Supporting Information

Biocompatible Polyethylene Glycol/Alginate Composite Hydrogel with Significant Reactive Oxygen Species Consumption for Promoting Wound Healing

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1. Characterization of purified poly (ethylene glycol) diacrylate.

Poly (ethylene glycol) diacrylate was purified and characterized by Agilent AV 400. ¹H NMR (400 MHz, CDCl₃) δ 6.45 – 6.37 (m, 2H), 6.18 – 6.09 (m, 2H), 5.86 – 5.79 (m, 2H), 4.34 – 4.24 (m, 4H), 3.72 (dt, *J* = 11.6, 4.5 Hz, 4H), 3.62 (d, *J* = 0.6 Hz, 72H).

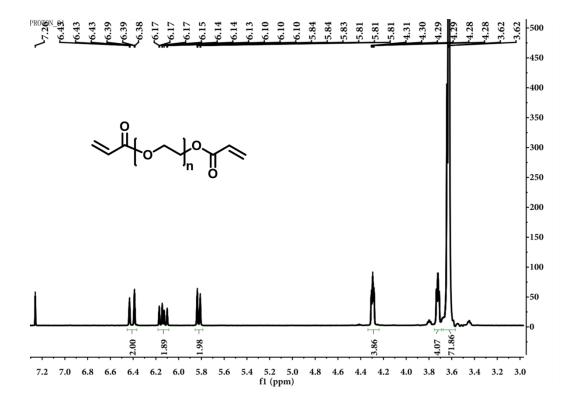


Figure S1. ¹H NMR spectrum of purified poly (ethylene glycol) diacrylate in CDCl₃.

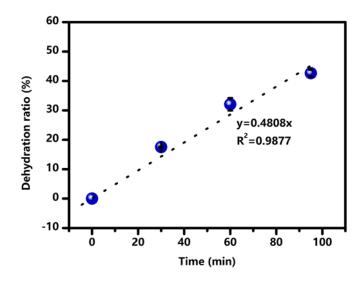


Figure S2. Simulation of *hdg*-PEGDA dehydration in the first 95 minutes.

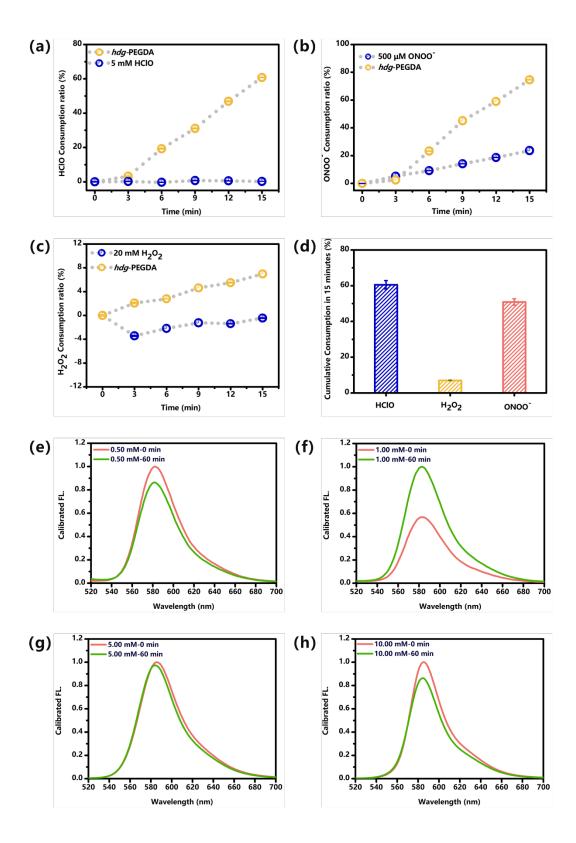


Figure S3. (a-c) *hdg*-PEGDA consumption on different types of reactive oxygen species (ROS).(d) The cumulative consumption in 15 minutes of hydrogel *hdg*-PEGDA on different types of

ROS. (e-h) The self-consumption of different concentrations of hypochlorous acid within 60

minutes.

HepG2 cells were stained with TRICT (Cytoskeleton–Red), Hoechst 33342 (Nucleus–Blue)

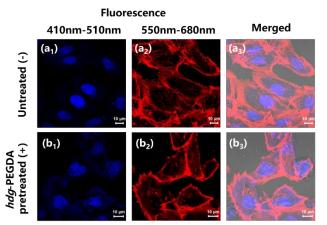


Figure S4. Nucleus and cytoskeleton morphology of HepG2 cells after hdg-PEGDA treatment

 (b_1-b_3) and untreated (a_1-a_3) .

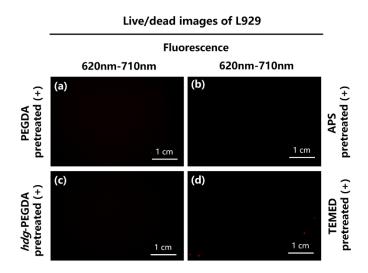


Figure S5. (a-d) Dead images of L929 cells treated with different conditions.

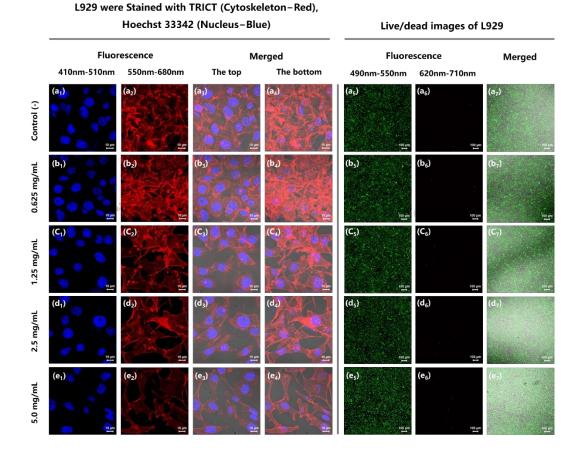


Figure S6. (a1-a4), (b1-b4), (c1-c4), (d1-d4) and (e1-e4): nucleus and cytoskeleton morphology of L929 cells after treated with different concentrations of SA degradation products. (a5-a7), (b5-b7), (c5-c7), (d5-d7) and (e5-e7): Live/dead images of L929 cells after co-cultured with SA degradation products for 24 hours.

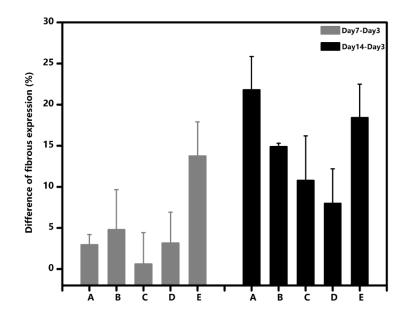


Figure S7. Analysis of the expression percentage of collagen fiber in Masson staining experiment.