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Supporting Information

Skin-Like Wound Dressings with On-demand Administration Based on *in Situ* Peptide Self-Assembly for Skin Regeneration

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Fig. S1. a) The chemical structure and MALDI-TOF-MS spectrum of GPLK. b) HPLC spectrum of GPLK.



Fig. S2. a) The chemical structure and MALDI-TOF-MS spectrum of GPK. b) HPLC spectrum of GPK.



Fig. S3. a) The chemical structure and MALDI-TOF-MS spectrum of GLK. b) HPLC spectrum of GLK.









Fig. S4. The SF-GPLK film was preserved in atmospheric moisture.



Fig. S5. TEM image of SF-GPK immersed in Tris·HCl buffer solution (pH 7.4) the addition of gelatinase for 6 h. Scale bars, 0.1μm.



Fig. S6. TEM image of SF-GLK immersed in Tris·HCl buffer solution (pH 7.4) the addition of gelatinase for 6 h. Scale bars, 0.2μm.



Fig. S7. The chemical structure and MALDI-TOF-MS spectrum of LK.



Fig. S8. CACs for LK in solution.



Fig. S9. TEM image of LK (200 μ M). Scale bars, 0.2 μ m.









Fig. S10. The picture of bacteria counting colony-forming units treated by SF-GPLK, SF-GPLK, SF-GPLK and SF group.



Fig. S11. S. aureus quantitative analysis of live/dead staining in Fig 3d.



Fig. S12. Live/dead staining of L929 and HUVEC cells after incubation in PBS for 6 h.



Fig. S13. H&E staining on day 15 of the newly regenerated skin tissues in uninfected wound. Scale bar: 250 μ m. (Black arrows: epidermal scaly skin, yellow arrows: hair follicle, red arrows: inflammatory cells.)



Fig. S14. Body weight changes of uninfected mice during the treatment.



Fig. S15. H&E staining on day 15 of the newly regenerated skin tissues in infected wound. Scale bar: $250 \ \mu m$. (Black arrows: epidermal scaly skin, yellow arrows: hair follicle, red arrows: inflammatory cells, green arrows: new capillary, purple arrows: cornification).



Fig. S16. The picture of colony-forming units of bacteria from mice tissue on day 6 treated by SF-GPLK+EGF, SF+EGF, 3M Tegaderm and PBS group.



Fig. S17. Body weight changes of infected mice during the treatment.