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ARTICLE TYPE

Self-Assembled Surfactant Cyclic Peptide Nanostructures as Stabilizing Agents

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A number of cyclic peptides, including [FR]₄, [FK]₄, [WR]₄, [CR]₄, [AK]₄, and [WK]_n (n = 3-5) containing L-amino acids were synthesized using solid-phase peptide synthesis. We hypothesized that an optimal balance of hydrophobicity and charge could generate self-assembled nanostructures in aqueous solution by intramolecular and/or intermolecular interactions. Among all the designed peptides, [WR]_n (n = 3-5) generated self-assembled vesicle-like nanostructures at room temperature as shown by Transmission Electron Microscopy (TEM), Scanning Electron Microscopy (SEM), and/or Dynamic light scattering (DLS). This class of peptides represents the first report of surfactant-like cyclic peptides that self-assemble into nanostructures. A plausible mechanistic insight of self-assembly of [WR]₅ was investigated by molecular modeling studies. Modified [WR]₅ analogues, such as [W_{Me}R]₅, [WR_{(Me)2}]₅, [W_{Me}R_{(Me)2}]₅, and [WdR]₅ exhibited different morphologies than [WR]₅ as shown by TEM observations. [WR]₅ exhibited significant stabilizing effect for generated silver nanoparticles and glyceraldehyde-3-phosphate dehydrogenase activity. These studies established a new class of surfactant-like cyclic peptides that self-assembled into nanostructures and could have potential applications for stabilizing of silver nanoparticles and protein biomolecules.

1. Introduction

Well-defined nanostructures have drawn significant attention in the fields of biomedicine and biotechnology, material science, and nanotechnology.^{1,2} The formation of ordered structures is facilitated by different types of noncovalent interactions, including electrostatic, hydrogen bonding, hydrophobic, and aromatic stacking.³⁻⁵ In this context, self-assembly of peptides at nanoscale has emerged as a subject of considerable interest in recent years. Various types of peptide nanostructures have been reported with applications in diverse fields, such as antibacterial agents,⁶ neurological regeneration,⁷ and biosensors.^{8,9} Ghadiri et al.¹⁰ showed that cyclic D,L-peptides composed of alternating D- and L- amino acid residues can self-assemble into nanotubes by

forming antiparallel hydrogen bonds between homochiral amino acid residues of adjacent rings.

Peptides have attracted much attention as an alternative to traditional surfactants for studies of membrane proteins, because of their tendency to augment the formation of ordered supramolecular assemblies.¹¹ There is an urgent need to develop more suitable surfactants that can provide natural alternatives for studying the membrane environment. Linear surfactant-like peptides bearing hydrophilic and hydrophobic residues have shown to act as membrane protein stabilizers.¹² Moreover, it has been reported that these peptides can spontaneously undergo self-assembly to form dynamic nanostructures in a pure solution, similar as other lipids and surfactants¹². Compared to linear peptides, cyclic peptides have greater potential as stabilizing agents due to their enhanced chemical and enzymatic stability.¹³ Microbial cyclic peptide surfactants have been previously reported in the literature.^{14,15} Random discovery by screening and challenges in production of these natural surfactants by synthetic or genetic engineering methods have hindered further development of these analogs. Thus, the applications of natural cyclic peptides as surfactants and protein stabilizers remain less explored. We hypothesized that self-assembled cyclic peptides with stabilizing and surfactant properties can be designed rationally. Herein, we report the discovery of nanosized surfactant-like L-cyclic peptides and their potential application as a new class of silver nanoparticle and biomolecule stabilizers. To the best of our knowledge, this class of peptides represents the

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first report of surfactant-like cyclic peptides that self-assemble into nanostructures. Furthermore, this is the first application of any surfactant-like synthetic cyclic peptide as a stabilizer for silver nanoparticles and an enzyme.

2. Results and Discussion

2.1. Self-Assembly of Peptides

A number of cyclic peptides, including [FR]₄, [FK]₄, [WR]₄, [CR]₄, [AK]₄, and [WK]₅ (Fig. 1) containing L-amino acids were synthesized using solid-phase peptide synthesis. These peptide sequences were selected to consist of alternating hydrophobic and positively charged residues or alternating neutral and positively charged residues as controls. We hypothesized that an optimal balance of hydrophobicity and charge could generate self-assembled nanostructures in aqueous solution by intramolecular and/or intermolecular interactions. We have previously reported that among all the synthesized cyclic peptides, amphipathic L-cyclic peptide containing tryptophan and arginine efficiently delivered cargoes intracellularly.¹⁶

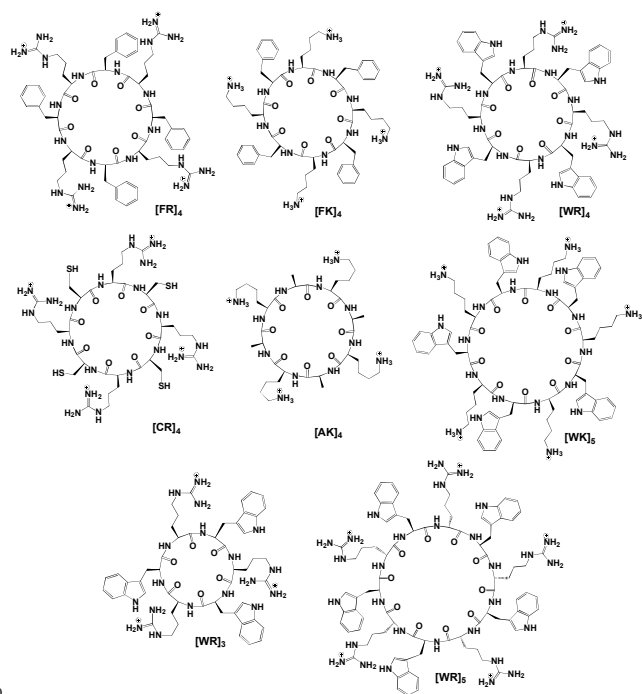


Fig. 1. Chemical Structures of synthesized cyclic peptides (F = phenylalanine, R = arginine, K = lysine, W = tryptophan, C = cysteine, A = alanine).

All the cyclic peptides (2 mM) were dissolved in water and their structural morphologies were investigated using Transmission Electron Microscopy (TEM) after a few days. Among all the designed peptides, [WR]₄ (Fig. 1) generated self-assembled nanostructures after a specific incubation time at room temperature. Six cyclic peptides had the combination of positively-charged residues and other amino acids. The indole ring of tryptophan in [WR]₄ appears to play crucial role in self-assembly because of high hydrophobicity and generating potential hydrophobic interactions. Moreover, the guanidine

group of arginine residues in [WR]₄ could take part in hydrogen bonding interactions. Other 5 cyclic peptides did not have optimal combination of hydrophobic and positively-charged residues required for self-assembly process.

Initial conventional-TEM imaging of [WR]₄ (2 mM aqueous solution) showed vesicle-like structures after 4 days (Fig. S1, Supplementary information). Larger particles were observed by using conventional TEM microscope after a prolonged incubation.

Further evidence for the aggregation of [WR]₄ was observed in a concentration-dependent fluorescence study. At lower concentrations, the fluorescence intensity was proportionally increased (excitation 280 nm, emission 360 nm). However, the intensity remained constant at a higher concentration of 1 mM (Fig. S2). The results suggest that fluorescence-quenching occurs at the concentration that is sufficiently high enough for aggregate formation. Since the hydrophobicity of Trp is weak, this peptide self-assembly occurs at a high concentration (i.e. ≥ 1 mM) possibly through intermolecular hydrophobic interactions between Trp residues.

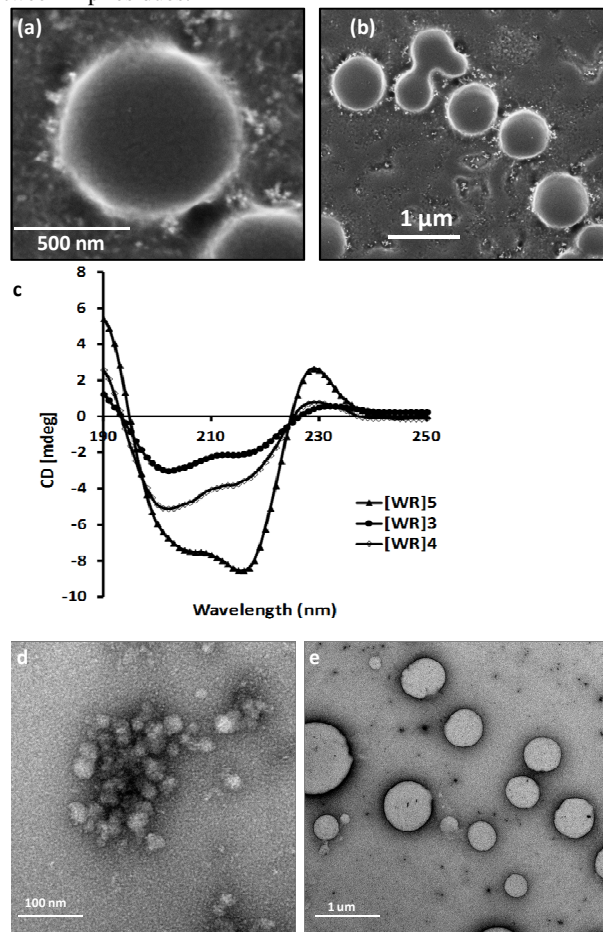


Fig. 2. (a) and (b) FE-SEM image of self-assembled structure of [WR]₅ (2 mM) in water after 20 days; (c) CD spectra of [WR]_n (100 μM) in water, where n = 3-5; (d) TEM images of self-assembly of [WR]₅ (100 μM) in PBS after 48 h; (e) TEM images of [WR]₅ (100 μM) in PBS after 12 days.

As a result of the unusual behavior of [WR]₄, other derivatives in this class ([WR]₃ and [WR]₅ (Fig. 1)) were synthesized and evaluated to investigate whether the ring size has any effect in the

self-assembly process. Structural investigations of analogues of [WR]₃ and [WR]₅ were accomplished by TEM. Vesicle-like structures were formed for all compounds in this class. TEM of [WR]₃ demonstrated small structure sizes of ~100 nm after 14 days incubation. [WR]₄ and [WR]₅ exhibited larger structures (~550 and 930 nm respectively) with much higher frequency (Fig. S1).

Systematic studies on the self-assembly behavior of [WR]₅ were carried out using different tools. Direct dissolution of the peptide in water did not reveal any specific structures after 24 h. TEM images showed the formation of a few micelle-like structures after 48 h. Nanovesicle structures with a diameter between 50 and 100 nm were clearly observed after 12 days (Fig. S4). The formation of vesicles was also confirmed by Scanning Electron Microscopy (SEM) after 20 days incubation (Fig. 2a,b). SEM results clearly show self-assembly by aggregation of small particles to the larger particles.

Zeta-potential measurement by dynamic light scattering (DLS) showed that self-assembled [WR]₅ and [WR]₄ had positive ζ potential values ($+56.2 \pm 7.62$ mV) and ($+31.4 \pm 6.83$ mV), respectively, which further demonstrate the formation of the positively charged surfaces by the guanidinium of R residues.

It is well known that the secondary structure of the peptides plays a critical role on the self-assembly pattern. Thus, circular dichroism (CD) was used to assess the secondary structure of these peptides. CD spectra of aqueous solution of the peptides [WR]_n (n = 3-5, 100 μ M) at 25 °C (Fig. 2c) revealed two negative bands at approximately 202-205 nm, 214-216 nm and a positive band around 229 nm. To estimate the secondary structure content, CD spectra were analyzed using an estimation program.¹⁷ The analysis showed that peptides [WR]_n n = 3-5 to have 37.6-38.8% random and 31.6-35.7% sheet component with the remaining assigned to turn (12.4-12.5%) and helix (11.5-13.1%) (Table S1). The CD spectra of [WR]_n (n = 3-5) displayed a distinct minimum of ellipticity around 202-205 nm, indicating that the peptides exist mostly in random coil states. The appearance of negative band around 214-216 nm and positive band at 229 nm is attributed to the strong exciton coupling produced by the aromatic chromophores of indoles tryptophan residues stacking against one another.¹⁸ In addition, all peptides showed the strong negative bands around 214-216 nm and positive band at around 190, which are characteristic of the formation of distorted β -sheet structures. In general, the backbone-backbone intermolecular hydrogen bonding interactions proceed through parallel or antiparallel β -sheet like stacking arrangements in linear peptides.^{10b} The results indicate that the self-assembly of these cyclic peptides is driven mostly by the hydrophobic force, hydrogen bonding by partial β -structures, and/or π - π stacking interactions between tryptophan residues. These data are consistent with those obtained with the previously reported cyclic peptides¹⁹ where the peptides did not show any specific secondary structure, and they self assembled into bilayer vesicular nanostructures. The CD spectra demonstrated a significant increase in spectral ellipticity of [WR]₅ compared to that of [WR]₃ and [WR]₄, suggesting stronger intramolecular/intermolecular interactions in this peptide. These data are consistent with TEM exhibiting high population of nanostructures formation by [WR]₅. Considering these facts,

[WR]₅ was chosen as a model peptide for further self-assembly studies in detail.

Dynamic light scattering (DLS) was employed to identify the self-assembled structures in solution by determining the size distribution of structures. DLS results showed that the majority (80%) of nanostructures of [WR]₅ (100 μ M) were found in the size range of 150-250 nm after 14 days (Fig. S3, Supplementary Information). Furthermore, it was observed that the size of the vesicle structures became larger over time as confirmed by TEM and SEM studies. This can be attributed to the fusion or aggregation of two or more vesicles as observed in the SEM image (Fig. S4, Supporting Information). This pattern of aggregation resembled with traditional surfactants.

To get a better understanding of the impact of time on the aggregation process, the time-dependent studies by DLS were conducted for [WR]_n (n = 3-5). The samples were prepared in water at the concentration of (1 mM) and the size distribution was evaluated on the first, third, and fourteenth day. The first day presents the immediate size of the formed nanoparticles after dissolving in water. Initially, the peptides will have interactions with water. Additionally, intramolecular interactions contribute to the size distribution at this stage. The intramolecular and water hydrogen bonding interactions are generated mainly by the appropriate functional groups, such as guanidine residues. Subsequently, the size of nanoparticles was compared after 3 days. The self-assembly trend in a short period is mainly related to the initial intramolecular interactions of peptide functional groups followed by the intermolecular interactions. However, the size distribution was also investigated after a relatively longer period (14 days) to determine the behavior of nanoparticles after a prolonged incubation time. At a longer period, it is expected that intermolecular interactions are the dominant driving force of self-assembly.

As depicted in Fig. 3, the majority of [WR]₅ nanoparticles were found in 319 ± 54 nm after 1 day. However, after 3 days, the self-assembly triggered, and [WR]₅ nanoparticles showed larger particles (395 ± 53 nm). This trend was continued, and after 2 weeks as [WR]₅ nanoparticles were observed in 543 ± 41 nm, suggesting that time is a critical element for the size distribution of nanoparticles. Similar pattern was observed for [WR]₄ nanoparticles. The size of [WR]₄ nanoparticles were found to be 306 ± 102 nm, 424 ± 46 nm, 641 ± 175 nm at days 1, 3, and 14, respectively. However, [WR]₃ displayed a completely different pattern. [WR]₃ nanoparticles started increasing from day 1 (241 ± 100 nm) to day 3 (352 ± 7 nm). However, the nanoparticles size was decreased to 216 ± 0.01 nm after 2 weeks. Several factors could be responsible for the collapsing of [WR]₃ nanoparticles, such as nanoparticles instability in the long term due to the small ring size and insufficient intramolecular or intermolecular interactions due to the high strain in the cycle, or inappropriate orientation of amino acids in the structure of peptide for interactions. Smaller ring size in [WR]₃ leads to more puckered and more cup like nonplanar structure of monomer which is less suited for self assemblage. The conformation of monomer dictates the differences in shapes of aggregated supramolecules. These results suggest that although time is a critical element in the size of nanoparticles, stability of the nanoparticles could affect the self-assembly process in the long period.

These data demonstrated the effect of the ring size the nanoparticles size. At day 1, [WR]₃, [WR]₄, [WR]₅ were found to be in 241 ± 100, 306 ± 102, and 319 ± 54 nm respectively. This part of result showed that the number of involved amino acids is an important element in the immediate size distribution of nanoparticles since [WR]₅ led to a larger size of nanoparticles compared to that of [WR]₃. However, [WR]₄ nanoparticles were measured to be in the size of 424 ± 46 nm compared to the size of [WR]₃ (424 ± 46 nm) and [WR]₅ (395 ± 53 nm) after 3 days incubation, suggesting that the ring size with 8 amino acids is optimal for self-assembly. This trend was similar after 2 weeks as [WR]₄ nanoparticles (641 ± 175 nm) showed distinctly larger nanoparticles than [WR]₃ (215 ± 0.01 nm) and [WR]₅ (543 ± 41 nm). These results indicate that a [WR] cyclic peptide with eight amino acids has an optimal ring size for the maximal intermolecular and intramolecular interactions after 14 days. However, primary nanoparticles size at day 1 was found to be ring size dependent.

Additional characterization data were provided for these peptides, such as fluorescence spectroscopy (Fig. S17) and FTIR. Because of the highly sensitive indole chromophore in the structure of the tryptophan, this amino acid was used for its intrinsic fluorescence studies. Three cyclic peptides such as [WR]₃, [WR]₄ and [WR]₅ (100 μM in H₂O) were used for fluorescence studies. The data showed higher fluorescence intensity for peptide containing higher number of tryptophan residues [WR]₅.

ATR-FTIR was used to further characterize the peptide amide bond in [WR]₄. The spectrum of the dry powder of the peptide mixture revealed a two amide-I stretching at 1654 and 1640 cm⁻¹. In general, antiparallel β-structures exhibit a principle β-sheet amide-I stretching vibration maximum at 1630–1640 cm⁻¹ with a secondary absorption at a frequency normally 50–70 cm⁻¹ higher. The absence of the secondary absorption shows that [WR]₄ adopts partially a parallel β-structure orientation. This result is consistent with stacking observed from the self-assembly simulations (Fig. 3).

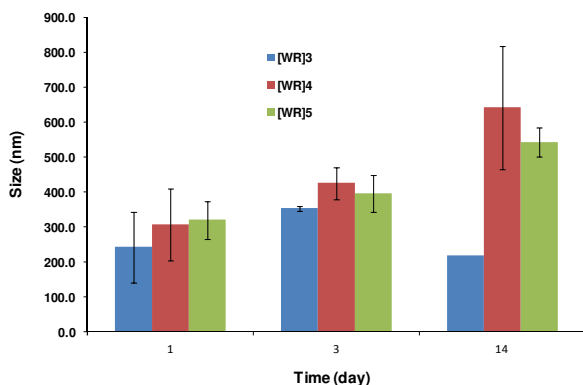


Fig. 3. DLS studies of [WR]_n (n = 3-5) (1 mM) at days 1, 3, and 14.

Furthermore, the effect of salts and pH was investigated in the self-assembly process. [WR]₅ showed self-assembled structures in phosphate buffered saline (PBS, pH 7.0) (Fig. 2e). The aggregation was faster in PBS compared to an aqueous solution resulting in the formation of a larger size of vesicles (500-1000

nm) after 12 days (Fig. 2f). This could be due to the presence of different salts that can facilitate the self-assembly. CD also revealed the development of additional positive band at 196 nm in PBS. The pattern of structures formation has a resemblance with surfactant-like linear peptides reported by Nagai et al.²⁰ Moreover, it is well known that NaCl favors surfactant aggregation.²¹ When PBS containing NaCl (167 mM) was examined in the self-assembly of this peptide, the aggregation of [WR]₅ was accelerated. NaCl decreases the repulsive force among the charged head groups, resulting in enhancement of hydrophobic interactions between the peptides. However, in acidic medium (0.02 N HCl), some irregular structures were observed initially but longer incubation time led to degradation of the structures (Fig. S5, Supporting Information).

In order to assess the different self-assembly behavior of [WR]₅ in water and PBS, turbidity measurements²² were carried out to determine vesicle formation at 350 nm. Difference in turbidity was clearly visible from the results. A sudden increase in the absorbance was observed above certain concentrations in both cases. Critical Aggregation Constant (CAC) values were determined from the intersection point obtained by extrapolating the lines. The CAC values for this peptide in water and PBS were found to be 0.9 mM and 0.45 mM, respectively (Fig. S6, Supporting Information), suggesting surfactant-like nature of the peptide in the presence of salts.

Fluorescence studies were performed for [WR]₄ to show the aggregation of the peptide at concentrations above 1 mM (Fig. S2) consistent with the data described in Fig. S6 for [WR]₅. The aggregation of [WR]₄ was observed in a concentration-dependent fluorescence study. At lower concentrations, the fluorescence intensity was proportionally increased (excitation 280 nm, emission 360 nm). However, the intensity remained constant at a higher concentration of 1 mM (Fig. S2, Supplementary Information).

As shown above, the self-assembly was promoted in the presence of salts. Molecular modeling studies were used to gain better insights of nanostructure formation with [WR]₅ in the presence of counterions. Initial coordinates for [WR]₅ were obtained using GAUSSVIEW.²³ Ab initio Hartree Fock molecular orbital calculations with complete geometry optimizations were performed with 6-31G²⁴ basis set on monomer [WR]₅ utilizing GAUSSIAN'03 software. Two conformations were obtained (Fig. S7, Supporting Information). The lowest energy conformation did not allow any intramolecular H-bonding. Higher energy conformation (25 kCal/mol) shows intramolecular H-bonding between cationic and neutral side chains. Side chains have to reorganize for efficient intramolecular H-bonding resulting in higher energy conformation. In the lowest energy conformation tryptophan side chains are disposed perpendicular to backbone. Higher energy conformation being more planar seems more suited for stacking. Stacking efficiency was explored utilizing supermolecule type intermolecular interaction calculations. Stacking efficiency = $E_{\text{stacked supermolecule}} - (E_{\text{monomer}} \times \text{no. of monomers} + E_{\text{counterion}} \times \text{no. of counterions})$.

[WR]₅ is an amphiphilic peptide. Cationic side chains may facilitate counterion assisted self-assembly. First, interaction of cationic side chains with small fluoride ions was studied (Fig. S8, Supporting Information). Fluoride counterions assisted self-

assembly propensity of [WR]₅ system and resulted in nanotubular formation (Fig. S9, Supporting Information). Then interaction of trifluoroacetate ions (CF₃COO⁻) with cationic side chain was calculated, and counterion-mediated stacking was explored. The data indicated that [WR]₅ can undergo self-assembly spontaneously in the presence of trifluoroacetate counterions. Trifluoroacetate counterions are larger in size. Thus, nanotubular stacking was not preferred. Instead, the system presented spiral noncovalent association (Fig. 4), giving rise to nanostructure formation. Longer period of self-assembly simulation generated structures with larger size as observed in TEM and SEM experiments. We assume that intramolecular along with intermolecular interactions, such as hydrogen bonding, π - π interactions, and counterion interactions work cooperatively during the self-assembly process of [WR]₅. Intramolecular hydrogen bonding between W and R residues results in the high energy conformation which is suited for stacking hence it indirectly helps. Intermolecular interaction is purely electrostatic between cationic R residues and counteranions which may be basically any anion small enough to squeeze between monomers and interact efficiently. In [WR]₅, self-assembly does not depend mainly on diffusion of the peptide, but certainly depends on diffusion of any medium i.e. without any counterions, [WR]₅ show only minimal aggregation as it is not thermodynamically favored.

We propose that the hydrophobic segments in the peptide initialize the supramolecular self-assembly and drive the peptide to form nanospherical water-filled structures. The positively charged portion of the peptide containing arginine residues orients to face the aqueous and counterion environment. Moreover, the hydrophobic residues stabilize the nanostructure as the particles become larger through inter- and intra-molecular hydrogen bonding. Additionally, the relatively weaker hydrogen bonded segments in the secondary structures facilitate the peptide to manage the assemblies at extremely lower concentrations (e.g. 100 μ M). As it is shown in Fig. 2c, increasing the hydrophobicity character by increasing the number of tryptophan residues from 3 to 5 for [WR]₅ induced the formation of completely different secondary structure as it is shown by CD. However, as it is exhibited in Fig. 3, the size distribution of [WR]₅ is relatively smaller than that of [WR]₄ after 14 days showing that other parameters are also involved in the self-assembly process.²⁵

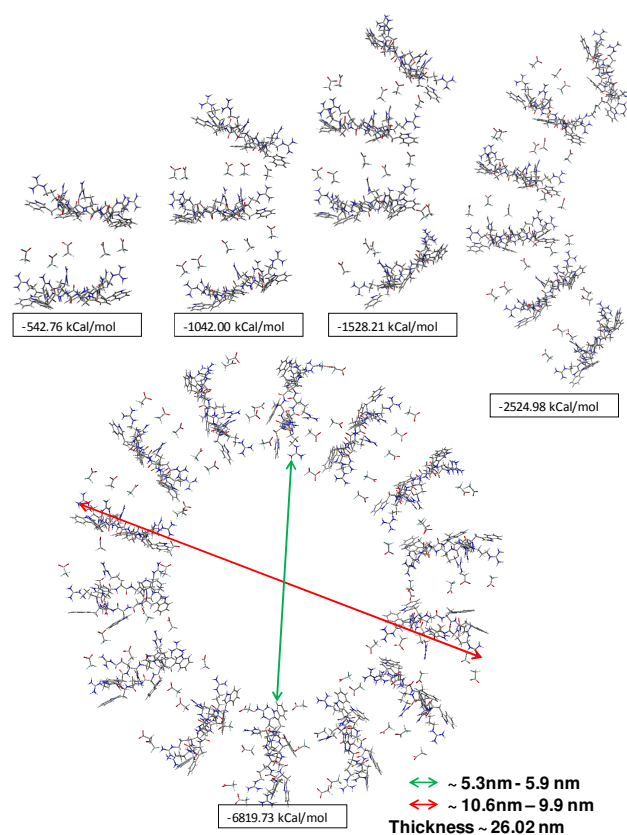


Fig. 4. Self-assembled nanostructure formation of [WR]₅ in the presence of CF₃COO⁻ ions. Stacking efficiency is given in kCal/mol.

Additional molecular modeling studies were then conducted in the presence of a 1-palmitoyl-2-oleoyl-sn-glycero-3-phosphocholine (POPC) membrane bilayer using SPC water model. Membrane was set up by system builder module. Then water was added. Whole system was subjected to simulated annealing followed by molecular dynamics at room temp for 5 ns. When the side view is observed carefully, it does show two distinct surfaces. Results of 5 ns trajectory showed that [WR]₅ dimer aligned at oil (mimicked by membrane)-water interface with hydrophobic surface containing tryptophan residues towards the hydrophobic residues in membrane, suggesting surfactant property of the peptide (Fig. S10). The stability of system at the interface and low RMSD for all heavy atoms with time indicated successful simulation of mode of action for [WR]₅ system as a biosurfactant or peptide surfactant (pepfactant). At oil-water interface mimicked by membrane-water interface in these simulations, this aggregated supramolecular form stays as such with hydrophobic groups towards membrane and hydrophilic groups towards water indicating the type of micellar formation

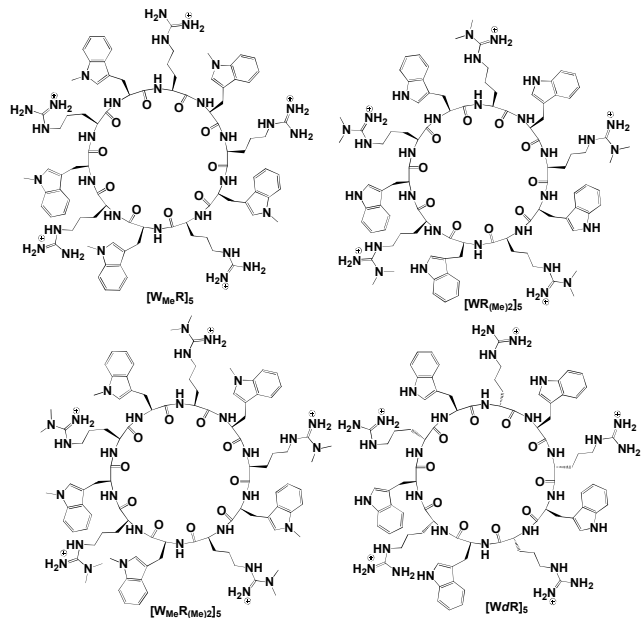


Fig. 5. Synthesized $[WR]_5$ analogues.

In general, the intrinsic properties of a peptide sequence including hydrophobicity, charge, secondary structure propensity, and sequence patterning are responsible for controlling the self-assembly process.^{26,27} Thus, in order to investigate the effect of hydrophobicity and charge in the self-assembly process and to gain further mechanistic insight into the packing of self-assembly of $[WR]$ series, a class of cyclic peptide analogues of $[WR]_5$ was designed rationally by using modified building blocks of arginine and tryptophan (Fig. 5), and their self-assembly pattern was investigated.

Modified $[WR]_5$ analogues exhibited different morphologies depending on the hydrophobicity and charge of the amino acids in the structure of the peptides as shown by TEM observations. To investigate the effect of hydrophobicity of tryptophan in the self-assembly, a new building block *N*-methyl-*L*-tryptophan (W_{Me}) was used as the hydrophobic residue in place of tryptophan. The resulting cyclic peptide $[W_{Me}R]_5$ showed nanotubular structures instead of spherical vesicles as evidenced by TEM (Fig. 6). Here, enhanced hydrophobicity of the peptide and the absence of NH moiety of tryptophan played critical roles in the self-assembly process.

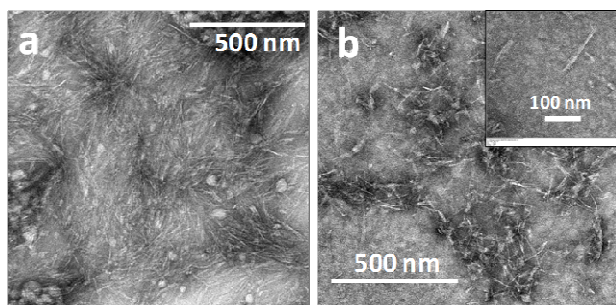


Fig. 6. TEM images of self-assembled structures of $[W_{Me}R]_5$ (2 mM in water) after (A) 6 days and (B) 12 days. Individual nanotube structure is shown in inset (scale: 100 nm).

To determine the role of positively-charged arginine in the

self-assembly process, another analogue $[WR_{(Me)_2}]_5$ was synthesized by replacing the arginine with *N,N*-dimethylarginine keeping tryptophan as a hydrophobic residue unchanged. Thus, the hydrophilicity of the charged residues was reduced that resulted in poor water solubility. Unlike the previous peptides, this compound (1 mM aqueous solution) generated micelle-like spherical nanostructures (Fig. S11, Supporting Information). It was found that the compound did not exhibit any large vesicles like $[WR]_5$ even after prolonged incubation (25 days), presumably due to the absence of one of free amine groups of guanidine. These data suggest that this free amino group contributes to the self-assembly process of $[WR]_5$ possibly through hydrogen bonding. On the other hand, a peptide containing both *N*-methyltryptophan (W_{Me}) and *N,N*-dimethylarginine ($R_{(Me)_2}$) ($[W_{Me}R_{(Me)_2}]_5$) did not produce any specific structures at 2 mM or 4 mM concentration. At higher concentration (4 mM), $[W_{Me}R_{(Me)_2}]_5$ exhibited a few micelle-like nanostructures along with patches of membrane-like sheets between the nanostructures and copper grid during sample preparation (Fig. S12, Supporting Information). This peptide might have undergone self-assembly on the surface of copper grid by hydrophobic interactions to form monolayer membranes, possibly due to the highly hydrophobic nature of the amino acids.

The formation of nanotubes through the self-assembly of D,*L*-cyclic peptides has been previously reported.¹⁰ The backbone-backbone interactions between peptides were found to be critical for the self-assembly of D,*L*-cyclic peptides. Our results indicated that hydrophobicity and the charge were crucial for the self-assembled structure formation for these cyclic peptides, as they were in the mechanism of linear surfactant-like peptides.¹¹ Thus, we compared the nanostructure formation between homochiral $[WR]_5$ with the corresponding heterochiral to evaluate the effect of stereochemistry. Thus, D,*L*-cyclic peptide $[WdR]_5$, where *dR* is *d*-arginine, was designed and synthesized for the comparative stereochemistry studies. Unlike the nanotube formation by D,*L*-cyclic peptides, $[WdR]_5$ exhibited micelle like nanostructures after prolonged incubation at room temperature (Fig. S13, Supporting Information). The detailed mechanism of the self-assembly for these cyclic peptides requires further investigation with modeling and simulation experiments.

Qualitative analysis of the conformation of these peptide analogues was obtained by CD. CD spectra of aqueous solution of the peptides $[W_{Me}R]_5$ and $[WR_{(Me)_2}]_5$ at room temperature (Fig. S14, Supporting Information) revealed two negative bands at approximately 203 and 217 nm. This does not resemble either a classic β -sheet or α -helical structures. Another analogue $[W_{Me}R_{(Me)_2}]_5$ also showed similar bands, but with a lower ellipticity. These three peptide analogues, namely $[W_{Me}R]_5$, $[WR_{(Me)_2}]_5$ and $[W_{Me}R_{(Me)_2}]_5$, showed negative band at around 217 nm and positive band at around 229 nm, which are characteristics of exciton coupling of tryptophan residues, indicating the presence of stacking interactions. However, $[WdR]_5$ did not exhibit any characteristic signatures of exciton coupling of tryptophan side chain, suggesting the absence of stacking of indole rings.

2.2. Stabilization of Silver Nanoparticles by [WR]₅ and Modified Analogues

We have previously reported the application of [WR]₅ as a nuclear targeting molecular transporter for a wide range of cargos.¹⁶ It was found out that a physical mixture of [WR]₅ and HAuCl₄ led to the formation of peptide-capped gold nanoparticles and enhanced the cellular uptake of drugs significantly.²⁸ We investigated to determine whether [WR]₅ can be used for the stabilization of silver nanoparticles and proteins. Several methods for the synthesis of silver nanoparticles have been previously reported. However, the stabilization of silver nanoparticles is required for various applications. The stabilizing agents (e.g., surfactants, polymer) prevent the particle aggregation by protecting these particles by steric stabilization. Cationic surfactants are commonly used as stabilizing agents in silver nanoparticles synthesis.²⁹ Accounting the above facts, we postulated that [WR]₅ could behave as a cationic surfactant.

In order to explore whether [WR]₅ can mimic the role of cationic surfactants in the synthesis of nanoparticles, a model experiment was performed using silver nitrate (AgNO₃) as a source of Ag⁺ ion and sodium borohydride (NaBH₄) as a reducing agent in the presence and absence of the cyclic peptide. Sodium borohydride was added into the equal volume mixture of cyclic peptide [WR]₅ (1 mM) and AgNO₃ (2 mM) solution. To demonstrate the role of cyclic peptide as stabilizer, sodium borohydride was used in excess amount (8 mM). The resulting mixture produced yellow color immediately, indicating the formation of silver nanoparticles. UV-Vis spectra showed the characteristic plasmon band of silver nanoparticles formation with a band at 410 nm (Fig. 7A). The formation of spherical silver nanoparticles was confirmed by TEM (Fig. 7B). In order to monitor the stability of the silver colloid, absorption of the colloid was measured after four weeks. The evolution of UV-Vis spectra is shown in Fig. 7A (curve b). There was only a minor red shift in peak position even after four weeks (curve c), confirming the stability of the silver colloids. No extra peaks or shoulders were developed even after one month, eliminating the possibility of aggregation. On the contrary, when sodium borohydride was added to AgNO₃ (2 mM) solution in the absence of the peptide (Fig. 7A, curve a), stable silver nanoparticles were not formed. These data confirm the surfactant-like nature of the cyclic peptide [WR]₅. The stability of silver nanoparticles was also examined in the presence of the corresponding linear peptide. The linear peptide was not efficient in stabilizing silver nanoparticles (Fig. S18).

The stabilization of silver nanoparticles with [WR]₅ was compared with a traditional surfactant (cetyltrimethylammonium bromide, CTAB). CTAB is a well known cationic surfactant which is used as a stabilizer in nanoparticle synthesis. It is already reported in the literature that CTAB shows better stabilizing effect at low concentration.³⁰ The stability of silver nanoparticles in the presence of CTAB was checked after 4 days, and it showed aggregation of nanoparticles. The decrease in band (~410 nm) intensity and development of extra band approximately at 610 nm indicate particle aggregation (Fig. S19). Traditional surfactant or some polymeric compounds form micelles which are used to stabilize the metal nanoparticles.^{31,32} The micelles are porous cluster with deep water filled cavities. In

case of traditional surfactants, micelles control the shape of the nanoparticles by trapping the metal ion and reagent in the small cavities. Here, micelles formed by peptides, are water rich. Due to self-assembly process, cyclic peptides initially form micelles which stabilize the nanoparticles. Thus, when AgNO₃ and peptide solution are mixed, Ag⁺ ions may reside into the micelles. Once NaBH₄ is added, Ag⁺ ions are converted into Ag⁰ that remains stable without any aggregation. Small surfactant-like peptides surround the metal nanoparticles and protect them from aggregation. TEM images shows clear layer of the peptide on nanoparticle surface (Fig. S20). This follows the similar model shown by a traditional surfactant.

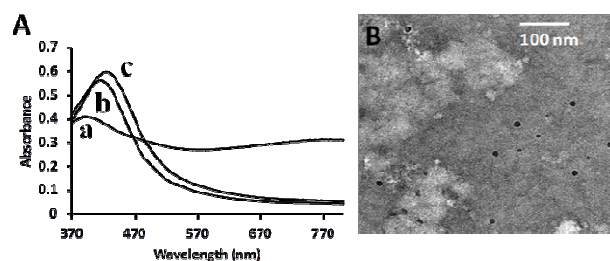


Fig. 7. (A) UV-Vis spectra of yellow color silver nanoparticles solution in the absence (a) and presence (b, c) of [WR]₅ peptide and comparison of the stability of silver nanoparticles in the presence of peptide after 30 min reaction (curve b) and one month (curve c); (B) TEM image of silver nanoparticles synthesized by sodium borohydride in the presence of [WR]₅.

The effect of the peptide concentration in stabilizing silver nanoparticles was investigated by using different concentrations of peptide. Visual observation showed that the typical color of silver nanoparticles changed from pale yellow to dark yellow and brown, respectively, indicating the alteration of nanoparticle morphology in the presence of different concentrations of peptide. This observation was consistent with UV-Vis spectroscopy (Fig. S15, Supporting Information) that showed a narrow band at 401 nm when 100 μM of the peptide was used. As the concentration of the peptide was increased from 100 to 200 μM, the absorbance curve did not change significantly. However, the absorbance band became short and shifted towards longer wavelength when a peptide concentration of 500 μM was employed, indicating the aggregation and formation of larger particles. The absorbance band became broad and significantly shorter at the peptide concentration of 1 mM. The broadening of the band suggests wide distribution (higher dispersity) of particles size. These results revealed that the optimization of the peptide concentration is necessary to achieve narrower particle size distribution. This pattern is similar to other synthesized surfactant-stabilized silver nanoparticles.

To investigate whether the other synthesized cyclic peptide analogues can show surfactant-like properties, they were explored as stabilizing agents in the silver nanoparticles synthesis. Among them, [W_{Me}R]₅ showed a great potential as a stabilizing agent. Although it shows slight red shift (6 nm) in the band after one week, but there is no sign of aggregation of silver nanoparticles (Fig. 8A). This result supports the surfactant-like nature of [W_{Me}R]₅. However, the intensity of absorbance was decreased in the presence of [WR_{(Me)2}]₅ after 7 days (Fig. 8B), indicating the

aggregation of nanoparticles. Furthermore, in the presence of $[W_{Me}R_{(Me)_2}]_5$, the plasmon band of silver nanoparticles shifted (46 nm) significantly to longer wavelength indicating the formation of large particles (Fig. 8C). Similar results were obtained for $[WdR]_5$ (Fig. 8D). This could be attributed to the irregular structure formation by these peptides.

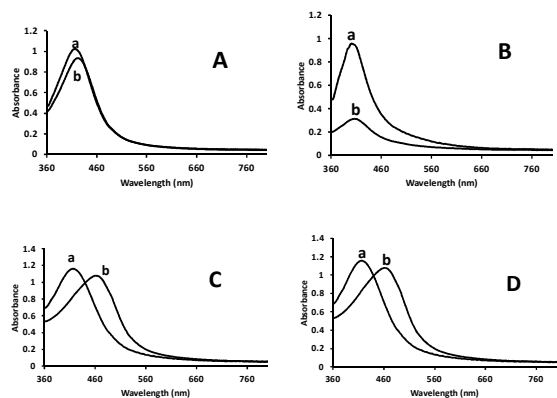


Fig. 8. UV-vis spectra of silver nanoparticles in the presence of (A) $[W_{Me}R]_5$, (B) $[WR_{(Me)_2}]_5$, (C) $[W_{Me}R_{(Me)_2}]_5$, and (D) $[WdR]_5$ at room temperature after 15 min (a) and 7 days (b).

2.3. Stabilization of Glyceraldehyde-3-Phosphate Dehydrogenase (GAPDH) by $[WR]_5$

It is already known that surfactants influence the enzymatic activities depending upon the interaction with the surface of the enzymes.³³ Thus, the capability of this surfactant-like peptide in stabilizing biomacromolecules was investigated using glyceraldehyde-3-phosphate dehydrogenase (GAPDH) as a model enzyme. The activity of GAPDH enzyme was determined by using KDalert GAPDH assay kit. The KDalert GAPDH assay determines the conversion of NAD^+ to NADH by GAPDH in the presence of phosphate and glyceraldehyde-3-phosphate (G-3-P). Under the suggested assay conditions, the rate of NADH production is proportional to the amount of GAPDH enzyme present. To evaluate the effectiveness of the peptide in stabilizing enzymatic activity, experiments were carried out by mixing the enzyme with various peptide concentrations in phosphate buffered saline (PBS, NaCl (167 mM), pH 7.0). GAPDH was mixed in PBS as a control. The activity of the enzyme at different time intervals in the presence and absence of the peptide was measured using the fluorescence-based method. Plots were made to show the increase in fluorescence due to NAD^+ reduction vs time. It was observed that enzyme containing 100 μM of $[WR]_5$ showed significantly enhanced activity compared to the control enzyme (Fig. 9A), suggesting surfactant-like nature of the peptide. Moreover, the enzyme activity was reduced at higher peptide concentration (500 μM), presumably due to the denaturation of the enzyme, indicating the ionic nature of the peptide-surfactant. A control experiment was also performed without the enzyme to rule out any effect of the peptide in the reduction process. Negligible fluorescence was recorded when the peptide solution was mixed with other components in the absence of the enzyme.

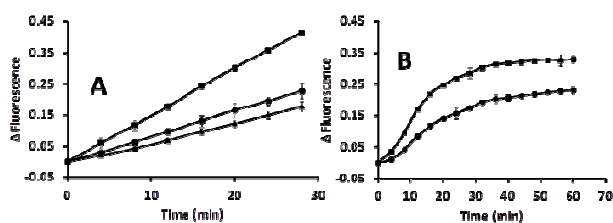


Fig. 9. Enzymatic activity of GAPDH. Changes in fluorescence due to NAD^+ reduction are the result of GAPDH activity. (A) Enzymatic activity shown till 30 min at room temperature; (B) Enzymatic activity shown till 60 min at 50 $^{\circ}C$. (●) GAPDH enzyme in PBS, (■) GAPDH enzyme in PBS in the presence of $[WR]_5$ (100 μM), (▲) GAPDH enzyme in PBS in the presence of $[WR]_5$ (500 μM).

The presence of peptide-surfactant enhanced the activity of the enzyme as described. Thus, it was essential to verify the influence of the peptide on the stability of the enzyme. To this notion, the activity of GAPDH was determined in the presence of the peptide at 37 $^{\circ}C$ and 50 $^{\circ}C$. The enzyme in the presence of the peptide showed 2-fold enhanced activity compared to control one at 50 $^{\circ}C$ (Fig. 9B) and 37 $^{\circ}C$ (Fig. S16, Supporting Information), indicating enhanced stability of GAPDH enzyme in the presence of the peptide-surfactant. These results suggest that this surfactant peptide could be a promising additive for stabilizing other biomacromolecules, such as membrane proteins.

3. Experimental

3.1. Synthesis of Cyclic Peptides

The synthesis of cyclic peptides was carried out according to the previously reported procedure by us.¹⁶ All reactions were carried out in Bio-Rad polypropylene columns by shaking and mixing using a Glass-Col small tube rotator in dry conditions at room temperature unless otherwise stated. In general, all linear peptides were synthesized by the solid-phase synthesis strategy employing *N*-(9-fluorenyl)methoxycarbonyl (Fmoc)-based chemistry and Fmoc-L-amino acid building blocks. 2-(1*H*-Benzotriazole-1-yl)-1,1,3,3-tetramethyluronium hexafluorophosphate (HBTU) and *N,N*-diisopropylethylamine (DIPEA) in *N,N*-dimethylformamide (DMF) were used as coupling and activating reagents, respectively. Fmoc-amino acid trityl chloride resins, 2-chlorotrityl chloride resin, coupling reagents, and Fmoc-amino acid building blocks were purchased from Chempep (Miami, FL). Other chemicals and reagents were purchased from Sigma-Aldrich Chemical Co. (Milwaukee, WI). Fmoc deprotection at each step was carried out using piperidine in DMF (20% v/v). Side chain protected peptides were cleaved from the resins by shaking the resins with a mixture of trifluoroethanol (TFE)/acetic acid/dichloromethane (DCM) (2:2:6, v/v/v, 15 mL) for 2 h. The resins were filtered off, and the liquid was evaporated to dryness to get side-chain protected linear peptide. The cyclization of the peptides was carried out in the presence of a mixture of 1-hydroxy-7-azabenzotriazole (HOAt, 163.3 mg, 1.2 mmol) and *N,N'*-diisopropylcarbodiimide (DIC, 187 μL , 1.2 mmol) in dry DMF (100 mL), and dry DCM (40 mL) for 24 h. DMF and DCM were evaporated. The side chain deprotection was carried with trifluoroacetic acid (TFA)/thioanisole/anisole/1,2-ethanedithiol

(EDT) (90:5:2:3 v/v/v/v) for 2 h. The crude peptides were precipitated by the addition of cold diethyl ether (75 mL, Et₂O) and purified by reversed-phase Hitachi HPLC (L-2455) on a Phenomenex Prodigy 10 μm ODS reversed-phase column (2.1 cm × 25 cm) with using a gradient system. The peptides were separated by eluting the crude peptides at 10.0 mL/min using a gradient of 0-100% acetonitrile (0.1% TFA) and water (0.1% TFA) over 60 min, and then, they were lyophilized. The chemical structure of [WR]₄ was characterized by nuclear magnetic resonance spectra (¹H NMR) determined on a Varian NMR spectrometer (500 MHz) as described previously.¹⁶ The chemical structures of all final products were confirmed by a high-resolution Biosystems QStar Elite time-of-flight electrospray mass spectrometer. The purity of final products (>95%) was confirmed by analytical HPLC. The analytical HPLC was performed on the Hitachi analytical HPLC system on a C18 Shimadzu Premier 3 μm column (150 cm × 4.6 mm).

3.2. Synthesis of Fmoc-Trp(Me)-OH

N-(9-Fluorenylmethoxycarbonyloxy)succinimide (Fmoc-ONSu, 11.55 g, 0.034 mol) was added to a round bottom flask, dissolved with 75 mL of DMF with constant stirring at 0 °C. 1-Methyl-L-tryptophan (5 g, 0.22 mol) was added to sodium carbonate solution (10%, 75 mL), and the mixture was stirred at room temperature to form a suspension. The suspension prepared above was added dropwise at 0 °C with constant stirring to the solution of Fmoc-ONSu in DMF. The precipitate formed immediately. The mixture was stirred for another 1 h and made at room temperature. Acetone (80 mL) was added to the mixture and the stirring continued overnight. The precipitate was removed by filtration, and the pH of the filtrate was adjusted to 7.0 with HCl. The solution was concentrated in a rotary evaporator. A 200 mL of ethyl acetate was added to the mixture and resulted solution was stirred for 6 h at room temperature. Ethyl acetate layer was washed with 100 mL of 0.1N HCl two times and then with brine (75 mL), and dried over anhydrous MgSO₄. The solvent was removed with rotary evaporator to yield crude creamy solid compounds. The compound was re-crystallized in ethyl acetate to afford 4 g of pure material, which was used without further purification for the synthesis of cyclic peptides [W_{Me}R]₅, [WR_{(Me)2}]₅, and [W_{Me}R_{(Me)2}]₅ according to the previously reported procedure.¹⁶

[FK]₄: HR-MS (ESI-TOF, m/z): [M]⁺ calcd for C₆₀H₈₄N₁₂O₈, 1100.6535; found, 1100.5836 [M]⁺, 551.2918 [M + 2H]²⁺, 367.8752 [M + 3H]³⁺.

[AK]₄: HR-MS (ESI-TOF, m/z): [M]⁺ calcd for C₃₆H₆₈N₁₂O₈, 796.5283; found, 796.4868 [M]⁺, 399.2434 [M + 2H]²⁺.

[CR]₄: HR-MS (ESI-TOF) (m/z): [M + H]⁺ calcd for C₃₆H₆₉N₂₀O₈S₄, 1037.4490; found, 1037.4096 [M + H]⁺, 345.1323 [M + 3H]³⁺, 259.1024 [M + 4H]⁴⁺.

[FR]₄: HR-MS (ESI-TOF) (m/z): [M]⁺ calcd for C₆₀H₈₄N₂₀O₈, 1212.6781; found, 1212.6052 [M]⁺, 607.3026 [M + 2H]²⁺, 405.5428 [M + 3H]³⁺, 304.1588 [M + 4H]³⁺.

[WR]₃: HR-MS (ESI-TOF) (m/z): [M + H]⁺ calcd for C₅₁H₆₇N₁₈O₆, 1027.5491; found, 1027.6030 [M + H]⁺.

[WR]₄: HR-MS (ESI-TOF) (m/z): [M + H]⁺ calcd for C₆₈H₈₉N₂₄O₈, 1369.7295; found, 1369.7934 [M + H]⁺, 685.3967

[M + 2H]²⁺, 457.2696 [M + 3H]³⁺, 343.2066 [M + 4H]⁴⁺; FT-IR of [WR]₄ was carried out using Thermo-Nicolet 380 FT-IR spectrophotometer using OMNIC software. FT IR (cm⁻¹) (ATR): 1654, 1640.36, 1531.22, 1494.39, 1431.63, 1198.35, 1170.79, 1119.02, 837.31, 799.83, 745.55, 719.71.

[WR]₅: HR-MS (ESI-TOF) (m/z): [M + H]⁺ calcd for C₈₅H₁₁₁N₃₀O₁₀, 1711.9099; found, 1711.6974 [M + H]⁺, 571.5658 [M + 3H]³⁺, 428.9286 [M + 4H]⁴⁺, 343.1493 [M + 5H]⁵⁺.

[W_{Me}R]₅: HR-MS (ESI-TOF) (m/z): [M + H]⁺ calcd for C₉₀H₁₂₁N₃₀O₁₀, 1781.9882; found, 1782.0165 [M + H]⁺, 594.6633 [M + 3H]³⁺, 446.4739 [M + 4H]⁴⁺, 357.4033 [M + 5H]⁵⁺.

[WR_{(Me)2}]₅: HR-MS (ESI-TOF) (m/z): [M + H]⁺ calcd for C₉₅H₁₃₁N₃₀O₁₀, 1852.0664; found, 1852.1065 [M + H]⁺, [M + 2H]²⁺, 618.3545 [M + 3H]³⁺, 463.9962 [M + 4H]⁴⁺, 371.4213 [M + 5H]⁵⁺.

[W_{Me}R_{(Me)2}]₅: HR-MS (ESI-TOF) (m/z): [M + H]⁺ calcd for C₁₀₀H₁₄₁N₃₀O₁₀, 1922.1447; found, 1921.5000 [M + H]⁺, 483.9131 [M + 4H]⁴⁺, 383.6360 [M + 5H]⁵⁺.

[W_dR]₅: HR-MS (ESI-TOF) (m/z): [M]⁺ calcd for C₈₅H₁₁₀N₃₀O₁₀, 1710.9021; found, 571.6383 [M + 3H]³⁺, 428.9571 [M + 4H]⁴⁺, 343.3879 [M + 5H]⁵⁺.

3.3. Transmission Electron Microscopy (TEM)

3.3.1. Conventional TEM

TEM sample was prepared by drop casting a 2 mM aqueous solution (10 μL) onto the formvar coated carbon grid of mesh size 300, which was allowed to sit for 2 min. Excess solvent was carefully removed by capillary action (filter paper). Grids were allowed to dry at room temperature for overnight and images were taken using JEOL Transmission Electron Microscope (Tokyo, Japan) maintained at 80 kv.

3.3.2. Negative Staining

After drop casting of peptide solution, grids were then stained with uranyl acetate (20 μL) for 2 min. Excess stain was removed, and the grids were allowed to dry for overnight. Images were taken using JEOL Transmission Electron Microscope (Tokyo, Japan) maintained at 80 kv.

3.4. Scanning Electron Microscopy (SEM)

SEM sample was prepared by drop casting a 2 mM aqueous solution (10 μL) onto the mica surface. Mica surface was allowed to dry at room temperature for overnight and analyzed in a Nova NanoSEM 450 using the Directional Backscatter (DBS) Electron Detector. All samples were imaged un-coated, in high vacuum mode.

3.5. Circular Dichroism

CD spectra were recorded on a JASCO J-810 spectropolarimeter using 1 mm path length cuvettes. The scan speed was 100 nm/min, and spectra were averaged over 8 scans. All samples were tested at 25 °C. The sample concentration was 100 μM in deionized water. The CD for background reference (water) was measured and subtracted from the sample.

3.6. Dynamic Light Scattering (DLS)

Dynamic light scattering and zeta potential experiments were conducted on a Malvern Instruments Zetasizer Nano ZS (Worcestershire, UK). The machine was calibrated using a 60 nm polystyrene standard. Each sample (1 mL, 100 μ M) was loaded into a cell, and the particle size and zeta potential were measured simultaneously in triplicate. To determine the size distribution, 1 mL samples were placed in PCS1115 glass cuvettes and DLS was performed at 25 $^{\circ}$ C at 173 $^{\circ}$ backscatter angle with 120 s equilibration time. The intensity-based z-averaged hydrodynamic diameters reported based on 11 scans. To measure zeta potential via electrophoretic mobility, the samples were added to a cuvette equipped with copper chip. The measurement was conducted at 25 $^{\circ}$ C and 3 cycles were performed (30 data points per cycle). DLS and zeta potential were conducted with 100 μ M [WR]₄ as a stock solution.

3.7. Synthesis of Silver Nanoparticles

A stock solution of silver nitrate was mixed with NaBH₄ in the presence of cyclic peptides. Typically, 100 μ L of 1 mM peptide solution was mixed with 100 μ L of 2 mM silver nitrate solution at room temperature. Then 100 μ L of 8 mM solution of NaBH₄ was added into the mixture to yield colloidal silver with a light yellowish color. The UV-Vis spectra of the solution were taken using SpectraMax M2 fluorescence spectrophotometer (Molecular Devices, CA) at different time intervals.

3.8. Enzyme Activity Assay

The activity of GAPDH enzyme was measured using KDalert GAPDH assay kit (Ambion, Austin, Texas). Briefly, 1 μ L of GAPDH enzyme was mixed with 99 μ L PBS (pH 7.0) in an eppendorf tube to create control enzyme solution of activity 0.26 U/ml. Same proportion of enzyme was mixed with different concentration of peptide solution (100 μ M, 500 μ M and 1 mM) in PBS. Then 10 μ L of enzyme solutions from each tube was taken in 96 well plates followed by the addition of 90 μ L KDalert master mix into it using multichannel pipettor. To determine the activity of the enzyme, fluorescence of the solutions was measured (Excitation: 560 nm, Emission: 590 nm). The reduction of NAD⁺ to NADH was continuously monitored on a SpectraMax M2 fluorescence spectrophotometer (Molecular Devices, CA) for 30 min at room temperature. Fluorescence measurements of NADH production with control enzyme solution and enzyme in 100 μ M and 500 μ M peptide solutions were carried out in triplicate. For thermal stability study, the fluorescence of the solutions was measured for 60 min at 50 $^{\circ}$ C.

3.9. Molecular Modeling

Ab initio Hartree Fock molecular orbital calculations were performed at the 6-31G basis set. Geometry of monomers was completely optimized (all bond lengths, bond angles and dihedral angles were allowed to relax without any constraint). Intermolecular interaction calculations using supermolecule approach were utilized to study stacking efficiency and interaction with counterions present in the reaction medium. Stacking efficiency = $E_{\text{stacked supermolecule}} - (E_{\text{monomer}} \times \text{no. of monomers} + E_{\text{counterion}} \times \text{no. of counterions})$. Molecular modeling studies without anion were conducted in the presence of a 1-palmitoyl-2-oleoyl-sn-glycero-3-phosphocholine (POPC) membrane bilayer using SPC water model. Simulation was performed at 300 K, and NPT ensemble was taken using Martyna-Tobias-Klein Barostat method. Molecular dynamics simulations were performed by Shaw's Desmond software. Membrane was set up by system builder module. Then water was added. Whole system was subjected to simulated annealing followed by molecular dynamics at room temp for 5 ns.

3.10. Fluorescence Spectroscopy

The steady-state fluorescence spectra were taken by a SpectraMax M2 fluorescence spectrophotometer (Molecular Devices, CA) in 96 well plate. The fluorescence of tryptophan residues in samples were measured using excitation and emission wavelength at 280 nm and 360 nm respectively. The measurement was performed by excitation and emission slits with a nominal band pass of 1 nm.

3.11. ATR-FTIR

FT-IR of [WR]₄ was carried out using Thermo-Nicolet 380 FT-IR spectrophotometer using OMNIC software. ATR-FTIR was used to collect the spectra of the sample.

4. Conclusions

In summary, a new class of surfactant-like cyclic peptides was reported. Among all the designed peptides, [WR]_n (n = 3-5) generated self-assembled vesicle-like nanostructures at room temperature as shown by TEM, SEM, and/or DLS. A plausible mechanistic insight of self-assembly of [WR]₅ was investigated by molecular modeling studies. Modified [WR]₅ analogues, such as [W_{Me}R]₅, [WR_{(Me)2}]₅, [W_{Me}R_{(Me)2}]₅, and [WdR]₅ exhibited different morphologies than [WR]₅ as shown by TEM. Surfactant-like cyclic peptide [WR]₅ exhibited significant stabilizing effect for generated silver nanoparticles and GAPDH activity. The data suggested that surfactant cyclic peptides, [WR]₅ and [W_{Me}R]₅, could be promising materials for stabilization of silver nanoparticles. These studies established a new class of surfactant-like cyclic peptides that self-assembled into nanostructures and could have potential application for stabilizing of silver nanoparticles and protein biomolecules.

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