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Target-driven supramolecular self-assembly for selective amyloid- β photooxygenation against Alzheimer's disease†

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Photo-oxygenation of β -amyloid (A β) has been considered an efficient way to inhibit A β aggregation in Alzheimer's disease (AD). However, current photosensitizers cannot simultaneously achieve enhanced blood–brain barrier (BBB) permeability and selective photooxygenation of A β , leading to poor therapeutic efficacy, severe off-target toxicity, and substandard bioavailability. Herein, an A β target-driven supramolecular self-assembly (PKNPs) with enhanced BBB penetrability and switchable photoactivity is designed and demonstrated to be effective in preventing A β aggregation *in vivo*. PKNPs are prepared by the self-assembly of the A β -targeting peptide KLVFF and an FDA-approved porphyrin derivative (5-(4-carboxyphenyl)-10,15,20-triphenylporphyrin). Due to the photothermal effect of PKNPs, the BBB permeability of PKNPs under irradiation is 8.5-fold higher than that of porphyrin alone. Moreover, upon selective interaction with A β , PKNPs undergo morphological change from the spherical to the amorphous form, resulting in a smart transformation from photothermal activity to photodynamic activity. Consequently, the disassembled PKNPs can selectively oxygenate A β without affecting off-target proteins (insulin, bovine serum albumin, and human serum albumin). The well-designed PKNPs exhibit not only improved BBB permeability but also highly selective A β photooxygenation. Furthermore, *in vivo* experiments demonstrate that PKNPs can alleviate A β -induced neurotoxicity and prolong the life span of the commonly used AD transgenic *Caenorhabditis elegans* CL2006. Our work may open a new path for using supramolecular self-assemblies as switchable phototheranostics for the selective and effective prevention of A β aggregation and related neurotoxicity in AD.

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Introduction

Alzheimer's disease (AD), the most prevalent type of dementia, affects more than 50 million people worldwide. Even worse, with the aging of the population, the number of cases of AD will increase rapidly. Increasing evidence has suggested that the aggregation of amyloid- β peptides (A β) is a critical step towards AD pathogenesis.¹ Accordingly, prevention of A β aggregation has been sought as a promising strategy to treat AD.^{2,3} Recently, photo-oxygenation of A β has been used for the suppression of A β aggregation with unique merits of low invasiveness, high selectivity, and spatiotemporal controllability.^{4–6} Until now, numerous molecular photosensitizers (such as porphyrins,⁷ riboflavin,⁸ and thioflavin T^{9,10}) have been reported for the

inhibition of A β aggregation by photo-oxygenation of A β , but none have achieved satisfactory therapeutic effects. The lack of efficacy is mainly attributed to the blood–brain barrier (BBB) with well-structured and dense paracellular tight junctions, which routinely impedes the entry of most therapeutic drugs into the central nervous system (CNS).^{11,12} In addition, these molecular photosensitizers also tend to aggregate and/or suffer from rapid elimination in the body,¹³ further resulting in a decrease in photo-oxygenation efficiency. Most recently, several photoactive nanomaterials with unique BBB penetration ability^{14–16} and physicochemical stability have been developed as promising alternatives to molecular photosensitizers.^{17–19} However, these nanoscale photosensitizers can cause unwanted off-target oxidative damage to healthy tissues due to the reactive oxygen species (ROS) generated under illumination.^{6,10} Hence, development of a novel photodynamic therapy (PDT) strategy with improved brain bioavailability and controllable ROS generation is highly desired.

In nature, the self-assembly of biomolecules into complicated and functionalized units utilizing multiple noncovalent interactions, including electrostatic, hydrophobic, π – π , and coordination interactions, affords a rationale to construct

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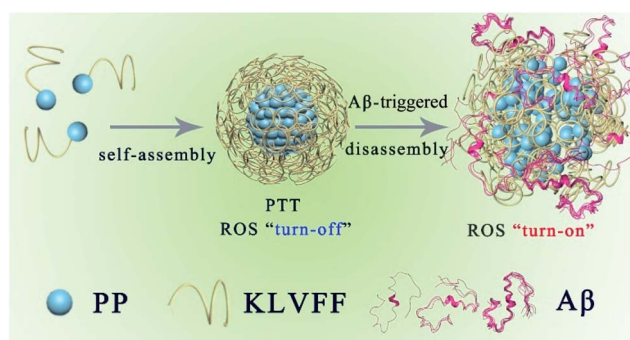
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nanostuctures.^{20–22} These reversible, controllable, and dynamic noncovalent interactions allow these self-assembled systems of biomolecules to adapt well to the physiological environment to fully optimize their biological functions.^{23–26} For example, the specific interaction among macromolecules causes a conformational change in the partner macromolecule to activate one of the binding partners to trigger a biological cascade.²⁷ In particular, low molecular weight peptides exhibit outstanding advantages (such as inherent biocompatibility, potential biodegradability, structural programmability, and easy preparation) compared to other existing self-assembly motifs.²⁸ Herein, a peptide-based porphyrin supramolecular self-assembly (PKNPs) with A β -responsive structural transformation was designed for selective photooxygenation of A β .

The PKNPs are prepared by the self-assembly of the US food and drug administration (FDA)-approved porphyrin derivative photosensitizer²⁹ (5-(4-carboxyphenyl)-10,15,20-triphenylporphyrin, PP) and A β -targeting peptide KLVFF.^{30,31} Porphyrin is selected as a photosensitizer due to its superior optical and electronic properties.^{13,32,33} Besides, porphyrin with intrinsically hydrophobic characteristics could function as a building block in the construction of supramolecular nanostructures. In the study, hydrophobic interactions and π - π stacking interactions facilitate porphyrin-peptide conjugate (PP-KLVFF) self-assembly into spherical nanostructured PKNPs and inhibit their fluorescence emission and ROS generation. Therefore, unsurprisingly, PKNPs exhibit an excellent photothermal effect under illumination, which is conducive to increasing their BBB permeability.^{34,49} More importantly, PKNPs permit disassembly upon specific interaction with A β , leading to smart transformation from photothermal activity to photodynamic activity.^{35,36} As a consequence, PKNPs achieve selective photooxygenation of A β without affecting non-specific proteins (insulin, bovine serum albumin (BSA), and human serum albumin (HSA)). In this way, PKNPs can realize both improved BBB permeability and highly selective photooxygenation of A β by the transformation of morphology (Scheme 1). Furthermore, PKNPs show no obvious toxicity both in rat pheochromocytoma (PC-12) cells and N2 wild-type strain worms. To the best of our knowledge, the use of a supramolecular self-assembly as a novel photosensitizer for activable PDT against AD has not been reported.



Scheme 1 Schematic illustration of the self-assembly process and A β -triggered disassembly process of PKNPs.

Results and discussion

KLVFF, a pentapeptide known to specifically target A β ,³⁷ was conjugated to hydrophobic PP to synthesize PP-KLVFF. The resulting PP-KLVFF was verified by ¹H nuclear magnetic resonance (¹H NMR, 600 MHz), Fourier transform infrared (FTIR) spectroscopy, mass spectrometry (MS), and high performance liquid chromatography (HPLC) (Fig. S1–S4†). Due to the amphiphilicity, PP-KLVFF could spontaneously assemble into nanostructured PKNPs in DMSO/H₂O (1 : 9, v/v) solution. Transmission electron microscopy (TEM) images showed that PKNPs possessed a spherical morphology (Fig. 1a). The dynamic light scattering (DLS) histogram of PKNPs revealed an average diameter of 70 nm (Fig. 1b). In addition, a distinct bathochromic shift and broadening of Soret and Q bands in the absorption spectra of PKNPs were observed (Fig. 1c), which could be ascribed to the face-to-face stacking of PP.³³ The corresponding shift for PKNPs was further investigated by circular dichroism (CD) spectroscopy. As shown in Fig. 1d, a distinct red-shift of the Soret band region in the CD spectrum of PKNPs was observed, which implied strong intermolecular interaction between PP molecules.^{38,39} As shown in Fig. 1e and f, the fluorescence emission and ROS generation of PKNPs were completely quenched, implying that porphyrin molecules were aggregated in aqueous solution. These results together confirmed the successful construction of PKNPs. More

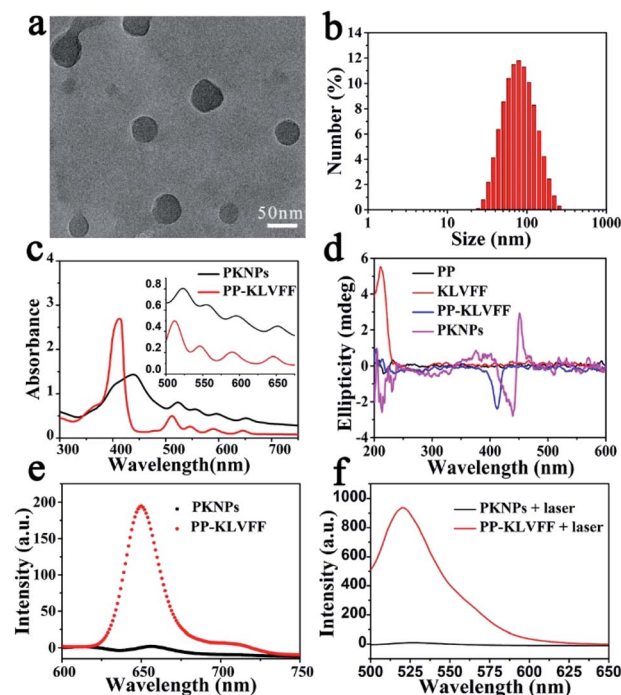
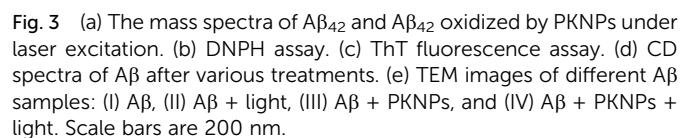
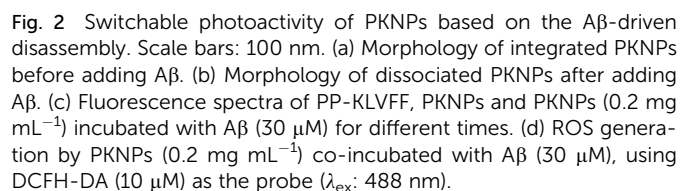


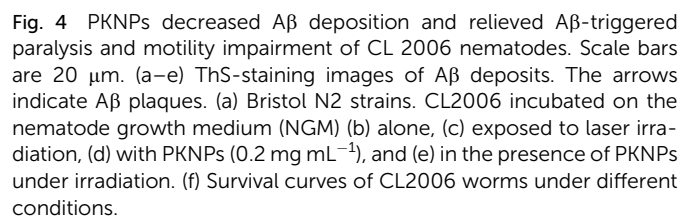
Fig. 1 Self-assembly of PP-KLVFF into PKNPs. (a) TEM image of PKNPs. (b) DLS histogram of PKNPs. (c) UV-vis absorption spectra of PP-KLVFF and PKNPs. The inset shows the amplified absorption spectra of Q bands. (d) CD spectra of PP, KLVFF, PP-KLVFF, and PKNPs. (e) Fluorescence emission spectra (λ_{ex} : 450 nm) of PKNPs and PP-KLVFF. (f) ROS generation by PKNPs and PP-KLVFF, using DCFH-DA (10 μM) as a probe (λ_{ex} : 488 nm).

Since the self-assembly is based on the synergy of non-covalent interactions, it is easily subject to environmental variations.^{36,43–45} Considering the high binding affinity of A β with KLVFF, A β deposition in the brain of AD patients could trigger the disassembly of PKNPs. To test this hypothesis, the morphology change of PKNPs was first detected by TEM. As shown in Fig. 2a and b, the morphology of PKNPs changed from regular solid spheres to amorphous and fluffy particles upon the addition of A β . This result suggested that A β could induce PKNP disassembly, which was attributed to the fact that this A β binding event disrupted the hydrophilic–lipophilic balance (HLB) of PKNPs.^{27,30} Moreover, the disassembly of PKNPs was further verified from the fluorescence spectra. As depicted in Fig. 2c, in the absence of A β , almost no fluorescence emission of

Next, selective photooxygenation of A β was monitored by matrix-assisted laser desorption/ionization time of flight mass spectrometry (MS). As described in Fig. 3a, a +16 Da modification was detected when A β was co-incubated with PKNPs and exposed to 450 nm laser irradiation, implying that A β was strongly oxidized by PKNPs. In contrast, PKNPs could not oxidize insulin under identical reaction conditions (Fig. S11†). These outcomes confirmed that PKNPs exhibited highly specific photooxygenation of A β . Furthermore, the photooxygenation of



The rigid BBB is the main obstacle that impedes the entry of most drugs into the brain.^{11,48} Recently, nanoparticles with intrinsic BBB-penetrability have been developed for neurodegenerative disease treatment.^{12,14,16} Therefore, we inferred that solid spherical nanostructured PKNPs exhibited great potential for traversing the BBB. Moreover, considering the photothermal effect of PKNPs, the BBB permeability of PKNPs could be improved under NIR irradiation.^{34,49} Thus, the ability of PKNPs to penetrate the BBB was studied by using *in vitro* BBB models based on a murine brain endothelial cell line (bEnd.3).^{50,51} As described in Fig. S15,[†] the penetration efficiency of PP and PKNPs across the BBB model was calculated to be 2.12% and 3.09%, respectively. Upon 638 nm laser irradiation, the penetration efficiency of PKNPs was further increased and reached 17.93%, proving that the BBB permeability of PKNPs could be enhanced by the photothermal effect. It should be noted that the transepithelial electrical resistance (TEER) values were stable throughout the experiment, suggesting that the mild photothermal effect did not impact BBB integrity.



Conclusion

In summary, we present the first example of A β -responsive activable PDT for treatment of AD by using nanostructured PKNP self-assemblies. Due to the photothermal effect of PKNPs, the BBB permeability of PKNPs under irradiation is 8.5 and 5.8 times higher compared to that of porphyrin and PKNPs alone, respectively. Moreover, attributed to the noncovalent interactions of the A β -targeting peptide KLVFF and PP, the spherical PKNPs show their ability for A β -driven disassembly and thus achieve A β -specific triggered ROS generation. As a result, PKNPs can act as an activable photosensitizer for selective photo-oxygenation of A β and inhibition of aggregation without off-target side effects. Furthermore, *in vivo* studies indicate that PKNPs can attenuate A β -mediated toxicity and extend the life span of CL2006 worms. Taken together, our designed switchable supramolecular self-assembly can realize selective and effective prevention of A β aggregation and related neurotoxicity in an AD model.

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