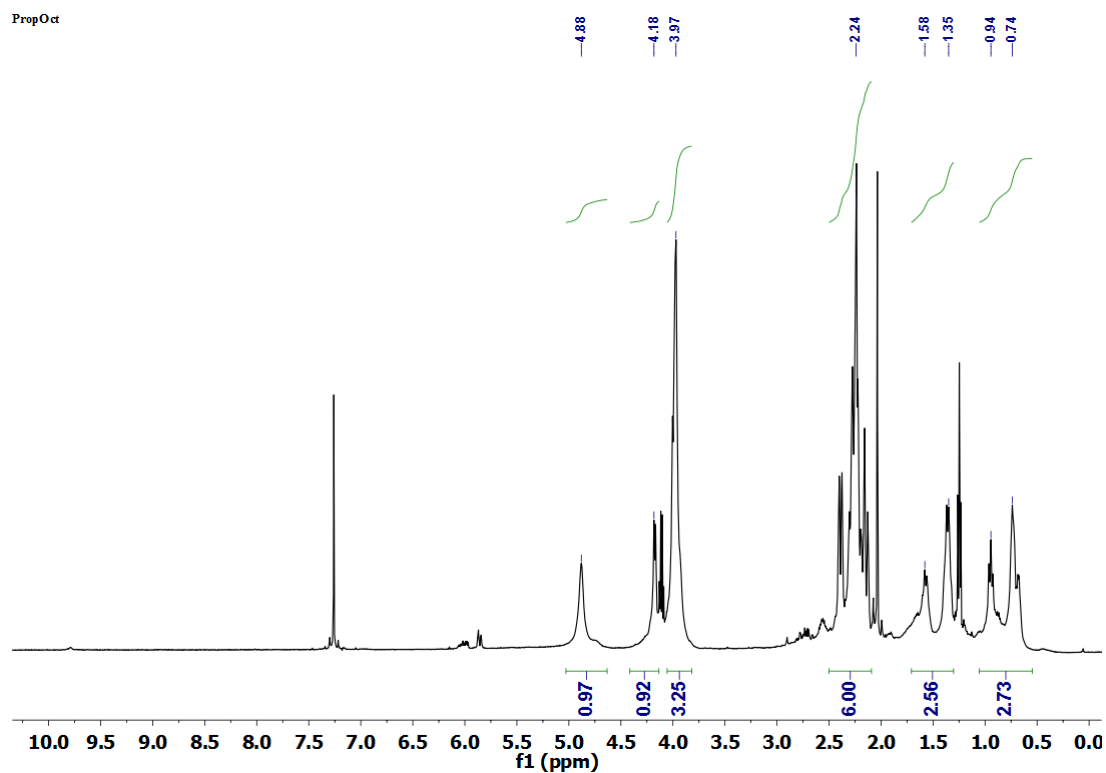


Figure S5. ^1H NMR of bicyclo[6.1.0]non-4-yn-9-ylmethyl *N*-(2-propyn-1-yl) carbamate

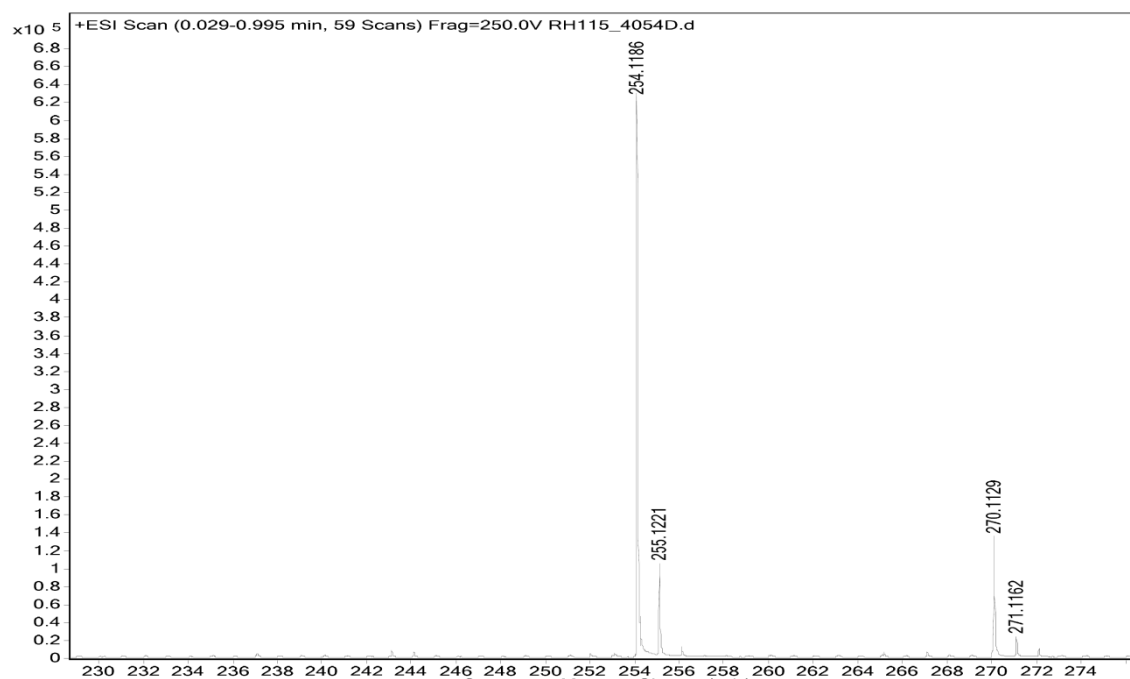


Figure S6. ESI MS of bicyclo[6.1.0]non-4-yn-9-ylmethyl *N*-(2-propyn-1-yl) carbamate

1.3. Characterization of PEG-PCL-DIC

1.3.1. Characterization of PEG-OH

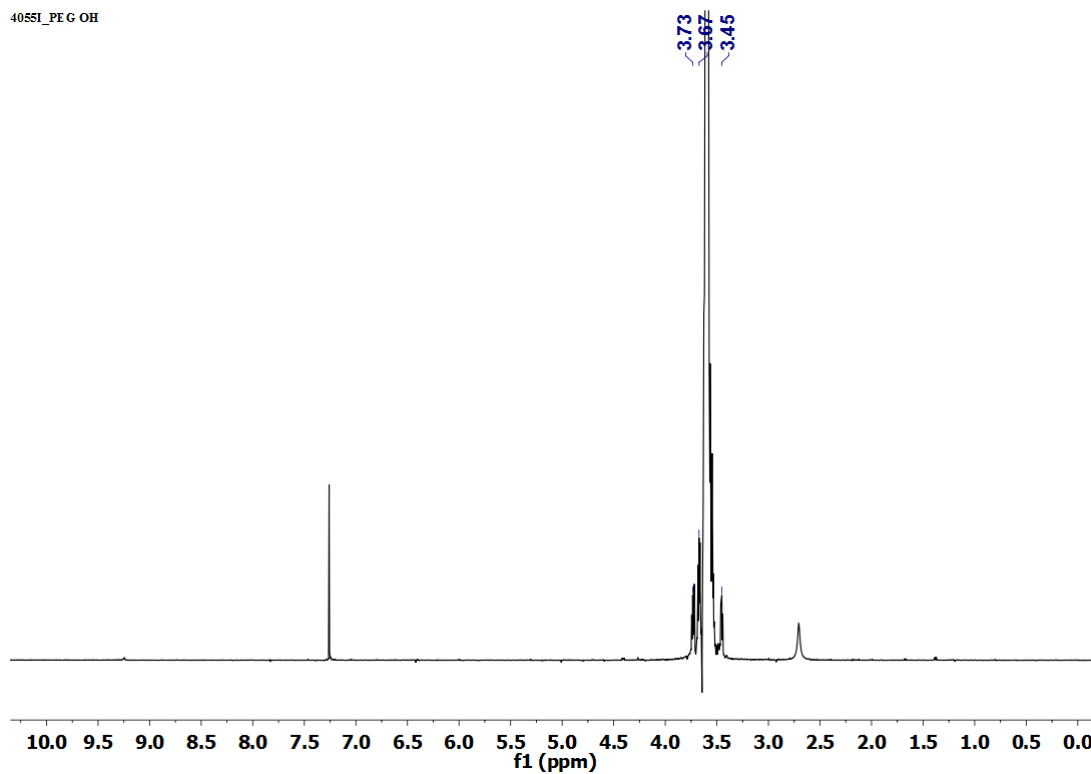


Figure S7. ^1H NMR of PEG-OH

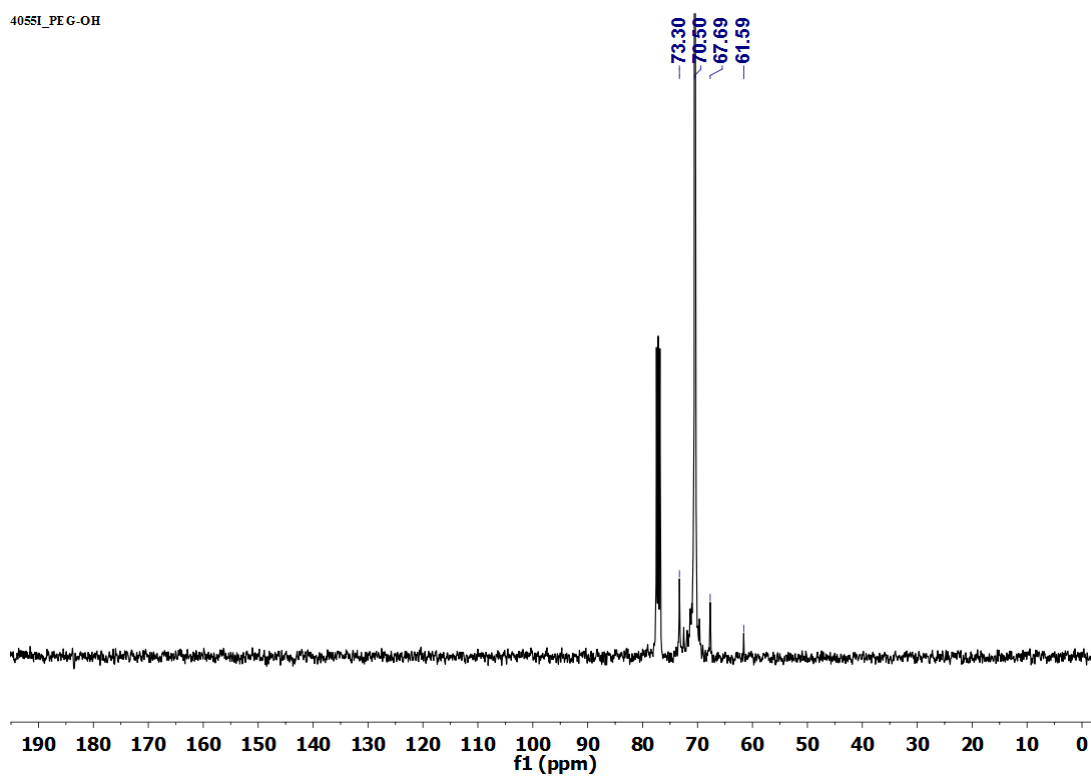


Figure S8. ^{13}C NMR of PEG-OH

1.3.2. Characterization of PEG-PCL-OH

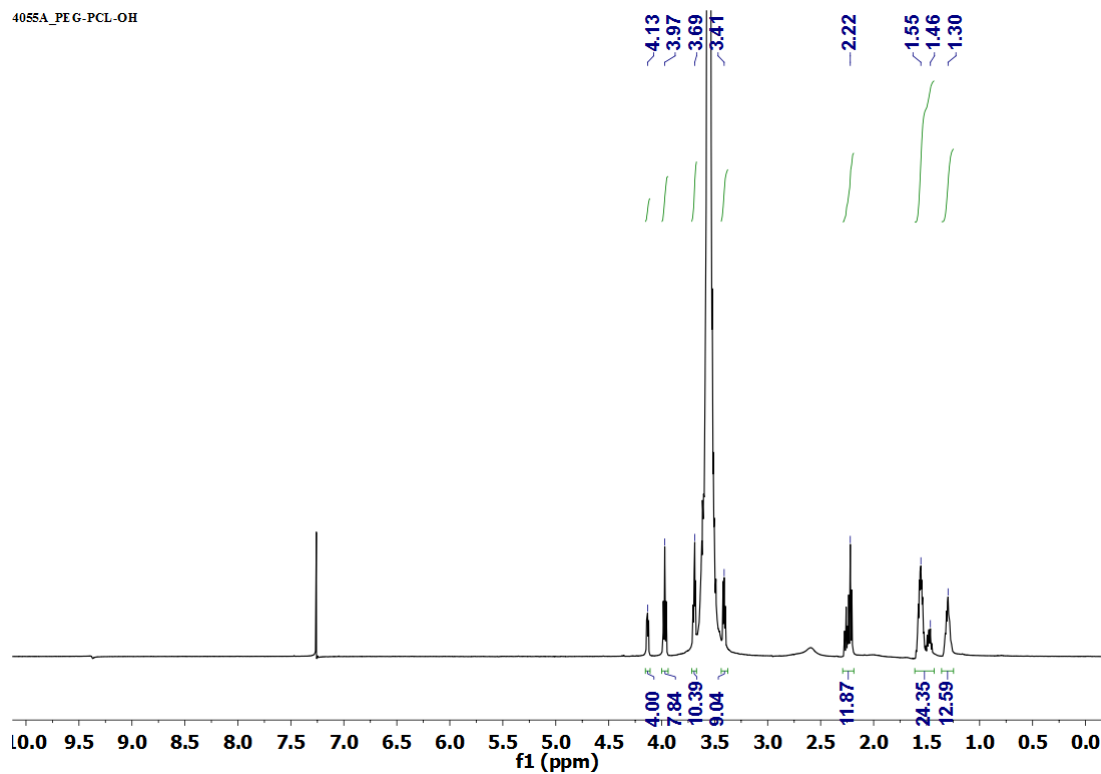


Figure S9. ^1H NMR of PEG-PCL-OH

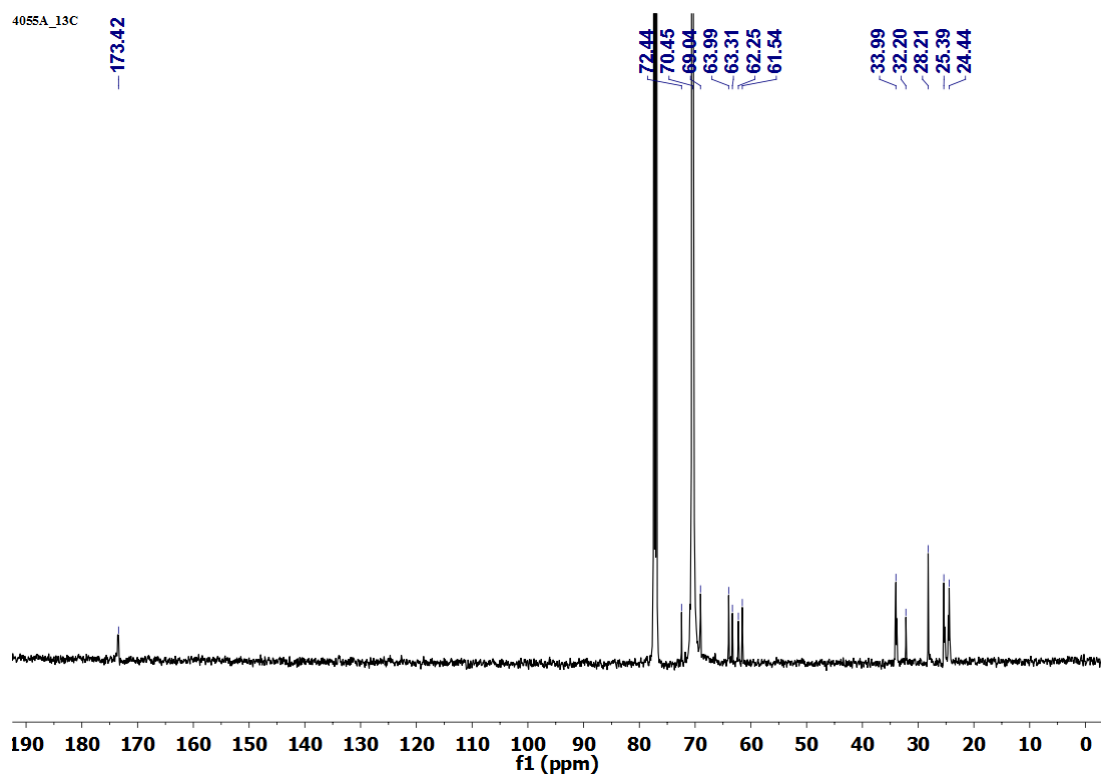


Figure S10. ^{13}C NMR of PEG-PCL-OH

1.3.3. Characterization of PEG-PCL-OMs

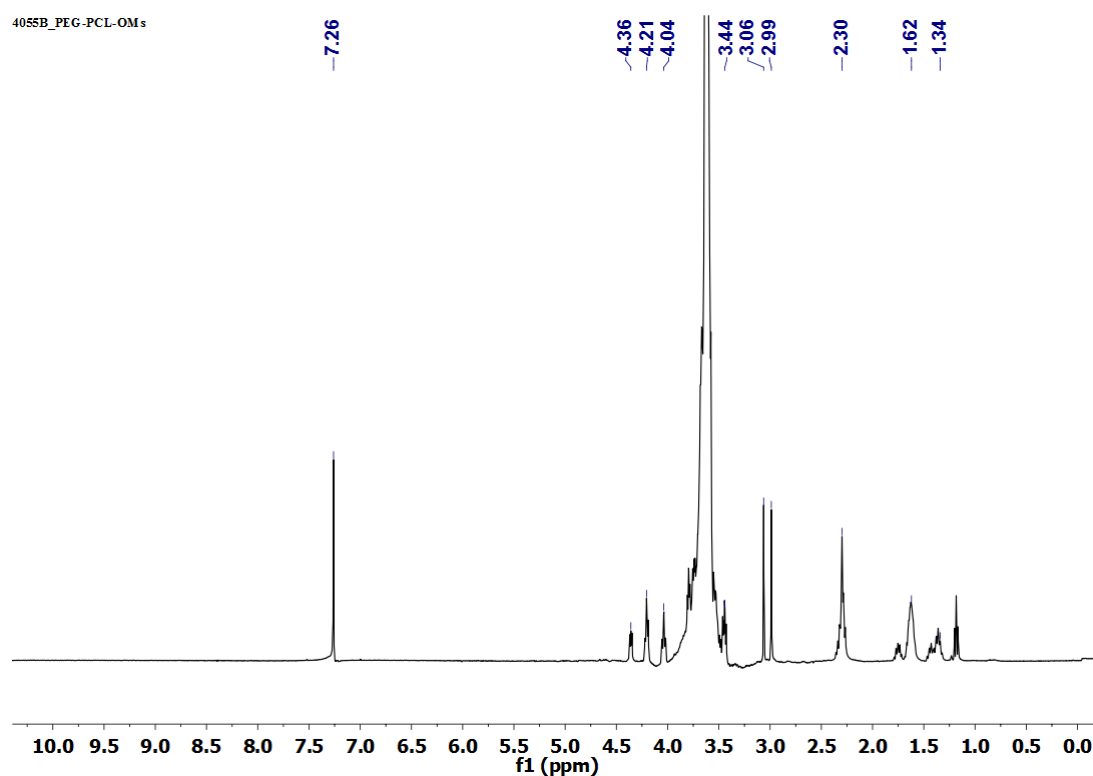


Figure S11. ^1H -NMR of PEG-PCL-OMs

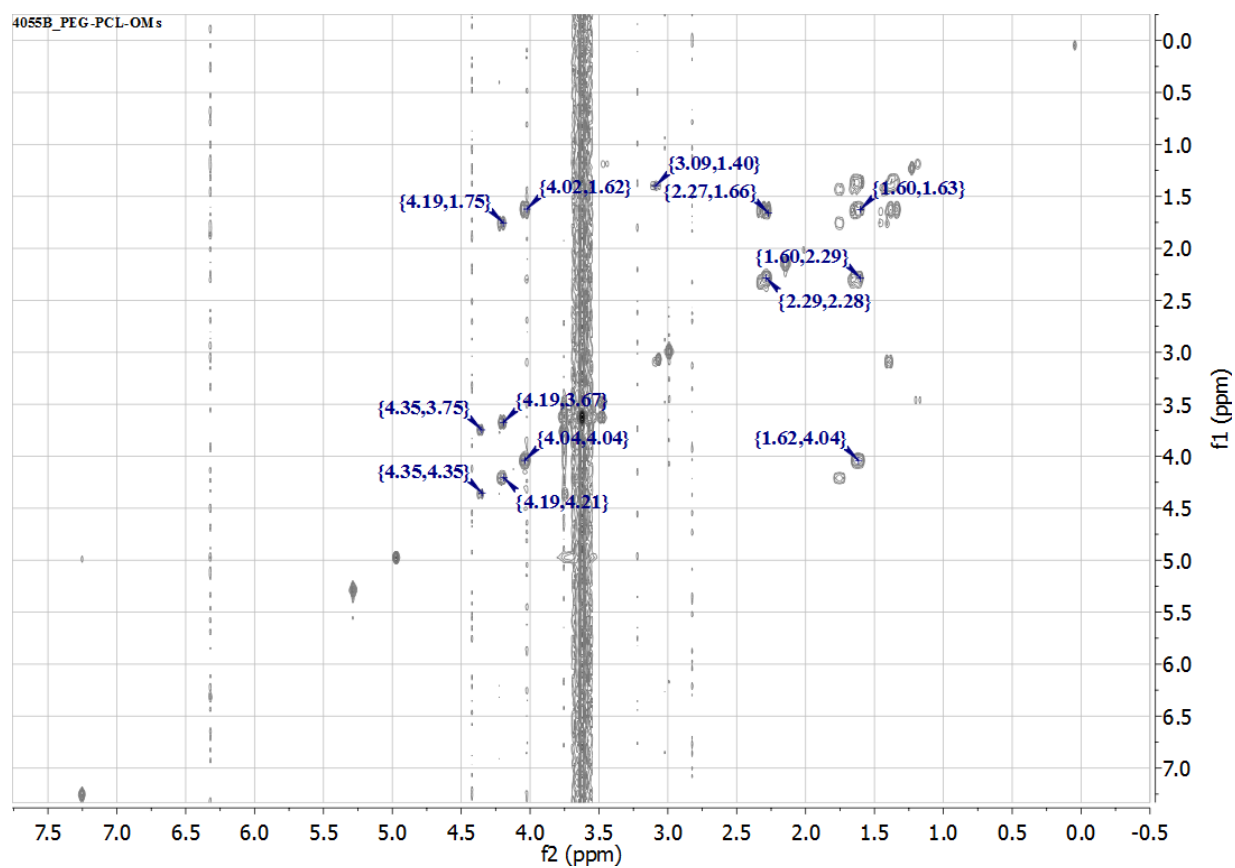
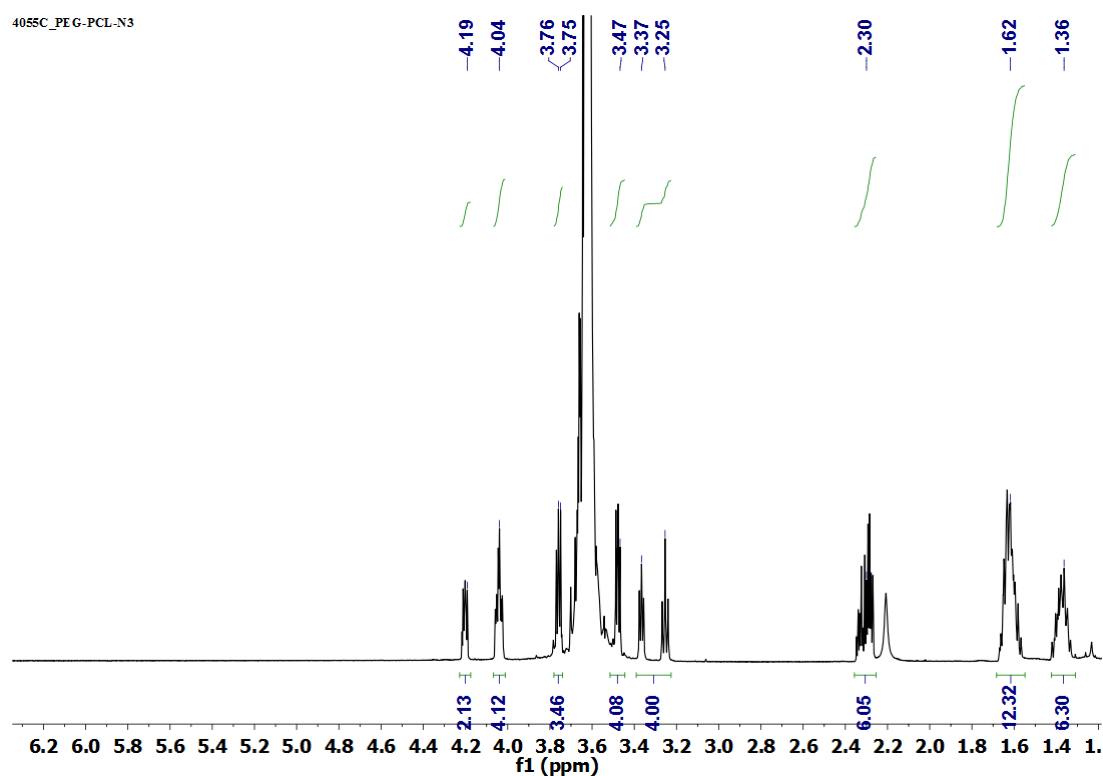
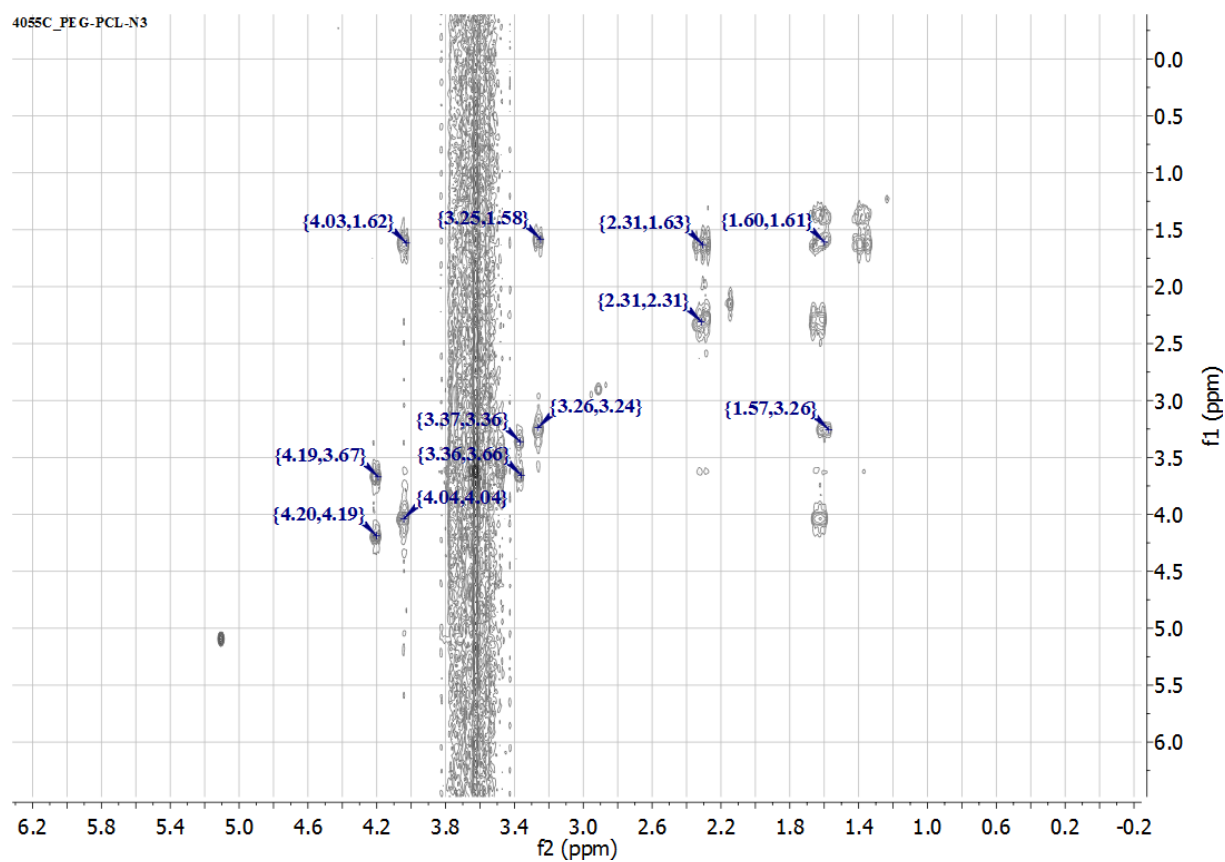


Figure S12. ^1H - ^1H correlation spectroscopy of PEG-PCL-OMs

1.3.4. Characterization of PEG-PCL-N₃Figure S13. ¹H NMR of PEG-PCL-N₃Figure S14. ¹H-¹H correlation spectroscopy of PEG-PCL-N₃

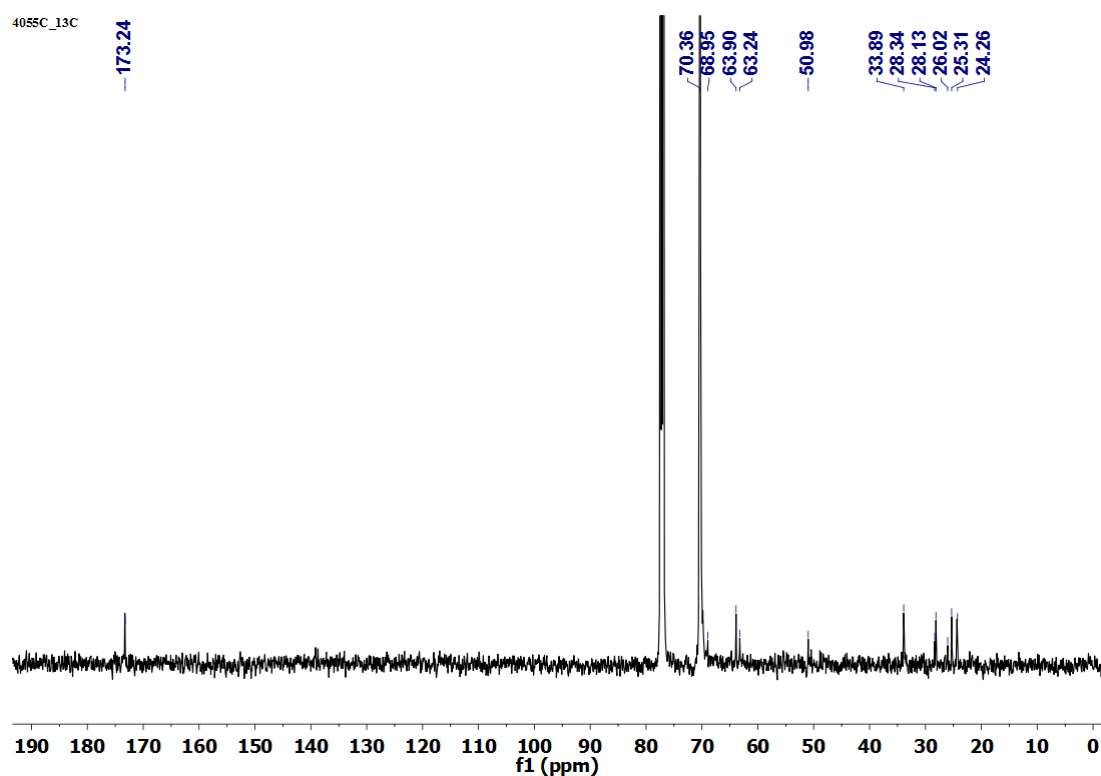


Figure S15. ^{13}C NMR of PEG-PCL- N_3

1.3.5. Characterization of PEG-PCL-DIC

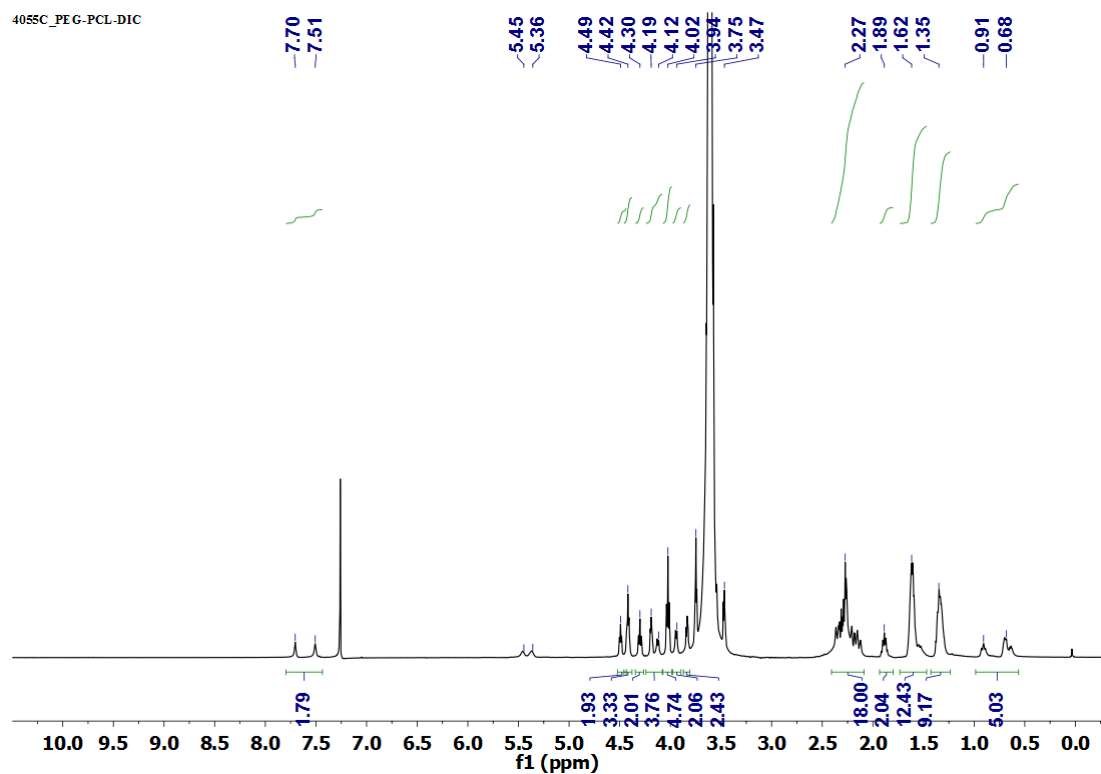


Figure S16. ^1H NMR of PEG-PCL-DIC

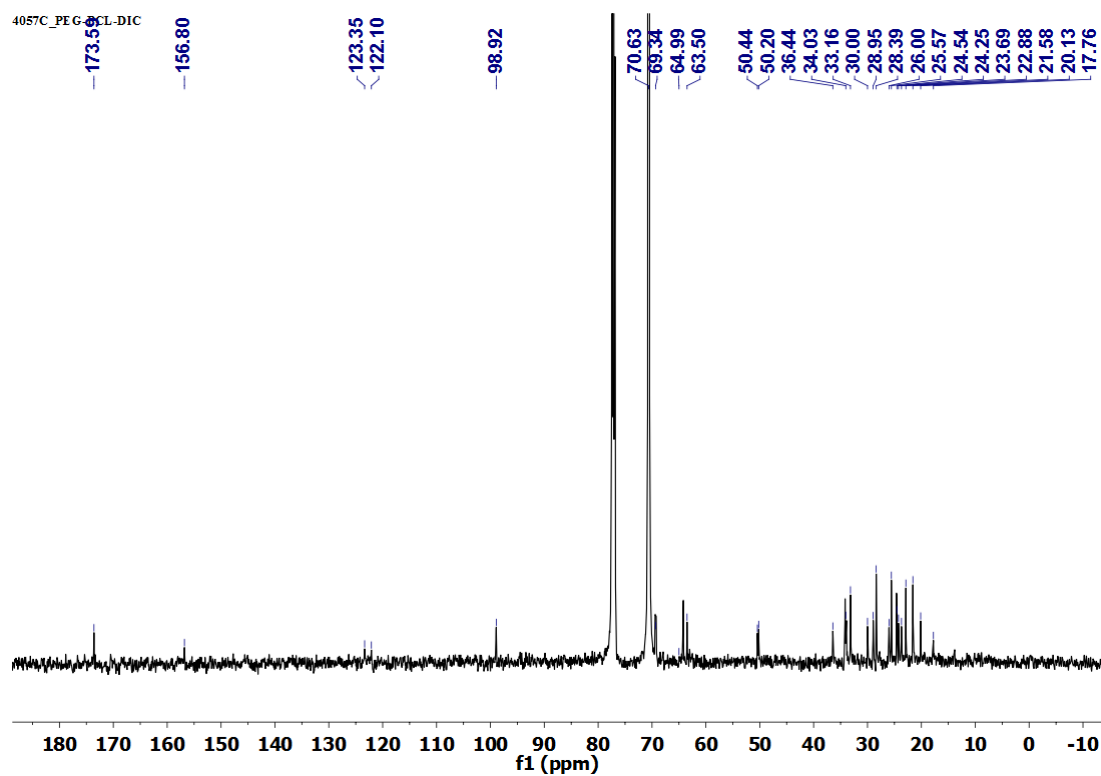


Figure S17. ^{13}C NMR of PEG-PCL-DIC

1.4. Morphology of hydrogels

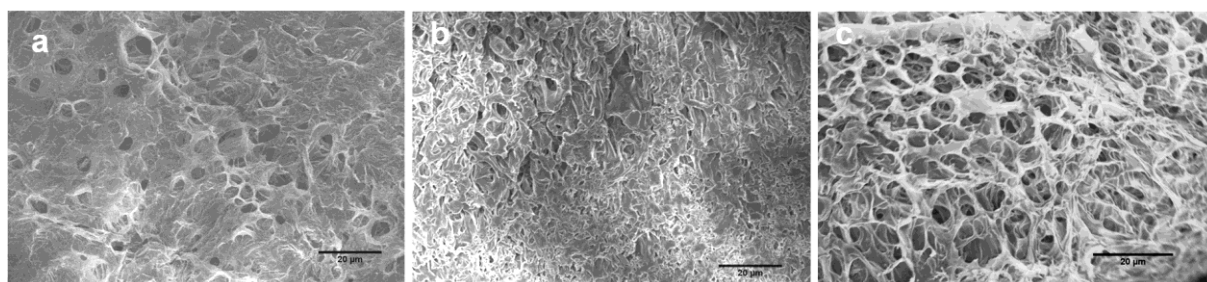


Figure S18. SEM micrographs of gel 1(a), gel 2 (b), and a cross-sectional image of gel 2 (c). The scale bar is 20 μm .

1.5. Cyto-compatibility of non-degradable dPGS gel

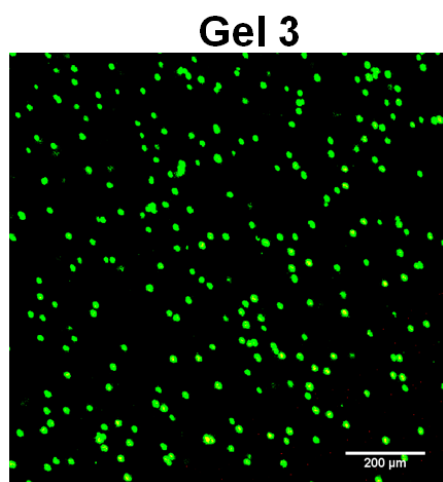


Figure S19. CLSM image showing mouse fibroblast L929 cells encapsulated in dPGS – PEG-DIC non-degradable hydrogels after 24 h culture. Cell seeding density: 20,000/ 50 μL of gel (4×10^5 cells/ml). The live cells were stained with calcein (green) and the dead cells were stained by ethidium bromide. The scale bar is 200 μm .