VAPG peptide density gradient and its influence on selective adhesion and directional migration of smooth muscle cells

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Selective adhesion and directional migration of smooth cells (SMCs) over fibroblast cells (FIBs) plays a significant role in the wound healing after blood-contacting implants, in particular for the scar formation. In this study, a uniform cell-resistant layer of polyethylene glycol (PEG) was firstly immobilized on the glass slides surface. Then a density gradient of alkynyl functionalized Val-Ala-Pro-Gly (VAPG) peptide was immobilized onto the PEG layer via click chemistry.



**Figure 1.** Schematic illustration to show the structure of the SMCs selective peptide (VAPG) density gradient on a uniform PEG brush layer. VAPG peptides were grafted onto cell-resistant layer through click chemistry.



**Figure 4.** Migration trajectories SMCs and FIBs on PEG, 1 mm, 5 mm, 9 mm on VAPG density gradient. The numbers represent percents of cells migrated toward +X direction.





**Figure 2.** Numbers of SMCs and FIBs being cultured for 8 h on TCPS and different positions of VAPG density gradient.





**Figure 5.** Migration rates of SMCs and FIBs on TCPS, PEG and 1 mm, 5 mm and 9 mm on VAPG density gradient.

## Conclusion

The gradient surface selectively enhanced the SMCs adhesion, preferential orientation and directional migration. The migration rate of SMCs was enhanced to 1.5-folds compared to that on TCPS. The gradient significantly weakened the adhesion of FIBs to 33% of that on TCPS, but had no obvious impact on the migration rate and directionality of adherent SMCs. The success of the gradient relies on the appropriate interplay between the cell-resistant layer and the cellspecific ligands, enabling the selective guidance of adhesion and migration of SMCs.

**Figure 3.** CLSM images of SMCs (a-d) and FIBs (e-h) on VAPG density gradient at (a,e) PEG, (b,f) 1 mm, (c,g) 5 mm and (d,h) 9 mm positions.

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## Reference

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