

Figure S1. Lipid phosphorus concentration in the supernatant and pellet for each multilamellar vesicle sample used to assess the binding to model membranes. Evaluation of the lipid sedimentation following a 30 minute incubation with several concentrations of the 1018-K6 peptide. **(a)** Zwitterionic lipids (PC); **(b)** Eukaryotic cell (PC₄₀:POPE₄₀:SM₁₅:PS₅); **(c)** *S. Typhimurium* (DOPE₇₈:POPG₁₈:CL₄); **(d)** *Escherichia coli* (PE₈₀:CL₅:POPG₁₅); **(e)** *S. aureus* (CL₄₂:POPG₅₈). The results correspond to the mean \pm standard error (SE) of three independent experiments per model membrane.

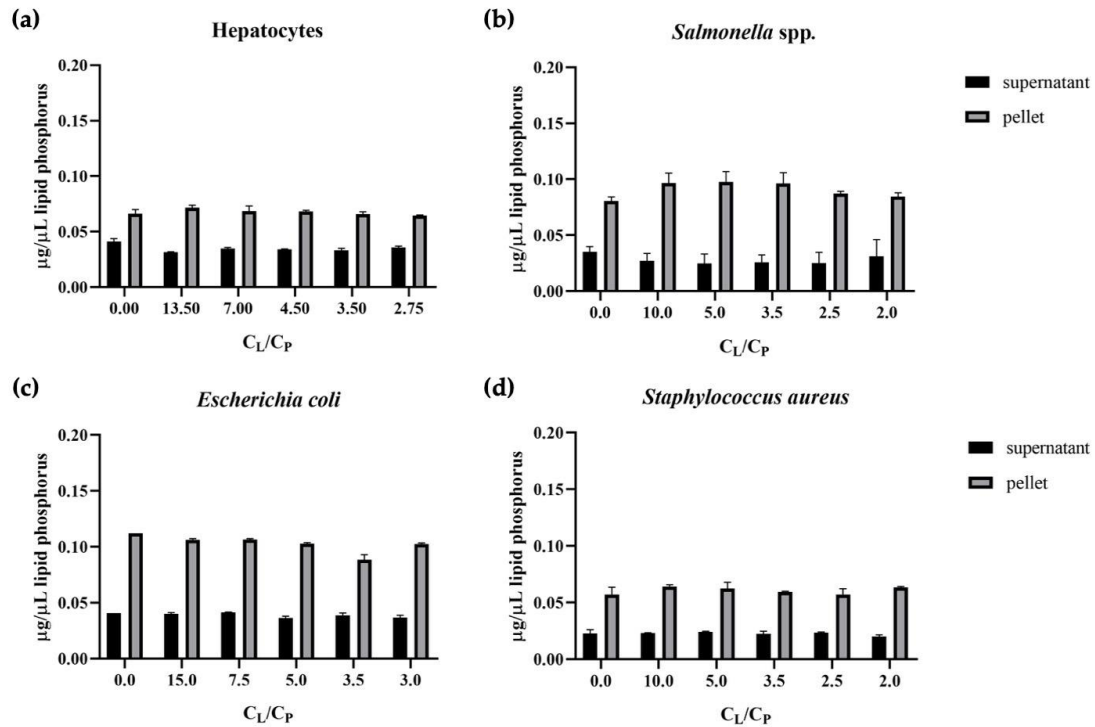


Figure S2. Lipid phosphorus concentration in the supernatant and pellet for each multilamellar vesicle sample used to assess the binding to biological membranes. Evaluation of the lipid sedi-

mentation following a 30 minute incubation with several concentrations of the 1018-K6 peptide. (a) Hepatocytes; (b) *Salmonella* spp.; (c) *E. coli*; (d) *S. aureus*. The results correspond to the mean \pm standard error (SE) of three independent experiments.

Table S1. Binding potential (BP) of AMP 1018-K6 towards multilamellar vesicles of model membranes.

Cell type	Lipid mixture	BP	
		Fluorescence	UV-Vis absorption
Zwitterionic membrane	PC	4.29	2.02
Eukaryotic membrane	PC ₄₀ : POPE ₄₀ : SM ₁₅ : PS ₅	4.91	0.43
<i>Salmonella</i> Typhimurium*	DOPE ₇₈ : POPG ₁₈ : CL ₄	5.17	3.37
<i>Escherichia coli</i>	PE ₈₀ : CL ₅ : POPG ₁₅	4.24	3.62
<i>Staphylococcus aureus</i>	CL ₄₂ : POPG ₅₈	3.32	2.17

*The results refer to the experiments carried out with a higher concentration of peptide.

Table S2. Binding potential (BP) of AMP 1018-K6 towards multilamellar vesicles of biological membranes.

Biological membranes	BP	
	Fluorescence	UV-Vis absorption
Hepatocytes	7.22	18.61
<i>Salmonella</i> spp.	26.87	18.78
<i>Escherichia coli</i>	10.71	42.35
<i>Staphylococcus aureus</i>	31.05	23.42