Design and Preparation of “corn-like” SPIONs@DFK-SBP-M13 Assembly for Improvement of Effective Internalization

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Purpose: Superparamagnetic iron oxide nanoparticles (SPIONs) have exhibited preeminent diagnosis and treatment performances, but their low internalization severely limits pre-designed functions. The low cell internalization is now an urgent bottleneck problem for almost all nanomaterials. To achieve more internalization of SPIONs, recombinant M13 phage was designed for targeted delivery and smart release.

Methods: M13 phages were designed to co-express exogenous SPARC binding peptide (SBP) and cathepsin B cleavage peptide (DFK), formed recombinant DFK-SBP-M13. 3.37±0.06 nm of SPIONs were modified by 3, 4-dihydroxyhydrocinnamic acid (DHCA) to gain 10.80 ± 0.21 nm of DHCA-coated SPIONs, i.e., DHCA@SPIONs. Upon adjusting the proportions of DHCA@SPIONs and DFK-SBP-M13, the multi-carboxyl SPIONs assembled onto recombinant M13 phages via covalent bonding. The assemblies were co-cultured with MDA-MB-231 cells to interpret their internalization and smart release.

Results: The “corn-like” SPIONs@DFK-SBP-M13 (261.47±3.30 nm) assemblies have not been reported previously. The assembly was stable, dispersible, superparamagnetic and biocompatible. After co-cultivation with MDA-MB-231 cells, the SPIONs@DFK-SBP-M13 assemblies quickly bond to the cell surface and are internalized. The enrichment rate of SPIONs@DFK-SBP-M13 assembly was 13.9 times higher than free SPIONs at 0.5 h, and intracellular Fe content was 3.6 times higher at 1 h. Furthermore, the DFK peptides favored cathepsin B to cleave SPIONs from the M13 templates resulting in release of SPIONs inside cells.

Conclusion: The novel SPIONs@DFK-SBP-M13 assembly can rapidly deliver SPIONs to the targeted sites and enabled smart release. The combination of genetic recombination and nanotechnology is beneficial for designing and optimizing some new nanomaterials with special functions to achieve wider applications.

Keywords: M13 phage, superparamagnetic iron oxide nanoparticles, self-assembly, smart release, targeting

Introduction

SPIONs, based on their excellent magnetic properties, changeable surface characteristics and low cytotoxicity, are gradually exhibiting an attractive prospect in the diagnosis and treatment of tumors, such as imaging analysis, drug loading, magnetic hyperthermia, magnetic targeted chemotherapy and gene therapy.¹⁻⁵ Similar to biomedicines with large molecular weight, SPIONs usually performed their functions at cellular levels. Only 0.7% of nano-formulations can reach the targeted tumor site due to two main bottlenecks: (1) the physical barriers, e.g., flow,

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diffusion and shear force, result in the erosion and low utilization of nanoparticles during the delivery process; (2) physiological barriers, including polymerization, protein adsorption, phagocytosis and renal or liver clearance, causing increased leakage and premature release.^{6,7} Such low accumulation of SPIONs around the targeted region usually leads to low uptake in targeted tissues, which consequently reduces the effectiveness of SPIONs.^{8,9} Furthermore, low cell internalization is a major concern with all nanomaterials that needs to be solved to achieve their potential high efficacy.

Recently, nanoparticles conjugation to functional aggregates caused rapid enrichment of small-sized nanoparticles and reduced systemic toxicity compared with the same concentration of large-sized particles.^{10,11} Furthermore, the functional scaffolds of free nanoparticles, assembled by low molecular weight polymers,¹² polypeptides,¹³ nucleic acids,¹⁴ and viruses can provide sufficient binding sites for nanoparticles and improve their utilization in vitro and vivo.^{15–17} With the advent of nanotechnology, SPION assemblies can possess various morphologies and multifunctionalities. Moreover, some specific robust anchors have been designed to accumulate nanoparticles at the targeted tissues via shortening transport time in blood and improving EPR (the enhanced permeability and retention) effect (Scheme 1).^{18–21} Among multitudinous scaffolds, temperate M13 phage has attracted the attention of researchers. Interestingly, M13 phage possesses cyclic single stranded DNA and 11 kinds of known proteins which could specifically infect *E. coli* containing F factor and is safe for humans.^{22,23}

Based on the fibrous structure having 1 μm length and a diameter of 6–8 nm, M13 bacteriophage possesses abundant binding sites and favors nanoparticles to line up so as to effectively avert steric hindrance.^{24,25} Meanwhile, the positive charge region from active amino sites at the umbrella-like α -helix N-terminus of pVIII also facilitates nanoparticles to bind M13 scaffold.^{26,27} Moreover, M13 phage is easy to perform gene recombination to express exogenous peptides with special roles,^{25,28,29} especially, M13 phage can be rationally modified for selective targeting and smart release by gene recombination.^{30,31} Most recombinant M13 phages are genetically stable, bioengineered quickly and in a cost-effective way.³² In addition, the lysogenicity of M13 phage is beneficial to enrich itself and its recombinant phages during large-scale preparation.³³ So M13 phages have been widely applied in biological, food and medical fields. Integration of the unique capabilities of M13 phage into designing nanoparticle assemblies

may offer some novel capabilities to SPIONs, such as rational molecular design via genetics, spatial control on a nanometer scale, and hierarchical assembly with 2D or 3D architecture.^{29,34,35}

To realize the rapid targeted absorption and internalization of ultra-small SPIONs, reduce leakage and unexpected release, and prolong the working time at the targeted site, the safe and recombinant M13 phage was used as template to prepare SPIONs assembly (Scheme 1). To control the self-assembly process of SPIONs, highly monodispersed carboxyl-coated SPIONs were prepared. Meanwhile, two exogenous functional peptides responding to the particular targeting and controllable release were co-expressed on the surface of M13 phage. Generally, this new kind of nanoparticle assembly will ensure SPIONs effective internalization with no side effects. In comparison to other modification techniques, phage-based self-assembly of SPIONs is easy to perform with few chemicals, that significantly reduces the production cost.³⁶

Materials and Methods

Materials

FeO (OH), oleic acid (AR), octadecene (99.5%), N-hexane (AR), ethanol (99.5%), phosphorus oxychloride (AR), polyethylene glycol (molecular weight 2000), 3, 4-dihydroxyphenylpropionic acid (AR), tetrahydrofuran, etc. All reagents are analytically pure.

MDA-MB-231 human breast cancer cells were purchased from the Cell Bank of the Chinese Academy of Sciences. M13 phage purchased from New England Biolabs.

Cell Culture

MDA-MB-231 human breast cancer cells were grown in RPMI-1640 medium supplemented with 10% FBS (EverGreen) and 1% penicillin-streptomycin mixture (Hyclone) at 37 °C.

Preparation and Surface Modification of SPIONs

Highly monodispersed SPIONs were prepared by the thermal decomposition method of FeO (OH) in 1-octadecene as previously reported.² To transfer oleic acid-coated nanoparticles into hydrophilic system, oleic acid ligands were exchanged by small molecules of 3, 4-dihydroxyphenylpropionic acid.³³ Firstly, 3, 4-dihydroxy-phenylpropionic acid (50 mg) and tetrahydrofuran (6 mL) was added into a 25 mL three-necked flask. 20 mg of SPIONs were

Scheme 1 Schematic diagram of M13 templated-superparamagnetic iron oxide assembly. **(A)** Schematic diagram of the nanoparticle assembly. **(B)** Function of the nanoparticle assembly.

added prior to incubation for 3 h at 50 °C. The reaction mixture was cooled and 1 mL 0.5 M of NaOH was added so as to precipitate SPIONs. 3, 4-dihydroxyhydrocinnamic acid (DHCA) coated nanoparticles were collected by centrifugation (3000 g, 5 min) and then dispersed in ddH₂O. Finally, nanoparticles were dispersed by ultrasound and filtered through a 0.22-micron filter membrane three times.

Preparation of M13 Phage Template

M13 phage template was constructed by representative phage dual display technique.^{15,28} SBP was inserted into the N-terminus of pIII protein on M13 phage at enzyme sites of *Acc65I* and *EagI*. Oligonucleotides encoding for SBP were 5'(Phos)-GTA CCT TTC TAT TCT CAC TCT TCA CCA CCG ACT GGA ATT AAC GGA GGC GGG TC -3' and 5'(Phos)-GGC CGA CCC GCC TCC GTT AAT TCC AGT CGG TGG TGA AGA GTG AGA ATA GAA AG-3'. Ligated DNA coding for T4 ligase was transformed into competent cells of XL-1 Blue cells, then competent cells were cultivated in plates containing SOC media as well as IPTG (Isopropyl β-D-Thiogalactoside), Xgal (5-Bromo-4-chloro-3-indolyl β-D-galacto-pyranoside) and tetracycline. White plaques were selected out and incubated overnight at 37 °C. DNA was isolated, purified and sequenced to confirm SBP insertion. After the successful insertion of the

genes encoding SBP, genes encoding DFK peptide were inserted into the N-terminus of pVIII protein at enzyme sites of *PstI* and *BamHI*. Oligonucleotides encoding for DFK were 5' (Phos)-GAT TTC AAG 3' and 5'(Phos)-GAT CCT TGA AAT CTG CA-3'. All the primers and target genes were synthesized by Suzhou Hongxun Company. All DNA was isolated using standard mini-prep kit (TIANGEN). All sequences were confirmed by DNA sequencing by Beijing Qingke Company.

Amplification and Characterization of M13 Phage Template

M13 template, SBP modified M13 template (labeled as rM13) and SBP-DFK modified M13 template (labeled as rrM13) were amplified by host cells.¹⁶ Briefly, phage templates were amplified by infecting specific host bacteria ER2738 at 37 °C under stirring for 4 h at 250 rpm. Cultures were centrifuged at 4500 g for 15 min, and supernatants were collected. Later phages were precipitated overnight at 4 °C by a mixture four times volume of 2.5 M of NaCl and 20% PEG8000 (w/v). Finally, phages were centrifuged at 12,000 g for 20 min, and re-suspended in TBS buffer for plaque forming assay.

Agarose gel electrophoresis was done to separate and quantify DNA of M13, rM13 and rrM13 phages. To evaluate the surface charge of phage templates, M13, rM13

and rrM13 phages were dispersed in 0.5 M of NaOH and 0.5 M of HCl solution with pH range from 2 to 13, respectively. Then the zeta potential under different pH value was measured via a dynamic light scattering laser instrument. To assess the molecular weight and molecular weight distribution of the phage templates, a trace number of samples were mixed with a substrate in a proportion of 1 to 10,000, solvent was evaporated at room temperature, and measured by laser assisted parsing/time of flight mass spectrometry.

Complexion of M13 Template with Carboxylated SPIONs

To construct the nanoparticle assembly, carboxylated SPIONs (30 μ L, 27 mM) activated by 1-ethyl-(3-dimethylaminopropyl) carbodiimide hydrochloride (EDC; 10 μ L, 20 mM) were mixed with M13/engineered M13 phages (10 μ L, 0.25 mM) activated by N-hydroxy succinimide (NHS; 10 μ L, 25 mM). Phage-nanoparticle complex was centrifuged (12,000 g, 20 min), harvested, and re-suspended in TBS or deionized water. The load capacity of rrM13 phage template was quantified via Fe content determination through inductively coupled plasma-atomic emission spectroscopy (ICP-AES)² and protein content with BCA Protein Quantification Kit (TIANGEN).

Characterization of SPIONs and M13 Templated SPIONs

The morphology and size of SPIONs, SPIONs@M13 and SPIONs@DFK-SBP-M13 assemblies were measured by transmission electron microscope (TEM). The lattice stripes of 3 nm-sized SPIONs were measured by high-resolution TEM. The crystalline nanoparticles were measured by X-ray diffraction analysis using Cu target ($2\theta = 20\text{--}80^\circ$). Characteristic absorption peaks of SPIONs were determined by Fourier transform infrared spectroscopy (FTIR, Equinox55). Weight loss rate curves were measured using Q500 thermogravimetric analyzer. Hysteresis loop of the sample was measured at room temperature via Mo 735 vibration magnetometer.

Cytotoxicity Test of SPIONs@DFK-SBP-M13 Assembly

The effect of the material on cell survival was determined by DOJINDO cell counting kit-8 (CCK-8) protocol. Cell seeding density was 5.0×10^4 cell/mL. Material concentrations were 0, 40, 80 and 160 μ g/mL.

Analysis of Targeting and Internalization of SPIONs@DFK-SBP-M13 Assembly

MDA-MB-231 cells were seeded with density 1.0×10^5 cells/mL into 6-well culture plate. After 24 h, the culture medium was removed and then rinsed twice with PBS buffer. SPIONs, SPIONs@M13 assembly or SPIONs@DFK-SBP-M13 assembly diluted with culture medium were added to reach the concentration of 40 μ g Fe/mL. Stationary culture (37 $^\circ$ C, 5% CO₂) was performed. At 0.5, 1, 3, 7, 18 and 24 h, cells were rinsed twice with PBS buffer, then trypsin was added to digest the cells. Later, samples were centrifuged and re-dispersed as a single cell suspension for hemocytometer. Fe content was determined as mentioned above. All experiments were done in triplicates.

Release Characteristics of SPIONs@DFK-SBP-M13 Assembly

SPIONs, SPIONs@M13 and SPIONs@DFK-SBP-M13 assembly were co-cultured with cells at a concentration of 80 μ g/mL. Stationary culture (37 $^\circ$ C, 5% CO₂) was performed. The distribution and morphology of three materials in MDA-MB-231 cells were measured by JEM-1230 TEM, fixed by 1% citric acid (30 min) and embedded by 618 epoxy resin (40 min, 33 $^\circ$ C).

Results and Discussion

Nanoparticle Surface Modification and Characterization

SPIONs were prepared by dynamic simultaneous thermal decomposition reported previously,² 3.37 ± 0.06 nm of oleic acid-coated SPIONs were obtained having uniform shape and size (Figure 1A–D). These ultra-small SPIONs possess cubic spinel configuration corresponding to the standard card No. 19–0629 with 1.63 Å lattice spacing (Figure 1C and E). The SPIONs were superparamagnetic due to no remanence (Hr) or coercivity (Hc) at room temperature (Figure 1K and L).

Because DHCA possessing two hydroxyls could replace oleic acids on SPIONs surface via ligand exchange,³³ it was utilized to convert hydrophobic (oil soluble SPIONs) to hydrophilic nature, so that binding ability between SPIONs and coat protein pVIII of M13 phage could be enhanced. After phase transfer, the oleic acid-coated SPIONs dispersed in chloroform (deep brown) were changed into DHCA-coated SPIONs having 10.80 ± 0.21 nm diameter in water (labeled DHCA@SPIONs).

Figure 1 Characteristics of SPIONs and DHCA@SPIONs. (A–D) individually show the stability in oil phase, TEM and size calculation of SPIONs; (E) shows the XRD pattern of SPIONs; (F–I) individually show the stability in water phase, TEM and size calculation of DHCA@SPIONs; (J) and (K) are respectively FTIR spectrums and the hysteresis loops at room temperature of SPIONs and DHCA@SPIONs; (L) is the fractionated amplification of (K).

(Figure 1F–I). The DHCA@SPIONs could stabilize the nanoparticles in aqueous phase (e.g., deionized water, PBS buffer and culture media) (Figure 1H) due to having -COOH from DHCA.³⁷ FTIR data in Figure 1J showed DHCA@SPIONs absorption peak of Fe at 590 cm^{-1} , C=O bond symmetric stretching vibration peak at 1429 cm^{-1} , methyl C-H bond vibrational absorption peak at 2800 cm^{-1} ; they also possessed the characteristic peaks of DHCA, e.g., new hydroxyl group O-H bond stretching vibration peaks at 3200 cm^{-1} and 920 cm^{-1} , and carboxyl group C=O bond stretching vibration peak at 1750 cm^{-1} . As to superparamagnetic peculiarity of DHCA@SPIONs, the M_s distinctly dropped to 29.98 emu/g after surface modification, due to decreased weight ratio of magnetic core (Figure 1K and L).

Construction and Characterization of Recombinant M13 Phage

To endow M13 phage template selective targeting and smart release, SPARC (i.e., polycysteine acid secretory protein) binding peptide (SPPTGIN, termed SBP) and

cathepsin B (CTSB) cleavage (termed DFK peptide) were designed to express at the surface of pIII protein (tail protein of M13 phage) and pVIII protein (major capsid protein of M13 phage), respectively. Based on high affinity of SBP to SPARC proteins, the tail-terminal SBP facilitates M13 phage to recognize and bind SPARC proteins that are abundantly expressed at the surface of cancer cells.³⁸ DFK peptide on the capsid protein is sensitive to cytosolic CTSB protease with smart release of SPIONs and beneficial to abundant DHCA@SPIONs anchorage (Figure 2A). Two recombinants were designed via genetic manipulation. The one containing the gene encoding SBP was named rM13. The other introduced two exogenous genes individually encoding SBP and DFK peptide was named rrM13. Meanwhile, SBP and SBP/DFK peptides were successfully expressed by rM13 and rrM13 phage, evident from mass spectrometry (Figure 2B). The characteristic peak at 3564.3 Da with a difference of 657 Da to that at 2931.8 Da , indicated SBP modification on pIII proteins of rrM13 phages because of molecular weight of SBP 657 Da . The characteristic peak of DFK-pVIII protein appeared at

Figure 2 Characteristics of recombinant M13 phage. (A) shows the whole molecular design of recombinant M13 phage; (B) is the MALDI-TOF Mass Spectrometry of wild and recombinant M13 phage; (C) presents the changes of Zeta potentials of M13, rM13 and rrM13; (D) exhibits the Zeta potentials of M13, rM13 and rrM13 at pH 7.4.

6113.9 Da but wild pVIII protein appeared at 5845.3 Da. Further, the isoelectric points (IEP) of M13, rM13 and rrM13 were 8.7, 6.3 and 6.3, respectively (Figure 2C). The nonpolar amino acids of SBP in rM13 rrM13 resulted in decline of IEP compared with M13, while DFK peptide did not change IEP due to zero electrostatic charge.⁴¹ On the other hand, surface charges of M13, rM13 and rrM13 were respectively +4.7 mV, -7.48 mV, and -6.59 mV at pH 7.4, which were beneficial to their stability (Figure 2D).

Construction of Recombinant M13-SPIONs Assembly

The carboxyl groups of DHCA@SPIONs interact with amino groups of M13, rM13 and rrM13 to form SPIONs complexes. As shown in Figure 3, when 0.5 mg/mL of phage templates

along with nanoparticle concentration 0.15 to 0.75 mg Fe/mL, the filamentous SPIONs-rrM13 (i.e., SPIONs@DFK-SBP-M13) assemblies increased (Figure 3A). However, SPIONs with 0.3 mg Fe/mL, line up along the coat protein pVIII like corn grains to form a “corn-like” complex (Figure 3B). Interestingly, SPIONs were increased to 0.4 mg Fe/mL, SPIONs-rrM13 assembly showed “grape-like” dendritic structure (Figure 3A-c and C). The more DHCA@SPIONs were used, the more carboxyl-binding sites were provided to bond with amino groups on pVIII protein, and fibrous SPIONs assemblies were inter-grafted with each other (Figure 3D and E).³⁹ As the synthesis scheme shows in Figure 3F, binding of DHCA@SPIONs induced M13 phage to bend due to combined action of coordination, covalent bonds, electrostatic forces, hydrogen bonding and π - π stacking, and finally SPIONs assemblies seemed to be less than 1 μ m.⁴⁰ The rrM13-SPIONs assemblies appeared as

Figure 3 Characteristics of the assemblies constructed by recombinant M13 phage and nanoparticles. **(A)** presentation of the morphology of rrM13-SPIONs by TEM, in which, Picture a to Picture e, 0.1, 0.3, 0.4, 0.5 and 0.75 mg Fe/mL SPIONs were added respectively; **(B–E)** are the fractionated amplification corresponding to **(A(b–e))**; **(F)** presents the assembly mechanisms between recombinant M13 phage and nanoparticles; **(G)** is the size calculation of "corn-like" rrM13-SPIONs; **(H)** is the hysteresis loops of "corn-like" rrM13-SPIONs at room temperature; **(I and J)** respectively show the parameter T_1 of free SPIONs and "corn-like" rrM13-SPIONs; **(K and L)** respectively exhibit the parameter T_2 of free SPIONs and "corn-like" rrM13-SPIONs.

closed aggregation at 0.5 mg Fe/mL of SPIONs (Figure 3A-d and D), and resulted in uneven reticulate structure with nanoparticle concentration of 0.75 mg Fe/mL (Figure 3A-e and E). It was imperative to adjust the dosage of DHCA@SPIONs and recombinant M13 phages for the formation of ideal recombinant M13-SPIONs assemblies.

Characterization of Recombinant M13-SPIONs Assembly

From dynamic light scattering data in Figure 3G, the hydrodynamic dimension of "corn-like" SPIONs-rrM13 (i.e., SPIONs@DFK-SBP-M13) complex was approximately 261.47±3.30 nm, suggesting that these assemblies were

Figure 4 The key characters of assemblies constructed by recombinant M13 phage and nanoparticles. (A) describes the cytotoxicity of rrM13-SPIONs complex; (B) presents time profiles cell uptake of DHCA@SPIONs (labeled as S1), rM13-SPIONs complex (labeled as S2), “corn-like” rrM13-SPIONs complex (labeled as S3), “grape-like” rrM13-SPIONs complex (labeled as S4); (C) presents specific uptake rates of S1, S2, S3 and S4.

relatively stable without agglomeration. The number of SPIONs per phage as per the concentration of iron and rrM13 phages in assemblies, were approximately 70.⁴¹ SPIONs-rrM13 assemblies still possessed superparamagnetic peculiarity and T1 imaging (Figure 3H–L), while their Ms dropped significantly to 0.25 emu/g, attributing to the reduced ratio of magnetic core by rrM13 templates.⁴²

Cytotoxicity of Recombinant M13-SPIONs Assembly

As to the toxicity of the recombinant M13-SPIONs complex on somatic cells, 0–160 mg Fe/mL DHCA@SPIONs, “corn-like” M13-SPIONs complex (i.e., SPIONs@M13 assembly) and rrM13-SPIONs complex (i.e., SPIONs@DFK-SBP-M13 assembly) had negligible effect on cell growth and proliferation of MDA-MB-231 cells (Figure 4A). At low concentration (40 µg Fe/mL), the three assemblies were almost non-cytotoxic and did not affect cell proliferation. Even at 160 µg Fe/mL, the cell survival rate was greater than 75%. The excellent cell compatibility of DHCA@SPIONs, M13-SPIONs and rrM13-SPIONs complex facilitates their application in biology and medicine.

Internalization of SPIONs and M13-SPIONs Assemblies in vitro

S1 (DHCA@SPIONs), S2 (M13-SPIONs complex, i.e., SPIONs@M13), S3 (“corn-like” rrM13-SPIONs complex, i.e., “corn-like” SPIONs@DFK-SBP-M13 assembly) and S4 (“grape-like” rrM13-SPIONs complex, i.e., “grape-like” SPIONs@DFK-SBP-M13 assembly) were selected to investigate the effect of complexes on selective targeting. The S1, S2, S3 and S4, having equal Fe

concentrations, were co-cultured with MDA-MB-231 cells. The Fe contents outside and inside the cells were measured during the culture process. In the first hour, four complexes showed a linear increase on cell surface and inside cells, and the Fe contents of four complexes were as follows S3 > S1 > S2 ≥ S4 (Figure 4B). As expected, SBPs at the end of rrM13-SPIONs complex displayed specific targeting ability to SPARC proteins secreted by cancer cells.⁴³ Consequently, SPIONs@DFK-SBP-M13 assemblies were captured and enriched on cytomembrane of MDA-MB-231 cells, which was conducive to transmembrane transport of nanoparticles.⁴⁴ In comparison to grape-like assemblies, the steric hindrance of corn-like fibrous material was less, which was beneficial to binding and internalization of “corn-like” SPIONs@DFK-SBP-M13 assemblies. Then the cell uptake of “corn-like” S3 was higher than that of “grape-like” S4.

Significantly, the cell uptake rate of S3 was respectively 2.86 and 13.9 times higher than that of S4 and S1 at 0.5 h as shown in Figure 4C. In comparison to non-targeting S1 and S2, the cell uptake rate of S3 reached the maximum and far exceeded that of S1 and S2. The cell uptake of S3 was 2.09 and 3.56 times more than those of S2 and S1 at 1 h (Figure 4B). Correspondingly, the specific uptake rate of S3 is 1.74 and 3.6 times higher than those of S2 and S1 (Figure 4C). The specific targeting of recombinant M13 phages promoted the endocytosis of nanoparticles, and an ordered arrangement of SPIONs along the fibrous configuration of M13 phages benefited the endocytosis of nanoparticles.^{45,46} Furthermore, the weak repulsive force among the negative-charged rrM13 phage templates and the cells benefited the vertical alignment of assemblies on the cell surface so as to improve

binding.^{47,48} These results were further testified by visible TEM images (Figure 5). The membrane boundary was obvious and the organelle structure was complete. The SPIONs were found at the surface of the membrane and in the cytoplasm. On the other hand, some nanoparticles seemed to be agglomerated upon the interactions between nanoparticles and active cells.⁴⁹ The total amount of S1, S2 and S3 was in the following order $S3 > S2 > S1$, which was consistent with the quantification results exhibited in Figure 4.

At 1 h, cell uptake of “grape-like” S4 was equal to that of fibrous S2 having no target peptide attached, because steric hindrance from a mass of blind corners in the dendritic structure of S4 decreased the specific binding capacity of SBPs. However, SPIONs were enriched on M13 phages, which still contributed to a higher cell uptake than free SPIONs i.e., S1. With increasing time in culture, large size nanomaterials reduced the endocytosis of S4.⁵⁰ Finally, SPARC protein attained saturation state, and the

targeting activity of S3 assembly was weakened, as a result, the uptake of S1 verged to S3 gradually.

Evaluation of Release of SPIONs@DFK-SBP-M13 Assembly in vitro

The distributions of SPIONs inside cells were observed by TEM after being co-cultured with MDA-MB-231 cells for 1 h (Figure 5). Generally, SPIONs distributed inside vesicles and lysosomes. S2, nanoparticle assemblies still possessed blocked configurations in cells, indicating SPIONs bonded on M13 templates were not shed obviously at 1 h (Figure 5F). The distribution of S3 in lysosomes is shown in Figure 5I. The morphology of S3 was looser and finely fragmented in comparison to that of S2 in lysosomes. Theoretically, S3 was easier to fragment than S2 due to predesigned CTSB-cleavable bridging fragments (DFK peptides) between pVIII protein and DHCA@SPIONs in S3. After S3 entered into cells,

Figure 5 Smart releasing of SPIONs from assemblies constructed by recombinant M13 phages and nanoparticles. (A, D and G) present respectively the whole cells treated with S1, S2 and S3; (B, E and H) are respectively the fractionated amplifications of cell surfaces treated with S1, S2 and S3; (C, F and I) are respectively the fractionated amplifications of lysosomes treated with S1, S2 and S3.

CTSB cleaved the spherical small-sized nanoparticles from M13 templates, which enabled smart release of SPIONs inside cells. Based on selective targeting and innovative release, the systemic damage of nanoparticles to cells could be effectively reduced. When S3 was co-cultivated with cells for 1 h, the mRNA level of CTSB was detected to increase 2 times higher than that without SPIONs, which strongly verified the TEM results.

Conclusions

By dynamic simultaneous thermal decomposition and ligand exchange, 10.80 ± 0.21 nm of hydrophilic DHCA@SPIONs were prepared; by gene recombination techniques, recombinant DFK-SBP-M13 phages were devised and gained to serve as the scaffolds for binding DHCA@SPIONs, with enhanced capabilities of recognition and conditional fragmentation. Under the assistance of EDC and NHS, the carboxyl groups of DHCA@SPIONs (0.3 mg Fe/mL) could react with amino groups of phage templates (0.5 mg/mL), forming 261.47 ± 3.30 nm of “corn-like” SPIONs@DFK-SBP-M13 assembly. Based on functionalized M13 phages, the superparamagnetic SPIONs@DFK-SBP-M13 assemblies could enhance the effective internalization of SPIONs via increased affinity to the targeted cells (Scheme 1). Moreover, SPIONs could be intelligently released in cells by specific shearing of cathepsin B. These specific binding and splice sites were constructed by predesigned functional groups that were provided by surface modification with chemical and biological methods. Especially, various functional groups can quickly be introduced onto M13 phages without affecting their original bio-functions. This functionalized modification can be utilized to design some specific nano-diagnosis and nano-treatment agents for cancers. Furthermore, integrating the advanced material preparation and gene recombination technology to improve the flexibility and variability of assembly structure would also be the prospect for nanomaterials.

Abbreviations

SPIONs, superparamagnetic iron oxide nanoparticles; SBP, SPARC binding peptide; DFK, cathepsin B cleavage peptide; SPARC, polycysteine acid secretory protein; M13, M13 phage; rM13, SBP modified M13 template; rrM13, SBP/DFK modified M13 template; CTSB, cathepsin B; DHCA, 3, 4-dihydroxyhydrocinnamic acid; TEM, transmission electron microscope; XRD, X-ray diffraction analysis; FTIR, Fourier transform infrared spectroscopy.

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Disclosure

The authors report no conflicts of interest in this work.

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