

Drug-eluding microarrays for cell-based screening of chemical-induced apoptosis

Cheong Hoon Kwon^{a,b}, Ian Wheeldon^{a,b,c}, Nezamoddin N. Kachouie^{a,b}, Seung Hwan Lee^{a,b}, Hojae Bae^{a,b}, Shilpa Sant^{a,b,c}, Junji Fukuda^d, Jeong Won Kang^{e,*} and Ali Khademhosseini^{a,b,c,*}

^a Center for Biomedical Engineering, Department of Medicine, Brigham and Women's Hospital, Harvard Medical School, Boston, MA 02115

^b Harvard-MIT Division of Health Sciences and Technology, Massachusetts Institute of Technology, Cambridge, MA 02139

^c Wyss Institute for Biologically Inspired Engineering, Harvard University, Boston, MA, 02115

^d Graduate School of Pure and Applied Sciences, University of Tsukuba, Tsukuba, Japan 305-8573

^e Department of Chemical and Biological Engineering, Korea University, 5-Ga Anam-Dong, Sungbuk-Ku, Seoul, South Korea 136-701

* Corresponding authors: Ali Khademhosseini, alik@rics.bwh.harvard.edu, Jeong Won Kang, jwkang@korea.ac.kr

Supporting Information

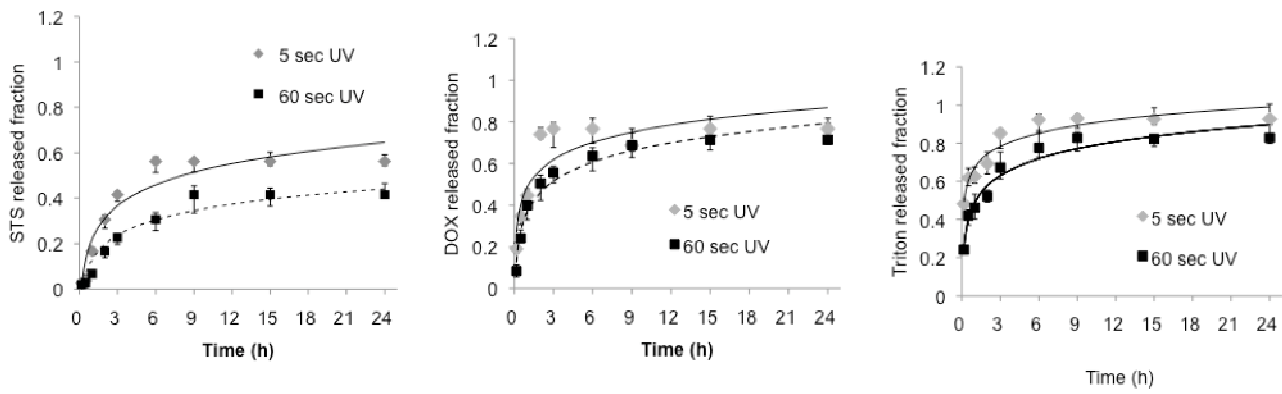


Figure 1. Chemical release from PEGDA hydrogels. Release profile of 10 μM STS (A), 10 μM DOX, and 0.1 % Triton X-100 (C), over a 24 hour period with 5 and 60 seconds of UV exposure. The release of each compound was measured from 1 μL gels placed in 100 μL of PBS solution. Chemical release was monitored by UV-Vis spectrometry at 495 nm for doxorubicin, 292 nm for staurosporine, and 280 nm for Triton X-100.

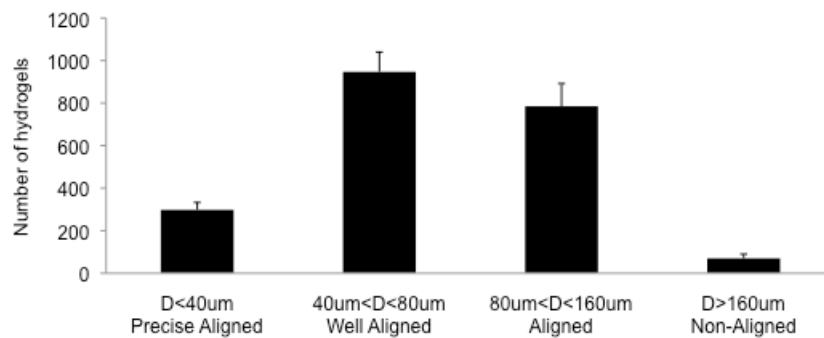


Figure 2. Device alignment accuracy. The number of hydrogels with a distance between the center point of the hydrogel and the center of the aligned microwell that is less than 40 μm , between 40 and 80 μm , between 80 and 160 μm , and greater than 160 μm . Hydrogels and microwells are considered

misaligned if the distance between the center of each is greater than 160 μm . The average alignment accuracy is 96.7% (n = 3).

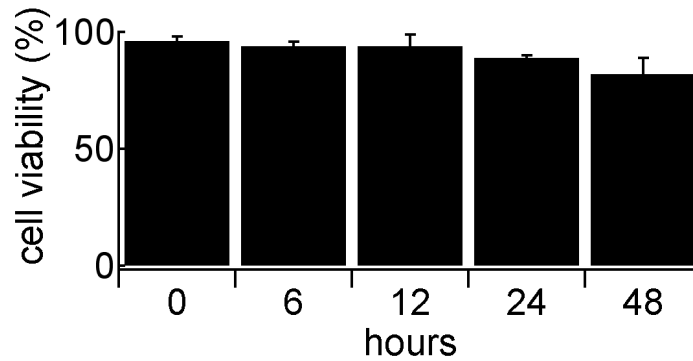


Figure 3. HEPG2 viability in sealed microwell cultures.

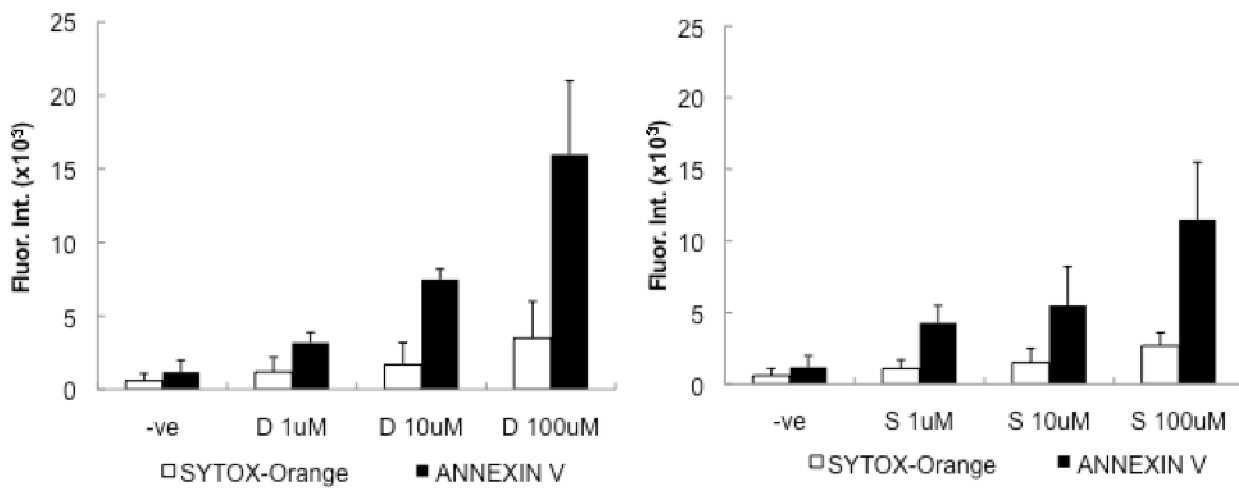


Figure 4. DOX and STS apoptotic assays in 96-well plate format. The fluorescence intensity of annexin V-APC and SYTOX Orange of MCF-7 cells exposed to 0, 1, 10 and 100 μM of DOX and STS for 12 hours.

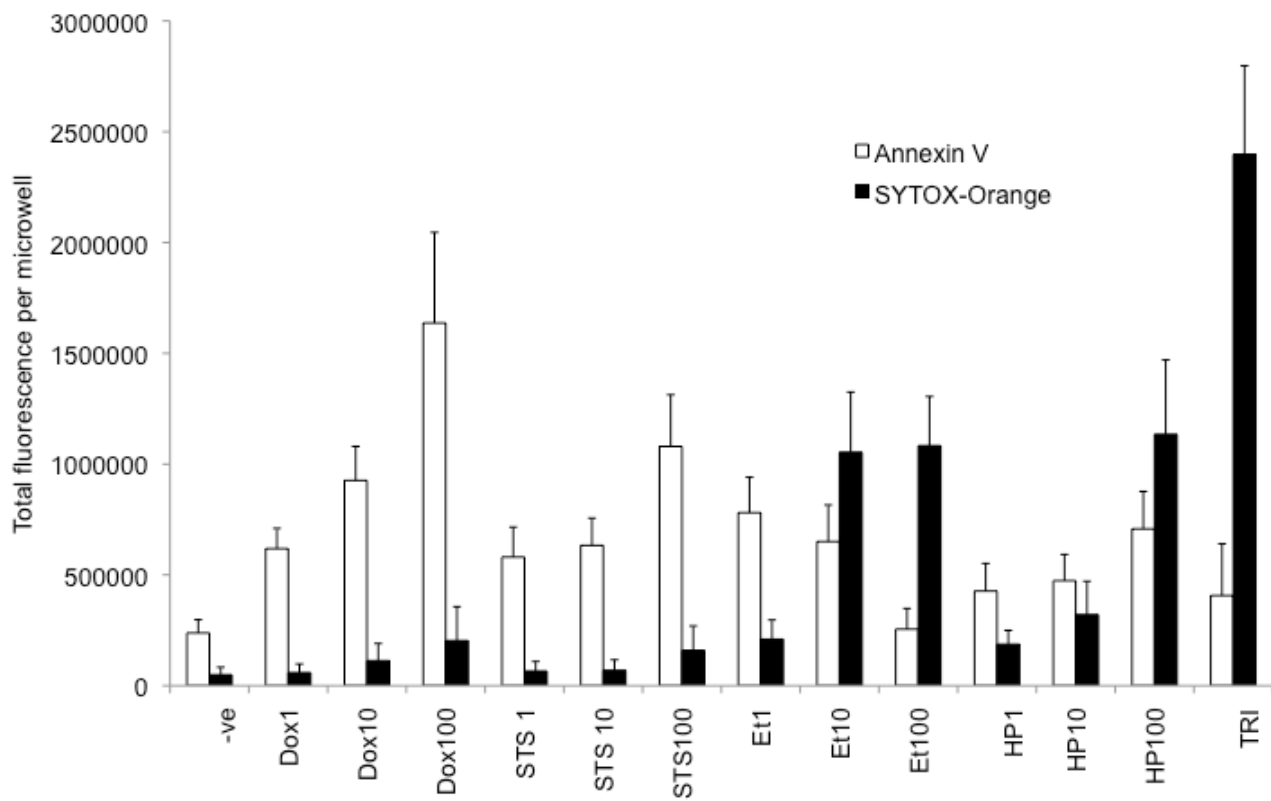


Figure 5. Annexin V-APC and STOX Orange fluorescent intensity for DOX, STS, ethanol, and hydrogen peroxide.