Identification and rational design of a novel antibacterial peptide dermaseptin-AC from the skin secretion of the red-eyed tree frog *Agalychnis callidryas*

Supplement materials



Figure S1 Inhibitory effects of DRP-AC4, DRP-AC4a and DRP-AC4b against (a) *S. aureus*, (b) *E. coli*, (c) *C. albicans*, (d) *P. aeruginosa*, (e) *E. faecalis*, (f) *K. pneumoniae* and (g) MRSA in a range of concentrations from 512 μM to 1 μM. Data represent means ± SEM.



MBICs against biofilm-forming S. aureus

Figure S2 Inhibition effects of DRP-AC4 (red), DRP-AC4a (green) and DRP-AC4b (blue) against the biofilm formed by *S. aureus*. Data represent means ± SEM.



Figure S3 Assessment of resistant induction of DRP-AC4 (red), DRP-AC4a (green) and DRP-AC4b (blue) in *S. aureus* after 16 passages. The 1/2*MIC bacterial suspension was further cultured after antibacterial assay. The vertical axis represented MIC data and the horizontal axis represented the number of passages.



Figure S4 Reverse-phase HPLC chromatogram (a) and full scan mass spectrum (b) of purified DRP-AC4. The acetonitrile gradient is indicated by solid line. Multiple charged ions: [M+2H]²⁺, [M+3H]³⁺ and [M+4H]⁴⁺.



Figure S5 Reverse-phase HPLC chromatogram (a) and full scan mass spectrum (b) of purified DRP-AC4a. The acetonitrile gradient is indicated by solid line. Multiple charged ions: [M+2H]²⁺, [M+3H]³⁺ and [M+4H]⁴⁺.



Figure S6 Reverse-phase HPLC chromatogram (a) and full scan mass spectrum (b) of purified DRP-AC4b. The acetonitrile gradient is indicated by solid line. Multiple charged ions: [M+2H]²⁺, [M+3H]³⁺ and [M+4H]⁴⁺.