High-Efficiency Fluorescence through Bioinspired Supramolecular Self-Assembly

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ABSTRACT: Peptide self-assembly has attracted extensive interest in the field of eco-friendly optoelectronics and bioimaging due to its inherent biocompatibility, intrinsic fluorescence, and flexible modulation. However, the practical application of such materials was hindered by the relatively low quantum yield of such assemblies. Here, inspired by the molecular structure of BFPms1, we explored the "self-assembly locking strategy" to design and manipulate the assembly of metal-stabilized cyclic (L-histidine-D-histidine) into peptide material with the high-fluorescence efficiency. We used this bioorganic material as an emissive layer in photo- and electroluminescent prototypes, demonstrating the feasibility of utilizing self-assembling peptides to fabricate a biointegrated microchip that incorporates eco-friendly and tailored optoelectronic properties. We further employed a "self-encapsulation" strategy for constructing an advanced nanocarrier with integrated in situ monitoring. The strategy of the supramolecular capture of functional components exemplifies the use of bioinspired organic chemistry to provide frontiers of smart materials, potentially allowing a better interface between sustainable optoelectronics and biomedical applications.

KEYWORDS: supramolecular fluorescence, cyclic dipeptide, self-assembly, optoelectronic, drug release monitor

Green fluorescent protein (GFP) has been extensively used as a genetically encoded fluorescent marker in biology. Specifically, a GFP mutant (BFPms1) that preferentially binds to Zn(II) has been developed. Zn(II) binding rigidifies the chromophore imidazole, reduces fluorophore mobility, and further limits the energy dissipation through thermal relaxation pathways, resulting in the enhanced fluorescence (Figure 1a).24 Inspired by this molecular structure, we aimed to design a metal-binding site on a short peptide to simulate metal coordination and design an electrostatic interaction site to generate supramolecular hosts of a β-fold barrel environment for self-locking, thereby fabricating the minimalist version of this sophisticated biological structure to provide a scalable technological solution (Figure 1b).14 Cyclic peptides derived from amino acid residues carrying complexing side chain substituents, such as imidazole, carboxylate, or thioether...
groups, can be used as models to mimic the coordination of metal ions in enzymes.\textsuperscript{25–30} Also, cyclic dipeptide is highly tunable due to hydrogen bonding capabilities of the skeleton and other noncovalent interactions that can be used to engineer artificial multifunctional scaffolds.\textsuperscript{31,32}

Here, we explored the assembly of cyclic(\textit{L}-histidine-\textit{D}-histidine) (CHH) and constructed highly fluorescent peptide dots with a large quantum yield (>0.7) through “self-assembly locking strategy”. As a proof of concept, we demonstrated CHH self-assemblies to show bright fluorescence, allowing their use as an emissive layer in the photo- and electro-luminescent light-emitting diodes (LEDs). Moreover, we utilized the “self-encapsulation” strategy to construct a nanocarrier to effectively deliver an anticancer drug into cancer cells with \textit{in situ} monitoring. Our studies show that, analogously to diphenylalanine, CHH is a type of pioneering minimalistic self-assembling peptide, which shows that bioinspired supramolecular functional components can be applied as multifunctional nanomaterials with exceptional features for optoelectronic or biological applications.

RESULTS AND DISCUSSION

To allow their coassembly, CHH and Zn(NO\textsubscript{3})\textsubscript{2} (CHH–Zn) were mixed under controlled experimental conditions, resulting in nanostructure formation. Atomic force microscopy (AFM) and transmission electron microscopy (TEM) imaging confirmed the presence of nanoparticles with an average diameter of \textsim 30 nm (Figures 1c and S1) which is in agreement with dynamic light scattering (DLS) data (Figure S2). Next, we explored the optical properties of the CHH–Zn nanostructures. Figure 1d shows the normalized UV–vis spectra of CHH–Zn, CHH, and Zn(NO\textsubscript{3})\textsubscript{2} (Inset: CHH–Zn under daylight (left) and UV lamp (365 nm right)). (e) Excitation–emission matrix contour profiles of CHH–Zn. (f) Comparison of the quantum yield of different fluorescent biometabolites with CHH–Zn marked with a green star (vitamin, riboflavin, \textalpha-tocopherol; amino acid, Phe, Tyr, Trp, Ala, His, Lys, Ser; Peptide, Trp-Phe+Zn(II), cyclo-WW+Zn(II), Phe-Phe-Phe,\textsuperscript{20,38,39} GFP, sg11, avGFP, sapphire, H9). (g) \textit{1}H chemical shifts of CHH–Zn, compared to the peptide alone. (h) Job plot analysis of CHH with Zn(NO\textsubscript{3})\textsubscript{2}.

Figure 1. Design of self-assembled fluorescent cyclic dipeptide by the biologically inspired molecular structure of BFPms1. (a) BFPms1 overall structure and its coordination geometry with Zn(II). (b) Scheme of self-assembly mechanism of CHH, Zn(II), and NO\textsubscript{3}-. (c) AFM image showing the presence of \textsim 30 nm nanoparticles. Scale bar = 400 nm. (d) Normalized UV–vis spectra of CHH–Zn, CHH, and Zn(NO\textsubscript{3})\textsubscript{2}. (Inset: CHH–Zn under daylight (left) and UV lamp (365 nm right)). (e) Excitation–emission matrix contour profiles of CHH–Zn. (f) Comparison of the quantum yield of different fluorescent biometabolites with CHH–Zn marked with a green star (vitamin, riboflavin, \textalpha-tocopherol; amino acid, Phe, Tyr, Trp, Ala, His, Lys, Ser; Peptide, Trp-Phe+Zn(II), cyclo-WW+Zn(II), Phe-Phe-Phe,\textsuperscript{20,38,39} GFP, sg11, avGFP, sapphire, H9). (g) \textit{1}H chemical shifts of CHH–Zn, compared to the peptide alone. (h) Job plot analysis of CHH with Zn(NO\textsubscript{3})\textsubscript{2}.
comparable to inorganic quantum dots or GFP.35 Moreover, a Job plot analysis of the CHH−Zn(II) coordination of Zn ion and imidazole ring was further verified. 

Figure 2. Structure analysis of the CHH−Zn assemblies. (a) Single-crystal structure of CHH−Zn(II) in Pbcn space group. Color scheme: gray, C; red, O; blue, N; green, Zn; and purple, I. (b) Single-crystal structure of CHH−NaNO₃ in P2₁/c space group. (c) Four snapshots from Movie S1 demonstrating the assembly kinetics of the CHH−Zn(NO₃)₂ single crystal. (d) PXRD pattern of a CHH−Zn(II) single crystal. (e) FLIM studies of the CHH−Zn(NO₃)₂ single crystal. (f) Lifetime distribution histogram derived from the FLIM study, showing two peaks centered at 1318 and 1890 ps. Scale bar is 50 μm.

Such red shift in the fluorescence emission spectra in response to a change in the excitation wavelength is termed as red edge excitation shift (REES).13,34 Notably, the maximum photoluminescence efficiency of the CHH−Zn self-assembly was about 70.6% (Figures 1f and S7), among the highest values reported so far for peptide-derived materials and even comparable to inorganic quantum dots or GFP.35−38

We further combined spectroscopic methods and control experiments to obtain specific chemical and structural information. We found that all Zn(II) present in the assembled system displayed a strong and stable fluorescence signal and similar absorption and emission spectra, whereas the sodium nitrate-related self-assembly system and CHH showed similar absorption spectra and weak fluorescence emission intensity (Figures S4 and S8). Therefore, it can be inferred that the Zn(II)-peptide coordinated structure is formed by supplying Zn(II) to the cyclic dipeptide system, which in turn determines the optical properties. NMR was further used to predict the possible coordination structure. As shown in Figure 1g and Figure S9, the imine protons of the side-chain imidazole ring downshifted Δδ = 0.143 ppm (a position) after the addition of Zn(II), implying strong coordination through imine−imidazole nitrogen.40 This coordination of the CHH imidazole ring with Zn(II) is also consistent with previous studies of carbonic anhydrase and β-amyloid peptide, where the histidine residue has a high affinity for the complexation of Zn.13,41,42 The coordination of Zn ion and imidazole ring was further verified by mass spectrometry analysis of CHH with Zn(NO₃)₂, showing an m/z 611.2 band corresponding to the oligomer of [2M_CHH + Zn²⁺] (Figure S10). Moreover, a Job plot analysis determined the CHH/Zn(NO₃)₂ stoichiometric ratio as 2:1, which is in contrast to the 1:1 stoichiometric ratio of CHH/Zn(II) (Figure 1h and Figure S11), thus indicating the presence of another assembled structure in addition to the Zn−peptide coordination. We therefore propose a specific self-assembly mechanism mediating CHH packing with nitrate.

To validate our hypothesis and further characterize the specific self-assembly mechanism, we crystallized both CHH−Zn(II) and CHH−NaNO₃ and analyzed the resulting structures via X-ray crystallography (Table S1). The CHH−Zn(II) crystallizes in orthorhombic space group Pbcn with one CHH molecule, one neutral [Zn(L)₂I₂] unit, and one isopropanol molecule per asymmetric unit. A perspective view of the Zn(II) center of the CHH−Zn(II) compound is illustrated in Figure 2a with a unit cell scheme. Each Zn(II) atom was coordinated with two ligands and two N-donor atoms from the imidazole groups of two different CHH molecules, occupying the apical coordination sites to generate a Zn(II) centered geometric tetrahedron. In turn, two adjacent cyclic dipeptides were connected through a β-bridgelike hydrogen bonding on the opposite sides of the backbone. The X-ray determined structure of CHH−NaNO₃ revealed a packing of the cyclic-dipeptides crystal in the monoclinic space group P2₁/c with two CHH and four nitrates in the unit cell (Figure 2b). The components assembled to form a 1D chain with a hydrogen bond (N−H···O=C) of 2.889 Å (donor−acceptor) via a parallel β-sheet hydrogen bonding network. The adjacent chains formed the extended structure through hydrogen bonds between the imidazole ring and nitrate groups.

We further examined CHH−Zn(NO₃)₂ single crystals through crystallographic analysis. By employing a microfluids technique (Figure 2c, Figures S12, S13, and Movie S1), it is possible to visually observe that the Zn(NO₃)₂ crystals are densely packed with a growth rate of 0.01 μm s⁻¹ along the a direction, ultimately forming a needle shape. The resulting powder X-ray diffraction (PXRD) pattern and unit cell...
parameters of the CHH–Zn self-assemblies highly resembled those of the formed CHH–Zn(NO3)2 crystals, indicating a similar molecular organization (Figure 2d and Figure S14). Surprisingly, the needle-shaped CHH–Zn(NO3)2 and the plaque-shaped CHH–NaNO3 crystal are isomorphous despite different crystal morphologies. Importantly, the Zn ion is not incorporated in the CHH–Zn(NO3)2 single crystal structure, suggesting that the growth of the CHH–Zn(II) oligomer may be mostly restricted by the CHH–nitrate self-assembly process. To validate our hypothesis and elucidate the effect of the CHH–Zn(II) oligomer on the CHH–Zn(NO3)2 crystal photodynamic properties, we performed confocal fluorescence lifetime microscopy (FLIM) studies of CHH–Zn(NO3)2 single crystals (Figure 2e). Figure 2f presents the lifetime distribution histogram using a multiexponential fit of the decays. Interestingly, a highly heterogeneous lifetime distribution could be observed for the CHH–Zn(NO3)2 single crystal, and the envelope of lifetime (τ) histogram could be divided into two independent fluorescence decay processes with two clearly differentiated Gaussian distributions centered at 1318 and 1890 ps. This data confirmed that the two lifetime values originated from the fluorescence decay of the two individual components, which could be assigned to the fluorescence emission within the CHH–Zn(II) oligomer and CHH–nitrate.43 In addition, CHH–Zn(NO3)2 forms yellow needle crystals, whereas CHH–NaNO3 gives rise to colorless transparent plaque crystals and the fluorescence emission intensity of CHH–Zn(NO3)2 is much higher than that of CHH–NaNO3 (Figures S15 and S16).
We computationally investigated the self-assembly of the two systems, independently comprising \( \text{CHH}^\text{−} \text{Zn(NO}_3\text{)}^2 \) and \( \text{CHH}^\text{−} \text{ZnCl}_2 \), in isopropanol, using multiple explicit solvent molecular dynamics (MD) simulations in CHARMM.\textsuperscript{44} In both systems, we observed the gradual formation of dimer structures having two \( \beta \)-bridge-bonded \( \text{CHH} \) coordinating with one \( \text{Zn(II)} \) ion (Figure S17), similar to the X-ray crystallography data, and the gradual formation of aggregate clusters formed by the constituent elements of each system. In line with the experimental findings of the Job plot analysis (Figure 1h), mature clusters in the \( \text{CHH}^\text{−} \text{Zn(NO}_3\text{)}^2 \) simulations showed a \( \text{CHH/Zn(NO}_3\text{)}^2 \) ratio of approximately 2:1, whereas mature clusters in the \( \text{CHH}^\text{−} \text{ZnCl}_2 \) simulations displayed a \( \text{CHH/ZnCl}_2 \) ratio of approximately 1:1 (Table S2). Within both sets of simulations, \( \text{CHH} \) molecules were observed to form antiparallel \( \beta \)-bridge conformations with the imidazole rings of two histidine molecules of opposing \( \text{CHH} \) dipeptides coordinated with the same \( \text{Zn(II)} \) ion, similar to the conformations observed in the \( \text{CHH}^\text{−} \text{Zn(II)} \) crystal structure of (Figures 2a and 3a and Table S3). In the simulations comprising \( \text{CHH}^\text{−} \text{Zn(II)} \) and \( \text{NO}_3\text{)}^− \), these antiparallel conformations were predominantly observed within the interior \( \text{CHH}^\text{−} \text{Zn(II)} \) nucleus of the clusters (Figure 3d). We also observed that the radius of gyration of \( \text{Zn(II)} \) within the clusters formed in the presence of \( \text{NO}_3\text{)}^− \) was lower compared to those formed in the absence of \( \text{NO}_3\text{)}^− \), showing that for clusters containing the same number of \( \text{CHH} \) molecules \( \text{Zn(II)} \) ions were more densely packed and concentrated in the clusters formed in the presence of \( \text{NO}_3\text{)}^− \) (Figure 3b). Importantly, in the clusters formed by \( \text{CHH}^\text{−} \text{Zn(NO}_3\text{)}^2 \) our computational analysis detected layers in which the exterior surface layer was composed of primarily \( \text{CHH} \) and \( \text{NO}_3\text{)}^− \), whereas the interior layer was composed primarily of \( \text{CHH} \) and \( \text{Zn(II)} \) (Figure 3c).

We further aimed to provide insight into the formation of the \( \text{CHH}^\text{−} \text{Zn(NO}_3\text{)}^2 \) clusters by performing free-energy analysis of the different pathways which may lead to their formation. On the basis of the free-energy analysis, \( \text{CHH} \) and \( \text{Zn(II)} \) coordinate first, forming the interior of the clusters, followed by individual pieces of the \( \text{CHH} \) and \( \text{NO}_3\text{)}^− \) exterior wrapping around the preformed interior (Figures 3d and S18a). This mechanism was supported through structural analysis in which the composition of the large \( \text{CHH} \) clusters was tracked as a function of time (Figure S19a). Thus, \( \text{CHH}^\text{−} \text{Zn(NO}_3\text{)}^2 \)
Zn(NO$_3$)$_2$ self-assembles through a "self-assembly locking strategy" in which the fluorescent CHH–Zn(II) nucleus is encased by the CHH–NO$_3$$_2$ scaffold. Figure 3d and Movie S2 show the initial formation of the internal CHH–Zn(II) nucleus, followed by the exterior CHH–NO$_3$$_2$ wrapping around the preformed nucleus. Our combined experimental and computational analysis suggests that the CHH–Zn(II) oligomer is encapsulated into CHH–NO$_3$$_2$ assemblies and that immobilization of the peptide oligomers by a high-stiffness scaffold would limit energy dissipation during thermal relaxation pathways for better quantum yield and fluorescence intensity.

On the basis of our experimental and computational analyses, a plausible self-assembly mechanism of CHH and Zn(NO$_3$)$_2$ is depicted in Figure 3e. The self-assembly of the CHH and Zn(II) can be observed at initial oligomerization step. Following the coordination of Zn(II) with the two histidine side-chains and stabilization of the dimer, the CHH monomers begin to form hydrogen bond interactions between their backbone atoms, forming a one-dimensional chain via β-sheet bridgelike interactions and subsequently generating an extended network through the linkage of nitrates. As the CHH one-dimensional chain grows, the chelation of CHH and Zn(II) is limited. Finally, the CHH–Zn(II) oligomer clusters are encapsulated and incorporated into CHH–NO$_3$$_2$ nanoassemblies.

Intrigued by the optical properties of the characterized assemblies, we sought to study their capability to serve as an emissive material in photo- and electroluminescent prototypes. As shown in Figure 4a, we prepared peptide-based phosphors by embedding CHH–Zn into polyvinylpyrrolidone (PVP) at the mass ratio of 1:70. The peptide-based phosphor converted LED emitted bright green light with Commission Internationale de L’Eclairage (CIE) color coordinates of (0.31, 0.45) and achieved high luminous efficiency of 56.62 lm W$^{-1}$ at 20 mA drive current (Figure S21). We further sought to study the utilization of CHH–Zn as a bio-organic light-emitting material in optoelectronics. A simple natural peptide derived bio-organic-Lessing (OLED) prototype was fabricated by using CHH–Zn-blended poly(N-vinyl carbazole) (PVK) as an emissive layer (Figures 4b and S22). As illustrated in Figure 4b, the operation photographs present a close-up view of the bright, uniform, and defect-free surface green electroluminescence emission from the peptide-based OLED. The typical luminance and current density curves as a function of the applied voltage for the OLED are demonstrated in Figure S23. The maximum EQE is 0.25%, corresponding to a current efficiency of 0.58 cd A$^{-1}$ (the corresponding $L_{\text{max}}$ is 1385 cd m$^{-2}$) (Figure S24). Because of the stable fluorescence, the bio-OLED showed no temporal degradation in the emission spectrum under the applied operating conditions, indicating significant potential for practical applications.

Because self-assembled peptide nanoparticles are composed of naturally occurring amino acids with inherent biocompatibility, peptide self-assembly with intrinsic fluorescence is suitable for bioimaging. High-resolution confocal fluorescence microscopy images of HeLa cells were collected following incubation with CHH–Zn and the DRAQ5 red DNA stain. The CHH–Zn structures were found to penetrate the cells and display bright green fluorescence under excitation of 405 nm (Figure S25). More notably, three-dimensional (3D) imaging analysis indicated that CHH–Zn could effectively transport through the nuclear pore complex of HeLa cells and accumulate within the nucleolus region (Figures S26 and S27). Also, in vitro cytotoxicity analysis showed in Figure S28 demonstrated the excellent cytocompatibility of CHH–Zn peptide nanoparticles toward HeLa cells. On the basis of the membrane permeability feature of the developed intrinsically fluorescent peptide structures, we further aimed to study their potential applications for drug delivery.

We experimentally confirmed the coassembly of CHH–Zn and Epirubicin, an anthracycline drug used for chemotherapy, through absorbance spectra, showing a 15.67% loading capacity of Epirubicin within the CHH–Zn nanoassemblies. We computationally investigated the self-assembly properties of CHH, Zn(NO$_3$)$_2$, and Epirubicin in water using multiple explicit solvent runs using CHARMM. In all simulations, we detected the gradual formation of clusters comprising CHH, Zn(II), NO$_3$$_2$, and Epirubicin (Figure S19d) in which Epirubicin and Zn(II) were primarily located in the interior nucleus whereas CHH and NO$_3$$_2$ were primarily at the exterior surface (Figures 4d, S19c, and S20), thereby depicting the self-encapsulation properties of the system. Free energy calculations (Figure S18b–d) and structural analysis (Figure S19c) show that individual or pairs of CHHs pulled Zn(II) from the solvent environment into a more peptide-like environment through an "environment-switching" mechanism. The Epirubicin molecules first aggregate and form the inner nucleus of the clusters which further facilitates the assembly of individual pieces of CHH and NO$_3$$_2$ exteriorly wrapping around the preformed Epirubicin (Figure 4 panels d ($\Delta G_f$) and e, and Movie S3).

To examine the drug delivery potential of the nanoassemblies, HeLa cells incubated with CHH–Zn+Epirubicin or Epirubicin alone were examined by live-cell confocal microscopy. The fluorescence intensity of intracellular Epirubicin in cells incubated with CHH–Zn+Epirubicin was significantly higher than that of Epirubicin alone, indicating efficient Epirubicin uptake and release into the nucleolus of HeLa cells via the CHH–Zn carrier (Figure 4e). The Epirubicin release profiles (Figures S29 and S30) suggested that the release of Epirubicin from the CHH–Zn can be efficiently triggered and accelerated by an acidic stimulus, which is favored for the acidic extracellular microenvironment of tumor tissues. In order to further monitor the Epirubicin release process and eliminate autofluorescence from the biological system, we applied the two-photon FLIM technique with phasor analysis. Pixels with similar lifetimes are selected in the phasor diagram and the FLIM image is separated and painted into four subcellular compartments: cell membrane (~3512 ps), cytoplasm (~2286 ps), nucleus membrane (~1595 ps), and nucleus (~1261 ps) (Figure S31). After internalizing of Epirubicin into the cells, changes in its fluorescence lifetime can thus indicate changes in the subcellular microenvironment, reflecting drug release and transport. With elongation of incubation time (Figure 4g), more Epirubicin was released and, consequently, the fluorescence intensity of Epirubicin gradually increased, along with a decrease in the average lifetime. These results indicated that the CHH–Zn+Epirubicin could accumulate around and bind to the cell membrane as early as 35 min of incubation with HeLa cells and then be released in the cytoplasm due to the acidic environment and eventually accumulate in the nucleus. In addition, the release behavior of CHH–Zn+Epirubicin could be monitored by the variation of the
fluorescent signal of CHH–Zn (Figure S32), showing that CHH–Zn not only promoted the transport of Epirubicin into HeLa cells but also can be acted as a real-time optical monitor for the drug release process. Thus, the fluorescence of peptide nanostructures can be used to investigate the drug release in spatiotemporal mode and metabolism kinetics of cancer drugs in a certain organ or tissue.

CONCLUSION
Within this study, inspired by the structure of BFPms1, we successfully constructed a fluorescent short peptide core encapsulated by the peptide scaffold building module to implement the concept of “self-assembly locking strategy”. We report the demonstration of a bright fluorescent peptide with quantum yields of up to 70% for green fluorescence, exemplifying the potential of such structures to serve as bioinspired, organic, supramolecular alternatives to complement the state-of-the-art inorganic counterparts. As a proof of concept, we demonstrate the utilization of these bright fluorescence peptide self-assemblies for eco-friendly optoelectronics and bioimaging. In particular, we show their capability to serve as an emissive layer for a bioinspired OLED prototype. Moreover, we employ the “self-encapsulation” strategy for fabricating an advanced nanocarrier for traceable intracellular drug delivery. These results allow us to envision CHH as a promising platform for further therapeutic or diagnostic applications, particularly in anticancer treatments. This efficient yet straightforward bottom-up approach for the design of highly efficient fluorescent peptide nanostructures may represent a strategy for developing peptide-based advanced nanomaterials.

METHODS
Atomic Force Microscopy. Five microliters of sample solution was dropped onto a freshly cleaved mica surface and dried by N2 purge (99.99%). A topographic image was recorded underwater a Dimension icon AFM (Bruker) in the tapping mode at ambient temperature with a 512 × 512-pixel resolution and a scanning speed of 1.0 Hz.

UV–vis Spectra. UV–vis spectra between 200 to 800 nm were recorded on an Agilent Cary 100 UV–vis spectrophotometer with a quartz cuvette of 1 mm path length.

Fluorescence Spectroscopy and Quantum Yield (QY) Measurement. Six hundred microliters sample solution was pipetted into a 1.0 cm path-length quartz cuvette, and the spectrum was recorded on an Agilent Cary 100 UV–vis spectrophotometer with a quartz cuvette of 1 mm path length.

Microfluidics Experiments. In a typical protocol, CHH–Zn(NO3)2 crystalline powder was inserted into the device. Then, a flow of fresh solutions was injected at a rate of 4 μL h−1 using Cetoni GmbH neMESYS Syringe Pumps (Korbussen, Germany) and glass HAMILTON LTL syringes, 1,725 μL of 250 μL. The process was examined under an Eclipse Ti-E inverted microscope (Nikon, Japan), equipped with a Zyla 4.2+ sCMOS camera (Andor, U.K.), and images were captured at different time points.

X-ray Crystallography. Crystals suitable for diffraction were coated with Paratone oil (Hampton Research), mounted on loops and flash frozen in liquid nitrogen. Single crystal X-ray diffraction data measurement was performed using a Rigaku XtaLabPro system with CuKα (λ = 1.5418 Å) radiation at 100(2) K. Data were collected and processed using CrysolisPro 1.171.39.22a (Rigaku OD, 2015). The structure was solved by direct methods using SHELXT-2016/4 and refined by full-matrix least-squares against F2 with SHELX-2013.

Photoluminescence Device Fabrication and Characterization. Commercially available InGaN chips were used at the bottom of the light-emitting diode (LED) base. For preparation of the color conversion layer, the CHH–Zn was blended into PVK at a mass ratio of 1:70, and the resulting mixtures were vacuum-dried at 60 °C for 30 min. The mixtures were applied on the InGaN chips and, following curing at 80 °C for 1 h, the LEDs peptide phosphors were obtained.

Organic LED Device Fabrication and Characterization. ITO-coated glass substrates were cleaned ultrasonically in organic solvents (acetone and isopropyl alcohol), rinsed in deionized water, and then dried in an oven at 150 °C for 10 min. The substrates were cleaned by a UV-ozone treatment to enrich the ITO surface with oxygen, thereby increasing its work function. The approximately 30 nm thick PEDOT:PSS hole injection layer was spin-coated at 3000 rpm for 30 s on the ITO, followed by annealing in an oven at 150 °C for 15 min. Subsequently, the emissive layer of CHH–Zn blended into PVK was spin-coated at 3000 rpm for 35 s over the surface of the PEDOT:PSS film from the solution of NMP, followed by baking on a hot plate at 80 °C for 15 min to form the active region of the peptide-derivated bio-OLED. Finally, the substrates were transferred to a vacuum chamber, and a 30 nm thick TPBI electron transport layer was thermally deposited with base pressure of 3 × 10−6 Pa. Next, a 20 nm Ca and 100 nm thick Al cathode was deposited using a shadow mask 2 mm in width. The active area of the devices was thus 4 mm2. The thermal deposition rates for TPBI and Ca/Al were 1, 1, and 3 Å s−1, respectively. The thickness of the films was measured using a Dektak XT (Bruker) surface profilometer and a spectroscopic ellipsometer (Suntech). The luminance–current–voltage (L–I–V) characteristics were measured using a computer-controlled Keithley 236 SMU and Keithley 200 multimeter coupled with a calibrated Si photodiode. Electroluminescence spectra were measured by an Ocean Optics 2000 spectrometer, which couples a linear charge-coupled device array detector ranging from 350 to 800 nm.

Live Cells Imaging Using Confocal Microscopy. HeLa cells were grown to 70–80% confluence in glass bottom cell culture dishes. Then, the cells were cultured with media containing the CHH–Zn + EPI at a concentration of 4 μg/mL for different durations. Next, the cells were stained using a DRAQ5 dye diluted 1:1000 in PBS for 15 min at room temperature in the dark to allow staining of the nuclei. The cells were then washed twice with PBS. Imaging was performed using SP8 inverted confocal microscope (Leica Microsystems, Wetzlar, Germany). Excitation and emission ranges were the following: λex = 405 nm, λem = 420–500 nm; EPI, λex = 543 nm, λem = 550–750 nm; DRAQ5, λex = 633 nm, λem = 750–780 nm.

FLIM Analysis of Cultured Cells. HeLa cells seeded in dishes were treated with CHH–Zn+EPI at a concentration of 4 μg/mL for 30, 76, 125, 194, 270, and 420 min, followed by washing with PBS. The time-resolved fluorescence signal was acquired using an LSM 7 MP two-photon microscope (Carl Zeiss, Weimar, Germany). Excitation and emission ranges were the following: λex = 405 nm, λem = 420–500 nm; EPI, λex = 543 nm, λem = 550–750 nm; DRAQ5, λex = 633 nm, λem = 750–780 nm.

ASSOCIATED CONTENT
Supporting Information
The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/acs.nanolett.9b01024.

Additional experimental details, including supplementary experimental methods, TEM image, DLS result, UV–vis
spectra, excitation–emission matrix contour profiles, photostability evaluation, NMR result, mass spectra, crystal morphology, XRD pattern, fluorescence microscopy images; MD simulations result: (molecular graphics image of the coordination-induced self-assembly of CHH with Zn(II), free energy of associate the clusters, time evolution percent composition of clusters); characterization of LED and OLED; confocal microscopy images, cytotoxicity result, in vitro drug release profiles, Phasor–FLIM analysis of HeLa cells cultured with CHH–Zn(NO₃)₂ (PDF)

Growth of CHH–Zn(NO₃)₂ single crystal by microfluidic (MP4)

MD simulation of CHH–Zn(II)–NO₃ (MOV)

MD simulation of CHH–Zn(II)–NO₃+EPI (MOV)

Crystallographic data (CIF)

Crystallographic data (CIF)

Crystallographic data (CIF)

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