Supporting Information for

Isosorbide Diallyl Based Antibacterial Thiol-Ene Photocured Coatings Containing Polymerizable Fluorous Quaternary Phosphonium Salt

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Cell viability assay

The proliferation of the L929 mouse fibroblasts in the presence of was evaluated by MTT assay. Briefly, cells were seeded at a density of 10^5 cells per well into a 96-well plate. PFQPS containing sample (30mg/ml) was prepared by adding Dulbecco's modified Eagle's medium (DMEM). Then the whole plate was incubated for 24h at 37°C under CO₂. After 24 hours of incubation, 20 µL of MTT (3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyl-tetrazolium bromide) solution (5 mg/mL in PBS) was added to each well. The formed dark blue formazan crystals in the cells were dissolved in DMSO (dimethyl sulfoxide) and the absorbance at 570 nm was measured with a microplate reader. 1% phenol solution was used as a positive control sample and for the negative control sample only the cell medium was used. The results are expressed as percentage with respect to the control sample that was not treated with PFQPS. All experiments were repeated 4 times.



Figure S1: FTIR spectra of a) isosorbide and b) isosorbide diallyl ether



Figure S2: ¹H-NMR of isosorbide diallyl ether



Figure S3: FTIR spectra of a) 17F-PI and b) PFQPS

a) 17F-PI



Figure S4: ¹H-NMR of a) 17F-PI and b) PFQPS



Figure S5: TGA spectrum of PFQPS



Figure S6: FTIR spectra of the photocured resins: a) before curing and b) after curing



Figure S7: DSC spectra of the photocured resins



Figure S8: XPS spectra of the selected photocured resins and the surface elemental percentages.



Figure S9: Viable L929 fibroblast cell fraction (%) in the presence of PFQPS. The cell viability was measured via MTT assay and the experiments were performed in quadruple.