

Electronic Supplementary Material (ESI) for Materials Chemistry Frontiers.
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Supporting information

Dynamic Display of Cell Targeting Motifs via Natural Glycopeptide Recognition for Cancer Cell Isolation

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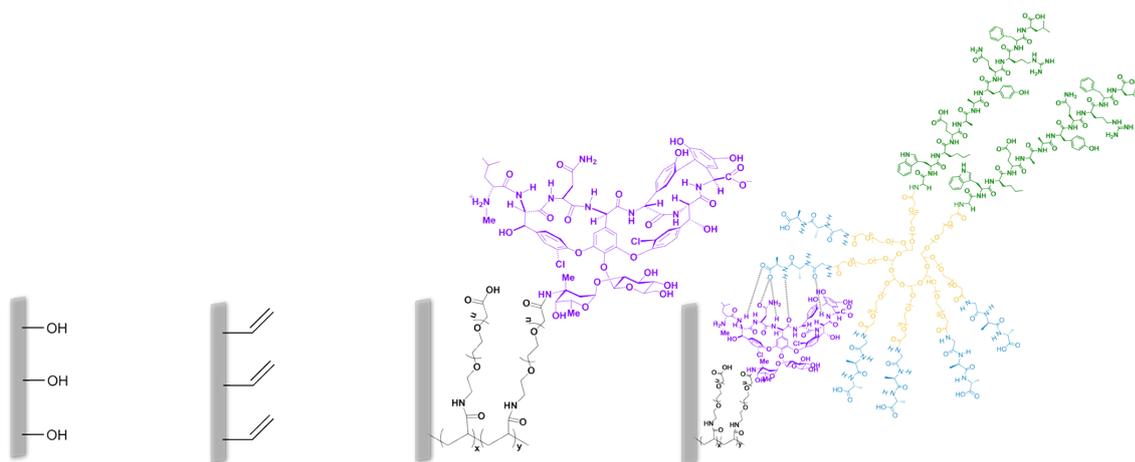


Figure S1. Schematic procedure for the preparation of Van brush grafted quartz substrate binding a target peptide.

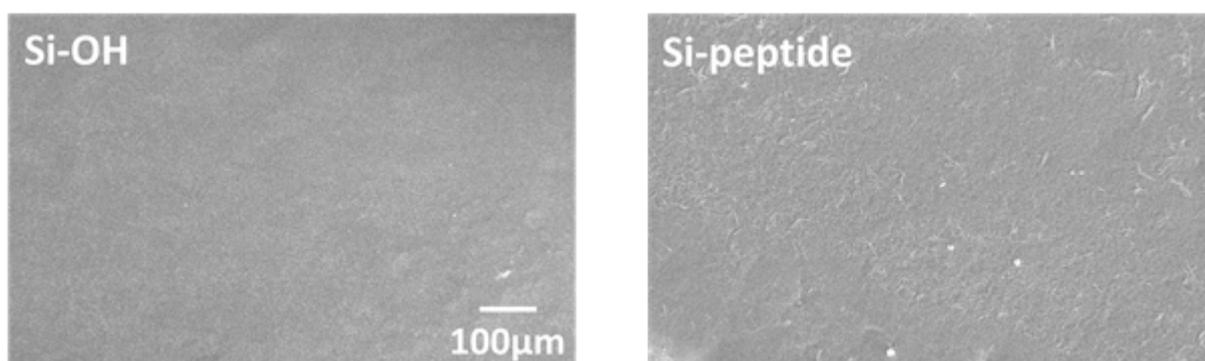


Figure S2. SEM images of different modified active surfaces.

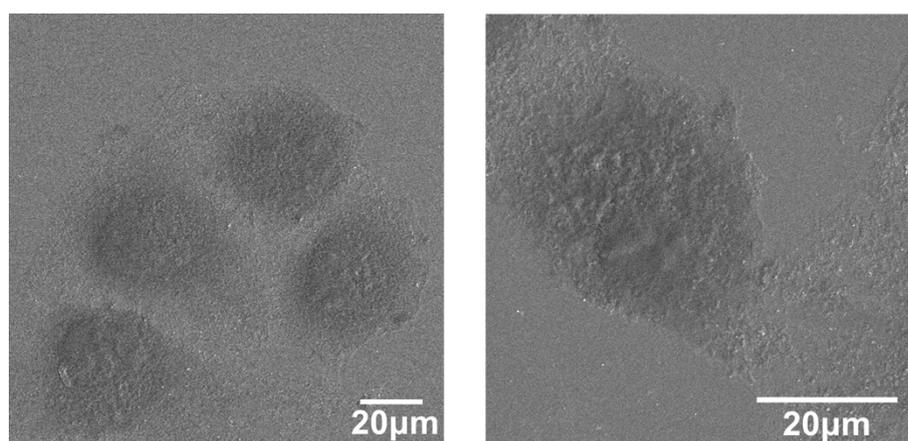


Figure S3. SEM image of MCF-7 cell adhered on the active surface.

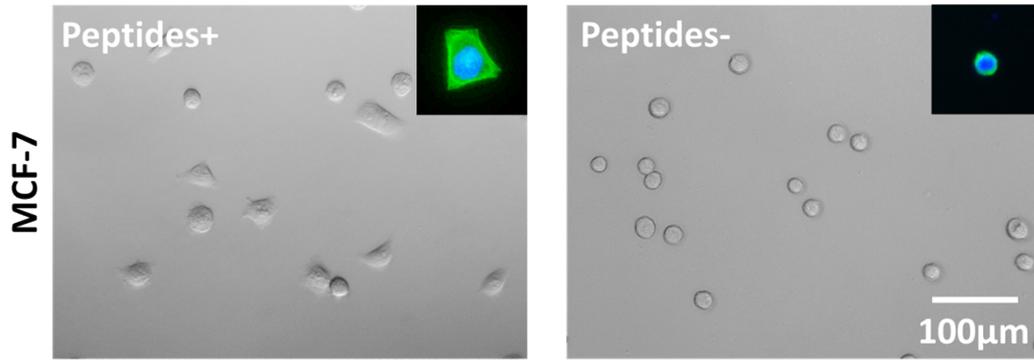


Figure S4. Representative micrographs and fluorescence staining images of PBS-incubated MCF-7 cells on different active surfaces.

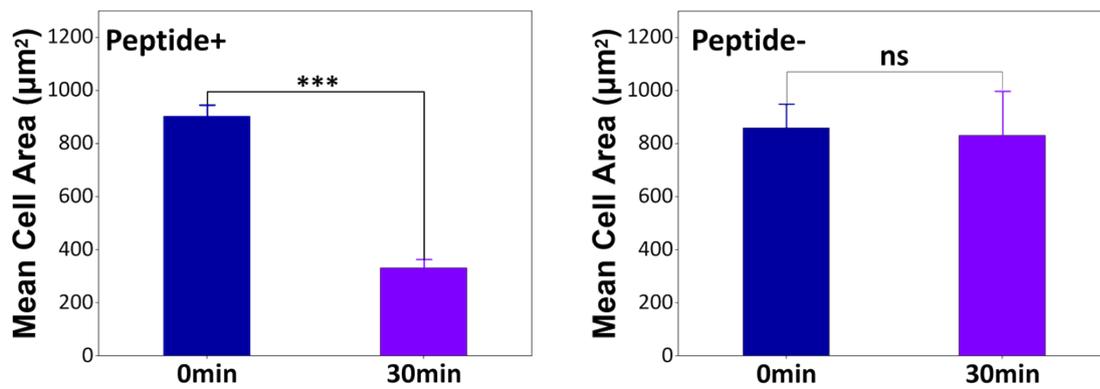


Figure S5. Mean cell area variations at different times after the addition of GAA to peptide+ and peptide-surfaces.

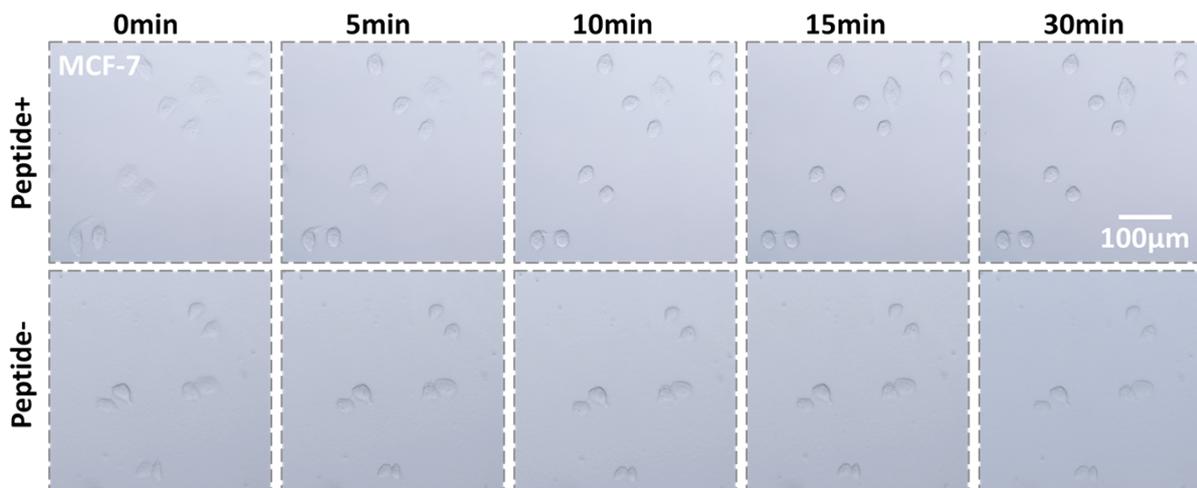


Figure S6. Microscopic images of cell morphology on peptide+ and peptide- surfaces with time after the addition of free AA (10mM).

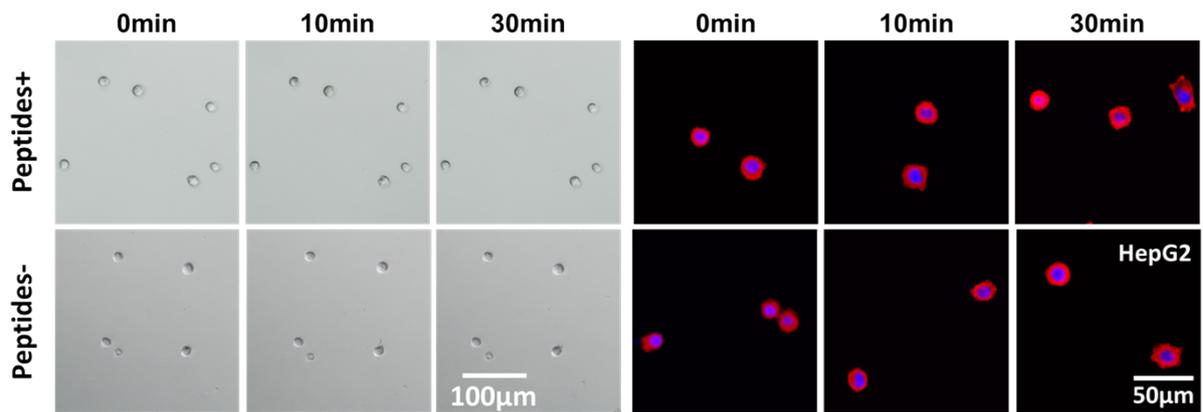


Figure S7. Real-time morphological changes of HepG2 cells on peptide+ and peptide- surfaces after the addition of free AA (10 mM). F-actin and nuclear were prestained by phalloidine (red) and DAPI (blue), respectively.

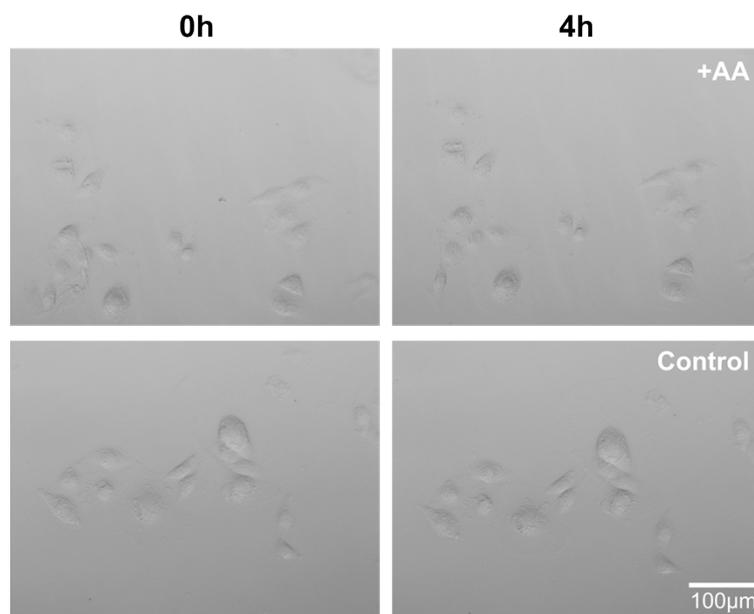


Figure S8. Cell morphology of MCF-7 co-cultured with free AA for 4h.

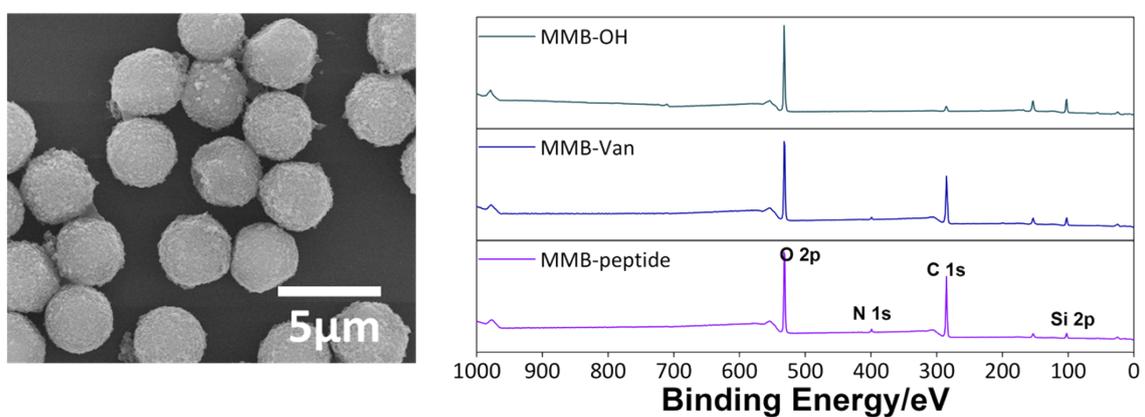


Figure S9. The SEM image and XPS spectrum of MMB-peptide.

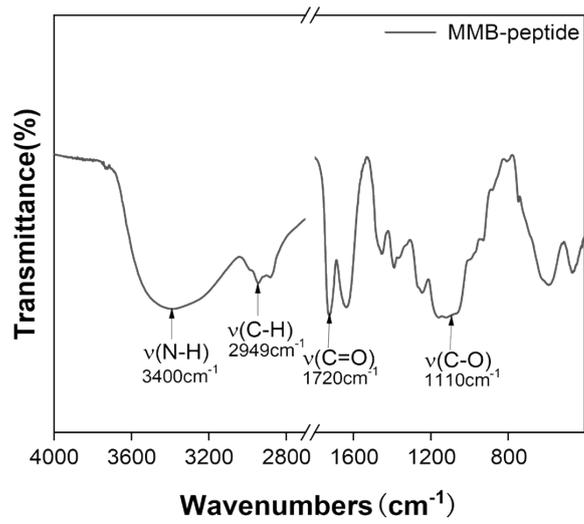


Figure S10. FT-IR spectrum of MMB-peptide.

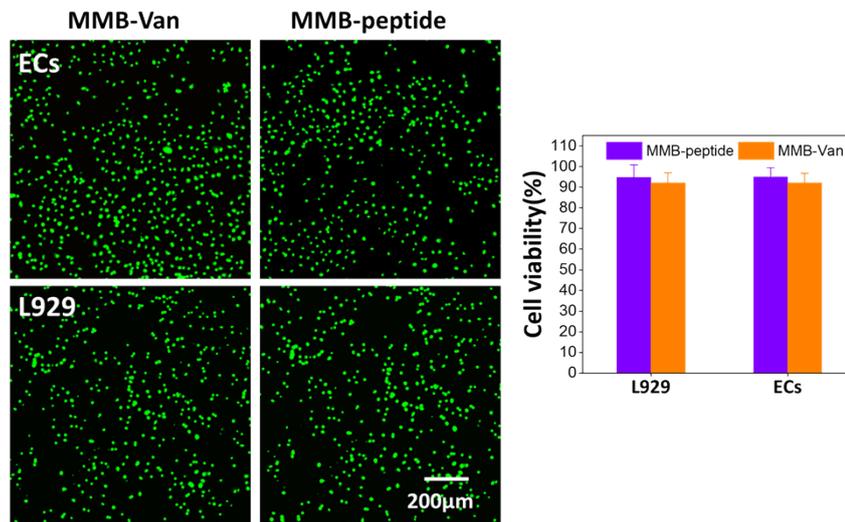


Figure S11. Live/dead staining and cytotoxicity of L929 and ECs co-cultured with different modified MMB.

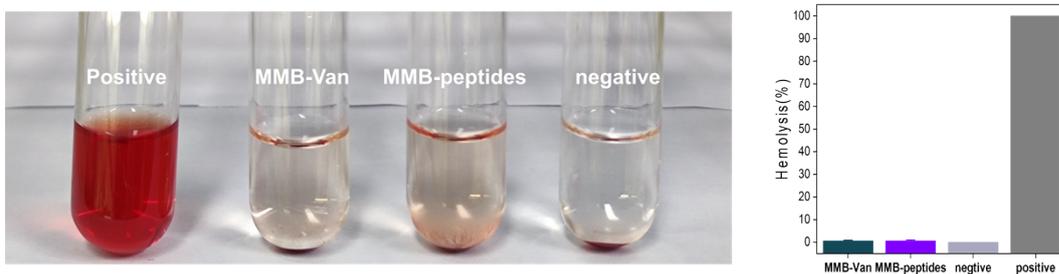


Figure S12. Hemolysis images of MMB after modification and their hemolysis rates.