

Electronic Supplementary Information

Study of endothelial cell apoptosis using fluorescence resonance energy transfer (FRET) biosensor cell line with hemodynamic microfluidic chip system

J. Q. Yu^a, L. K. Chin^a, X. F. Liu^b, A. Q. Liu^a and K. Q. Luo^{*b}

^a*School of Electrical and Electronic Engineering,
Nanyang Technological University, Singapore 639798*

^b*School of Chemical and Biomedical Engineering,
Nanyang Technological University, Singapore 637457*

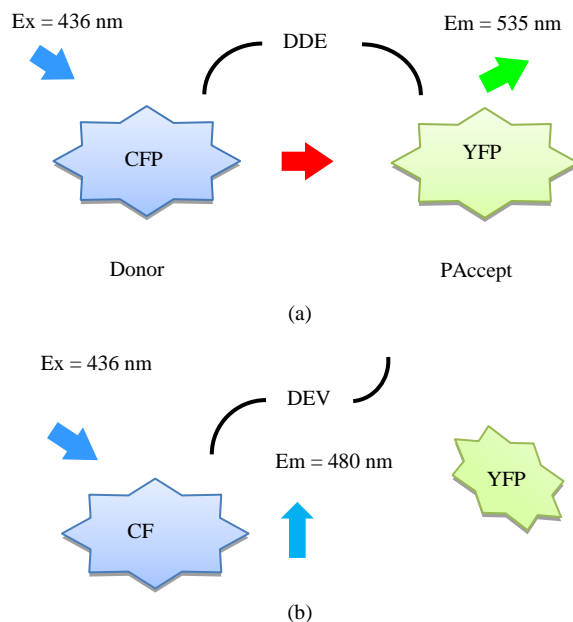


Fig. S1 Principle of a living cell biosensor in detecting caspase-3 activation based on FRET effect. (a) Living HUVEC-C3 with a 535-nm green fluorescent emission, and (b) dead HUVEC-C3 with cleaved linker peptide and a 480-nm blue fluorescent emission.

([†] Tel: +65 6790-4257; Fax: +65 6791-1761; Email: kluo@ntu.edu.sg)

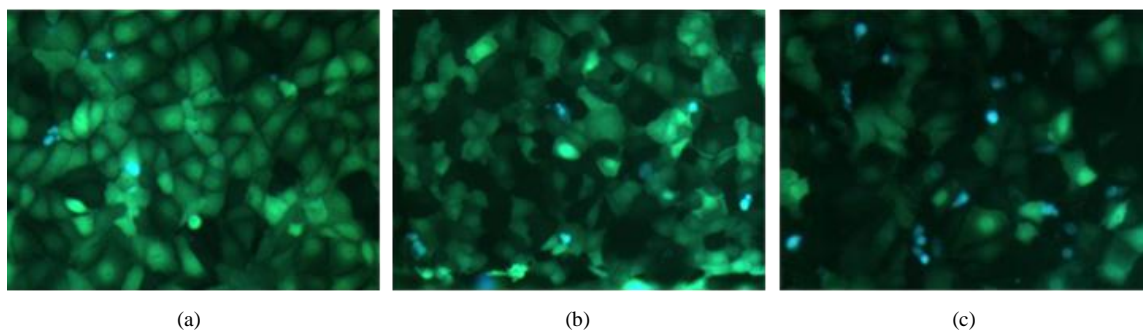


Fig. S2 FRET changes in response to caspase-3 activation with different glucose concentrations (a) 5 mM (b) 10 mM (c) 20 mM under shear stress 30 dyne cm^{-2} 2 h plus shear stress 15 dyne cm^{-2} 10 h.

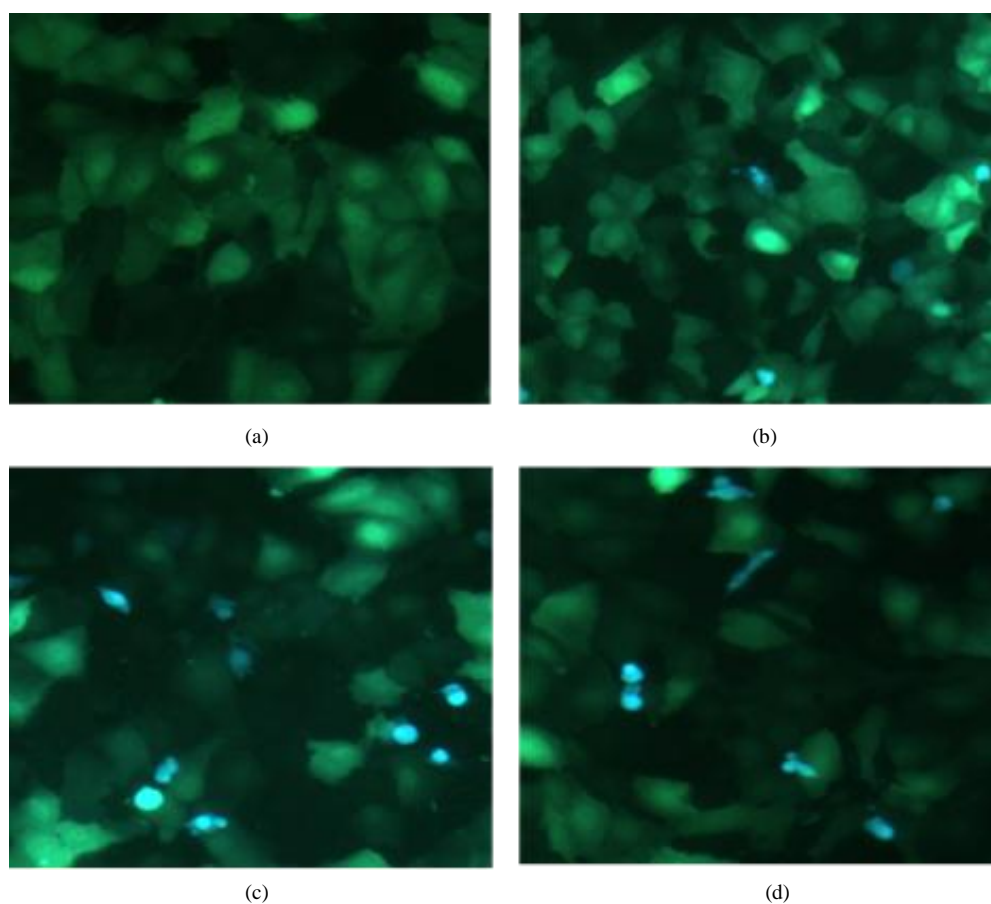


Fig. S3 FRET changes in response to caspase-3 activation under different conditions (a) 5 mM glucose under shear stress of 15 dyne cm^{-2} for 12 h, (b) 20 mM glucose under shear stress of 15 dyne cm^{-2} for 12 h (c) 20 mM glucose under shear stress of 30 dyne cm^{-2} for 1 h plus shear stress of 15 dyne cm^{-2} for 11 h, (d) 20 mM glucose under shear stress of 30 dyne cm^{-2} for 2 h plus shear stress of 15 dyne cm^{-2} for 12 h.

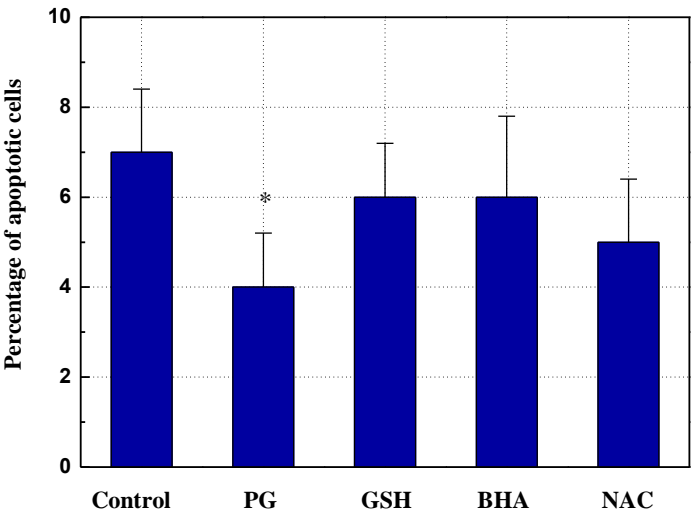


Fig. S4 Effect of antioxidant drugs on high glucose and shear stress induced cell apoptosis.