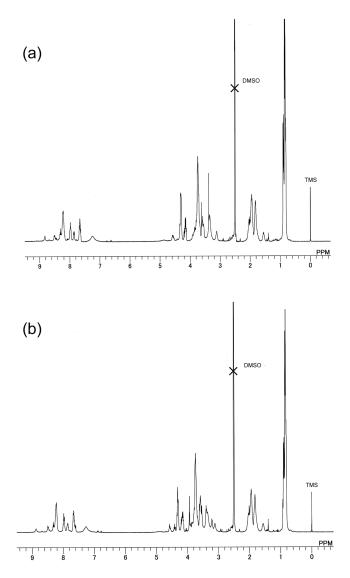
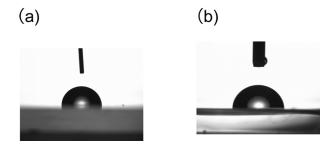
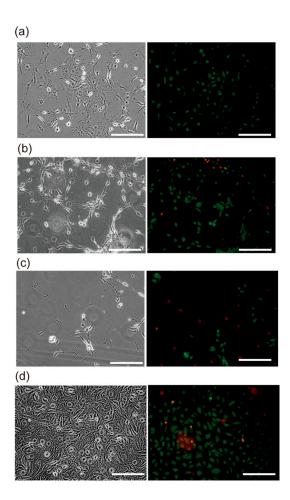
## **Supporting Information**



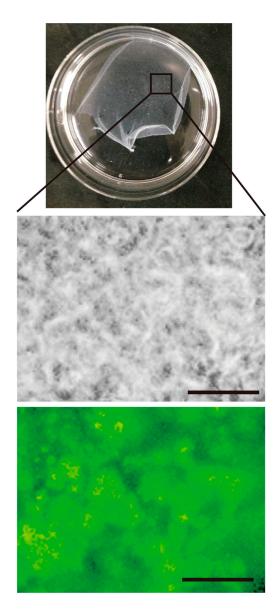
**Figure S1.** <sup>1</sup>H-NMR spectra of the RGDS-ELP (**a**) and RGDS-deg-ELP (**b**) in  $d_6$ -DMSO (TMS) at 20 °C.



**Figure S2.** Water contact angles of the **(a)** RGDS-ELP-covered PSt-dish and **(b)** RGDS-deg-ELP-covered PSt-dish.



**Figure S3.** The live/dead cell assay of NIH/3T3 cells (1.0 × 10<sup>4</sup> cells cm<sup>-2</sup>) cultured for 24 h on RGDS-deg-ELP-covered PSt-dish (**a**), RGDS-ELP-covered PSt-dish (**b**), bare PSt-dish (**c**) and tissue culture dish (Biocoat<sup>TM</sup> Fibronectin Cellware) (**d**). Left pictures: phase-contarast images, right pictures: merged fluorescence images. Calcein stain (green) identifies live cell, while EthD stain (red) identifies dead cells. Scale bar: 200 μm.



**Figure S4.** Phase-contrast and fluorescence images of the cell-sheet prepared on RGDS-ELP-covered PSt-dish. Calcein stain in fluorescence image (bottom) identifies live cell. Scale bar:  $50 \mu m$ .