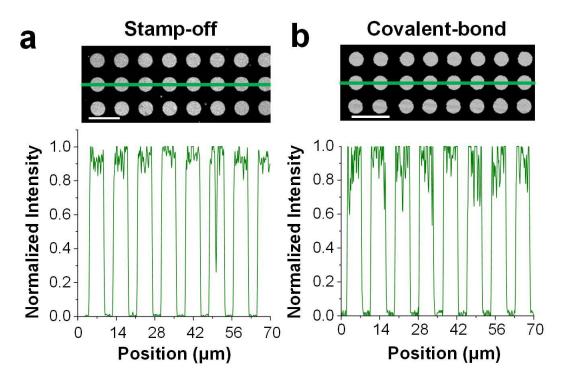
## Protein-Substrate Adhesion in Microcontact Printing Regulates Cell Behaviors

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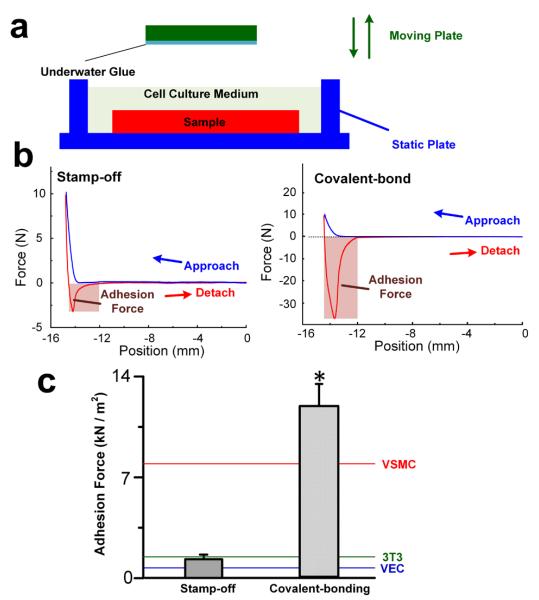
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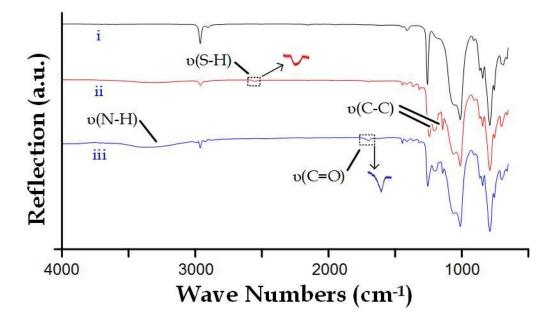


## **Supplemental Figures**

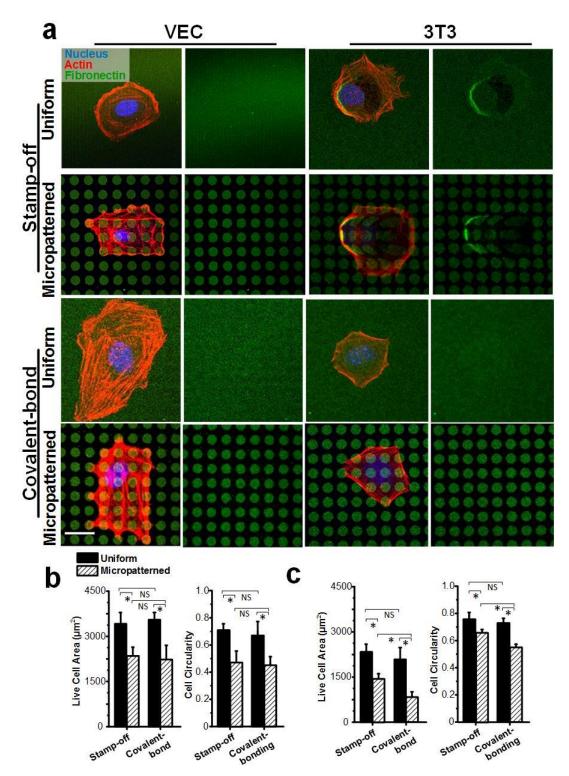
Figure S1. Quality of the  $\mu$ CP created patterns. High contrasts of the fluorescent brightness among the patterned areas and uncoated areas indicate the good quality of the created patterns for (a) stamp-off and (b) covalent-bond methods. Scale bar: 10  $\mu$ m.



**Figure S2**. Measurement of the adhesion force between the ECM protein and the substrates. (a) Set-up of the measurement. (b) Representative force-displacement curves for the measurements of the substrates coated by the stamp-off method and the covalent-bond method. (c) The adhesion force of protein coating by the stamp-off method and the covalent-bond method.



**Figure S3.** ATR-FTIR spectra at the key micropatterning steps. v(S-H) and v(C-C) adsorptions indicate the conjugation of MTPMS; v(N-H) and v(C=O) adsorption indicates the conjugation of GMBS.



**Figure S4.** Micrographs (a) and morphological statistics of (b)VECs and (c)3T3 cells seeded on the substrates coated by the stamp-off method and the covalent-bond method. NS stands for no significance. Scale bar:  $20 \mu m$ .

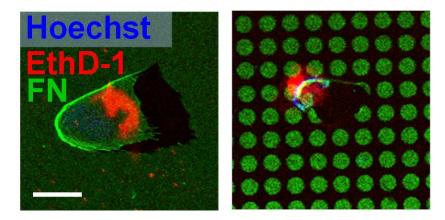
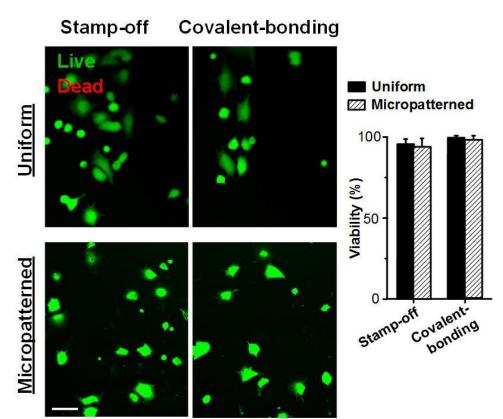


Figure S5. Correlations between the ECM protein delamination of the cell death for 3T3 cells. The red colour indicates the EthD-1 dye entered a dead cell. Scale bar:  $20 \mu m$ .



**Figure S6.** VMSCs viability was significantly improved (>90%) after treated with 50  $\mu$ M blebbistatin on the stamp-off and covalent-bonding surfaces. Scale bar: 50  $\mu$ m.