SUPPORTING INFORMATION for

Synthesis and targeting of PPP-type copolymers to breast cancer cells: Multifunctional platforms for imaging and diagnosis

Demet Goen Colak, Ioan Cianga, Dilek Odaci Demirkol, Ozge Kozgus, E. Ilker Medine,

Serhan Sakarya, Perihan Unak, Suna Timur*, and Yusuf Yagci*

*Corresponding Authors:

Prof. Y. Yagci

Department of Chemistry, Faculty of Science and Letters, Istanbul Technical University, Maslak, 34469, Istanbul, Turkey, E-mail: yusuf@itu.edu.tr

Prof. S. Timur Biochemistry Department, Faculty of Science, Ege University, Bornova, 35100, Izmir, Turkey, E-mail: suna.timur@ege.edu.tr



Spectral data for macromonomer and PPP copolymer;

Figure S1. ¹H-NMR spectral data: (a) macromonomer, and (b) PPP copolymer in CDCl₃.

Cytotoxicity of PPP



Figure S2. Time dependent cell viability of MCF7 cells. 5x105 cells/mL in the presence of PPP (0.05 mg/mL in the medium) and PBS (pH 7.4, 50 mM) as a function of time by typical MTT assay. Error bars represent the standard deviation of three measurements].

Fluorescence spectra of PPP after conjugation of Anti-MTDH, Lec and

Anti- MTDH-Lec



Figure S3. Fluorescence spectra of PPP after bioconjugation. Anti-MTDH, Lec and Anti-MTDH-Lec conjugation (in PBS pH 7.4), the excitation wavelength is 280 nm.