

## Electronic Supporting Information

### Antibacterial Hydrogels of Aromatic Tripeptides

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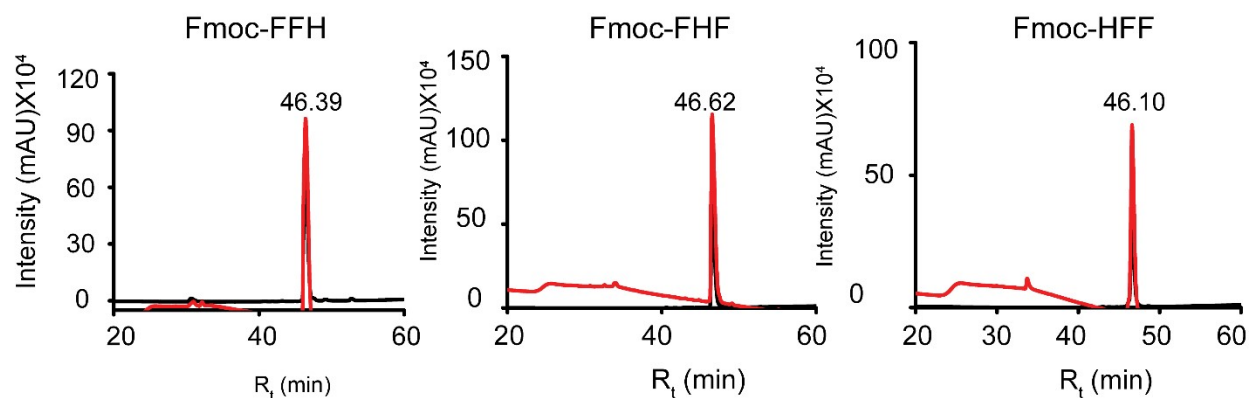
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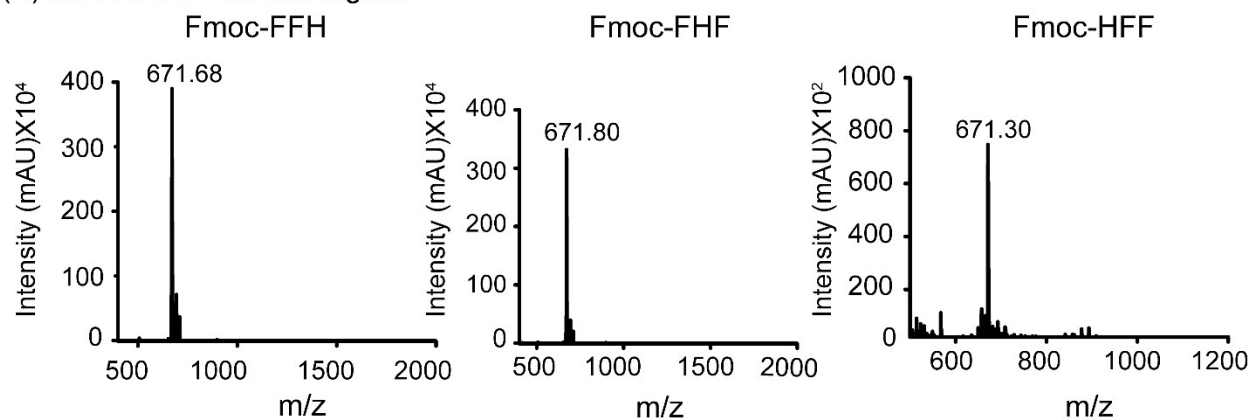
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## 1. Design, Synthesis and Characterization of peptides

### (A) HPLC chromatogram



### (B) MALDI/TOF chromatogram



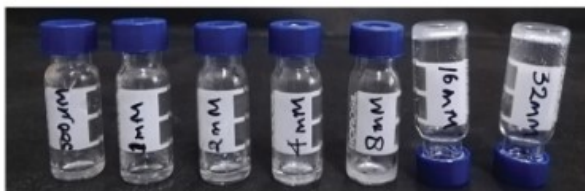
**Fig.S1** Characterization of synthesized peptides; (A) HPLC chromatogram of the synthesized peptide, and (B) Mass spectrum for primary characterization by verifying the mass of the synthesized peptide.

## 2. Characterization of hydrogel formation

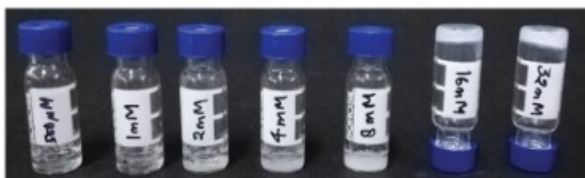
Peptides were dissolved in phosphate buffer (pH 7.4) at different concentrations, and gelation was observed by tube inversion test (Fig. S2)

(A) Concentration dependence of gelation

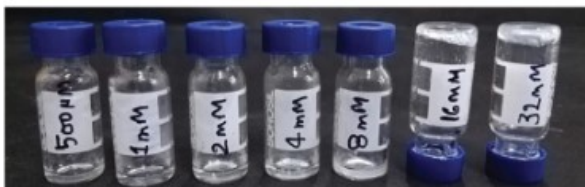
Fmco-FFH



Fmco-FHF

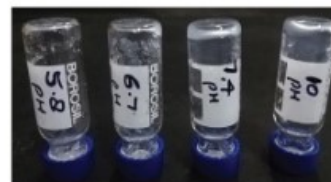


Fmco-HFF

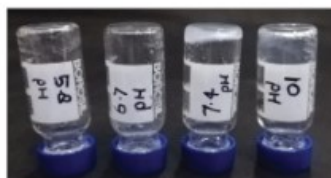


(B) pH optimization for gelation

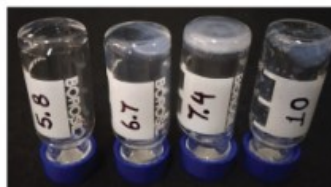
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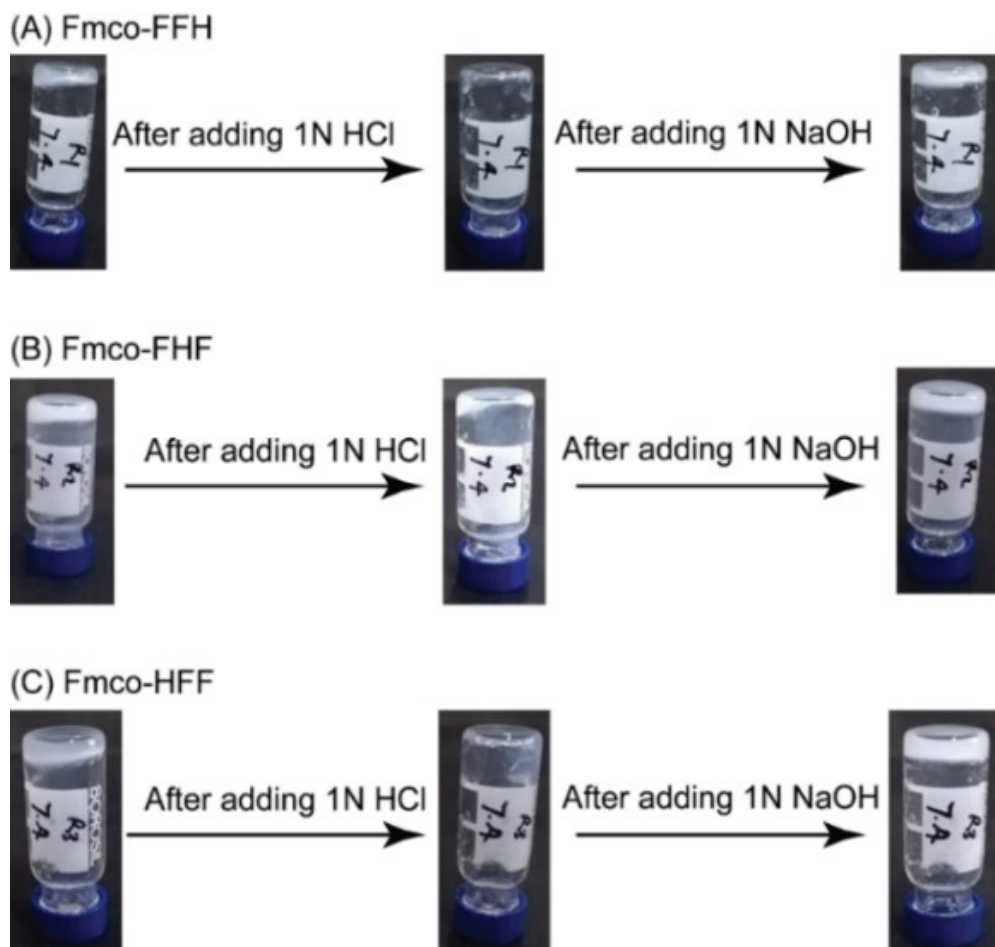
Fmco-FHF



Fmco-HFF



**Fig. S2.** (A) Concentration dependence of hydrogelation and (B) Optimization of pH for hydrogel formation.



**Fig S3.** Dependence of hydrogelation property with pH switch. Hydrogels lose their gelation property in acidic pH but regained again when switched to basic conditions.

**Table S1.** Determination of gelation concentration for peptide hydrogels

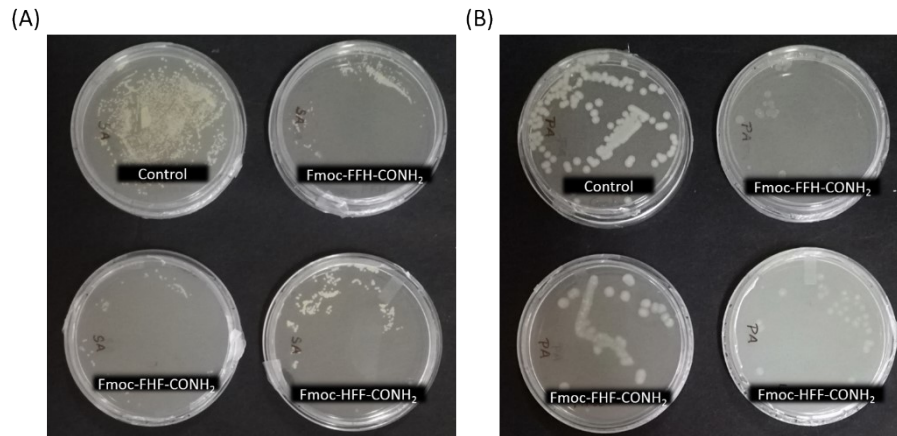
Concentration	Concentration % (w/v)	Gelation Property
500 $\mu$ M	0.03125%	No gel formation
1 mM	0.0625%	No gel formation
2 mM	0.125%	No gel formation
4 mM	0.25%	No gel formation
8 mM	0.5%	No gel formation
16 mM	1%	Gel formation
32 mM	2%	Immediate Gel formation

**Table S2.** Effect of pH on gelation

pH	Gelation Property	Gelation time
5.8	No gel formation	-
6.7	No gel Formation	-
7.4	Gel formation	Immediate
10	Viscous solutions	-

## Spread plate method

Spread plate method was used to validate the antibacterial effect of peptides.



**Fig. S4.** Spread plate method to validate the antibacterial effect of peptides on (A) *Staphylococcus aureus*, (B) *Pseudomonas aeruginosa*